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THE EVALUATION OF SMALL AIRWAY DISEASE IN THE
HUMAN LUNG WITH SPECIAL REFERENCE TO TESTS
WHICH ARE SUITABLE FOR EPIDEMIOLOGICAL
SCREENING.

By

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Abstract

It has been known for several years that small airways of the lung (<2 mm internal diameter) may be defective in the absence of either clinical symptoms or abnormal function tests such as Forced Expiratory Volume in one second (FEV₁), Forced Vital Capacity (FVC) or Peak Expiratory Flow Rate (PEFR).

Interest in this "silent zone" has led to the development of special tests of small airway function. Some of these special tests including Frequency Dependence of Compliance and Deposition Patterns using radionuclide tagged particles are not readily adaptable to epidemiological screening. Others such as measurements of instantaneous flow rates on Maximal and Partial Expiratory Flow Volume (MEFV and PEFV) curves and the determination of Closing Volume (CV), are more suitable since they are obtained from gas flow at the mouth.

A relatively inexpensive apparatus was constructed and used to measure CV, FEV₁, FVC and instantaneous flowrates on MEFV and PEFV curves produced by healthy volunteers (asymptomatic smokers and non smokers), whilst breathing air or a mixture of 80% helium and 20% oxygen (He-O₂).

The position of a characteristic flow volume curve discontinuity (notch) was analysed in 28 volunteers (11 smokers) and was used to effect a measure of separation between the smokers and non smokers.

From the systematic differences in MEF 50 and MEF 25 obtained on the MEFV and PEFV curves of 10 males (3 smokers) it is tentatively suggested that the wide predicted normal range for these indices may be related to changes in airway smooth muscle tone.

An index of radionuclide particle penetration (Initial Lateral Penetration Index) was as expected positively correlated with PEFR ($r = 0.52$), FEV₁ ($r = 0.57$) and MEF 50 ($r = 0.44$) and negatively correlated with CV ($r = -0.49$) in 11 adults,

confirming that depth of deposition of particles is a sensitive function of airway health.

A total of 13 indices were ranked according to the sensitivity and specificity with which drug induced reversible airway changes were detected in a group of 25 volunteers (14 non smokers). Partial (P) indices (obtained from submaximal inspiration) were ranked:-

- (1) MEF 25 (P) (Air) and MEF 40 (P) (Air).
- (2) MEF 25 (P) (He); MEF 40 (P) (He) and Isoflow volume point (IFVP).
- (3) The excess flow rate on He-O₂ compared to Air {MEF 25 (P) (He/A) and MEF 40 (P) (He/A)}.
- (4) CV.

Correspondingly the ratings for maximal (M) indices were:-

- (1) MEF 40 (He).
- (2) MEF 25 (He).
- (3) MEF 40 (Air).
- (4) IFVP and CV.
- (5) MEF 25 (Air).
- (6) FEV₁.
- (7) FVC.
- (8) CV/SVC (%), MEF 40 (He/A) and MEF 25 (He/A).
- (9) FEV₁/FVC (%).
- (10) Slow vital capacity.

MEF 40 on both air and He-O₂ was superior to CV and both of these were superior to FEV₁ in detecting reversible airway changes.

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THE ROLE OF MEASUREMENTS WHICH DEPEND ON GAS FLOW IN AIRWAYS IN
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THE MOUTH.

Introduction

By the late 1950's, pulmonary function tests capable of detecting lung disease of the obstructive and the restrictive types were in widespread routine use. Despite this, Mead (1970) has pointed out that a quiet zone of the lung exists where changes are not detectable by many regularly used lung function indices, such as forced expiratory volume (FEV_1) and peak flow rate (PFR). The anatomical site of this quiet zone is thought to be in peripheral airways having internal diameter less than about 2mm. Anthonisen et.al. (1968) found that some bronchitic patients whose routine pulmonary function tests were within normal limits had regional abnormalities and attributed these abnormalities to obstructive lesions in small airways. Hogg et.al. (1968) using casts of the bronchial tree, concluded that the site of obstruction in chronic airway obstruction is in airways smaller than 2 mm internal diameter. Macklem and Mead (1967) used a retrograde catheter to measure the airflow resistance in central and peripheral airways and concluded that the peripheral resistance was a small component of total pulmonary resistance. Macklem (1972) reasoned that disease which lead to chronic airway obstruction must pass through a stage in which considerable peripheral obstruction exists whilst airway resistance (R_{aw}) and PFR remain within the predicted normal range.

For population studies, tests which aid in the diagnosis of early asymptomatic impairment of small airways need to be sensitive, specific, simple to perform and readily acceptable to the majority of persons in the population. Tests such as those measuring the frequency dependence of lung compliance or those involving the distribution of radioactive

gases or of the penetration of radionuclide tagged particles in the lung will not find ready acceptance for population studies. Tests such as these will be discussed only briefly. The reasons are that the one is unpleasant (a small balloon is enplaced in the subject's oesophagus) and the other has the potential of increasing the genetic radiation hazard to the population at large. In the following discussion emphasis will be given to tests which use gas flow at the mouth as the basis of detecting abnormal small airway function.

Techniques

Two classical methods exist for the assessment of airflow obstruction: plethysmograph measurement of airway resistance (R_{aw}) and measurements on forced expirograms. R_{aw} is largely dependent on the flow patterns in the bronchial tree which in turn is related to the cross-sectional area of the airway. The dichotomous branching of bronchi (Weibel, 1963) account for the decrease in internal diameter of the airways the more distal (from the mouth) being smallest. The total cross-sectional area of each generation increases (Thurlbeck, 1970) and in peripheral airways total cross-section is large (Horsefield, 1974). Linear gas velocities in distal airways must always be smaller than in airways nearer to the mouth. If laminar flow patterns can exist in the lung airways, (Pedley et.al. 1970) they are more likely to be found in distal than in central airways. In almost all circumstances therefore the major contribution to R_{aw} will be from central rather than from peripheral airways. Only gross obstruction of distal airways is likely to be revealed in this way.

During forced expiration, a rapid increase in flow is followed by a steady decrease as lung volume decreases. Mead (1967) has shown

that at a given lung volume maximum expiratory flow is dependent upon the elastic recoil pressure of the lungs and the resistance of airways between the alveoli and the "equal pressure point" (EPP). The EPP occurs where the pressure within the airway (intraluminal pressure) is equalled by the pressure outside the airway (pleural pressure). The EPP is not stationary but moves in the direction toward alveoli as lung volume decreases. The EPP divides the airways into an upstream portion (on the alveoli side) and a downstream portion (mouth side).

BODY PLETHYSMOGRAPHY

Airway resistance, (R_{aw}) and thoracic gas volume (Vtg), were measured in a constant volume body plethysmograph using the method described by Dubois (1956). Briefly, Boyle's law for gases at constant temperature is used to measure the change of lung volume in a subject as breathing manoeuvres are carried out from a sitting position inside an airtight box. A pneumotachograph detects box volume change as a pressure change. By panting against a closed shutter the pressure and volume of gas in the thorax is altered. If Vtg is considered to be mainly in the lungs (ignoring any negligible amount below the diaphragm), and dV is the change in volume due to compression of the chest by respiratory muscles, P and dP the alveolar pressure and change in alveolar pressure whilst the airways are occluded then:- $Vtg = P \cdot dV/dP$. Under static conditions P is effectively the barometric pressure less the vapour pressure of water at body temperature. During panting against a closed shutter dP is measured as mouth pressure (dP_m). Box pressure change dP_{box} and dP_m are displayed on an oscilloscope fitted with a protractor to measure the angle θ . The relationship:- $\tan \theta = dP_m/dP_{box}$ is used to estimate Vtg at or near to functional residual capacity.

When with the shutter open, the subject breathes quietly (flow rates about 0.5 l s^{-1}) box pressure change and pneumotachograph output (\dot{V}) are related:- $\tan \phi = \dot{V}/dP_{\text{box}}$. ϕ is measured from the oscilloscope trace. Raw is calculated from:- $\text{Raw} = dP_m/dP_{\text{box}} dP_{\text{box}}/\dot{V} \text{ kPa L}^{-1}\text{s}$. SRaw obtained from the multiplication of Raw by the $V_t g$ at which it was made results in a more stable index. θ and ϕ are the mean of five replicates.

CLOSED CIRCUIT HELIUM DILUTION METHOD FOR LUNG VOLUMES.

This involves the subject rebreathing from a closed circuit spirometer containing a known concentration of helium. This initial concentration of helium falls as the spirometer gas is mixed with the gases in the lungs, the fall in concentration being proportional to the volumes of gas in spirometer and lungs respectively. The helium concentration is measured by a katharometer, the spirometer is attached to a kymograph which records the respiratory movements. A pump ensures mixing as well as steady flow through the katharometer. The subject, seated upright and wearing a noseclip breathes into a mouthpiece which allows connection to room air or the closed circuit apparatus. When a regular breathing pattern has been established on room air the subject is turned in to the apparatus at end tidal level. He continues to breathe quietly, the CO_2 is removed by sodalime absorber and O_2 is added to the closed circuit to keep end tidal level constant. The helium concentration falls exponentially as the lung gases are mixed with those of the closed system and finally attains a steady state at which the value is noted. The subject is then instructed to expire to residual volume (RV) and inspire rapidly to maximum inflation (TLC). This is repeated before returning to normal breathing.

If V is the volume of gas in the closed circuit and He_1 and He_2

are the initial and final concentrations of helium, then expressing volumes at body temperature and pressure saturated with water vapour (BTPS), $FRC = V(\text{He}_1 - \text{He}_2)/\text{He}_2$ L BTPS.

Because the gas in the spirometer is at atmospheric temperature and pressure saturated with water vapour (ATPS) a correction factor is applied. For a spirometer temperature $t^{\circ}\text{C}$, barometric pressure P_B , and partial pressure of water vapour at $t^{\circ}\text{C}$ of $P_{\text{H}_2\text{O}}(t)$, the correction factor is given by:-

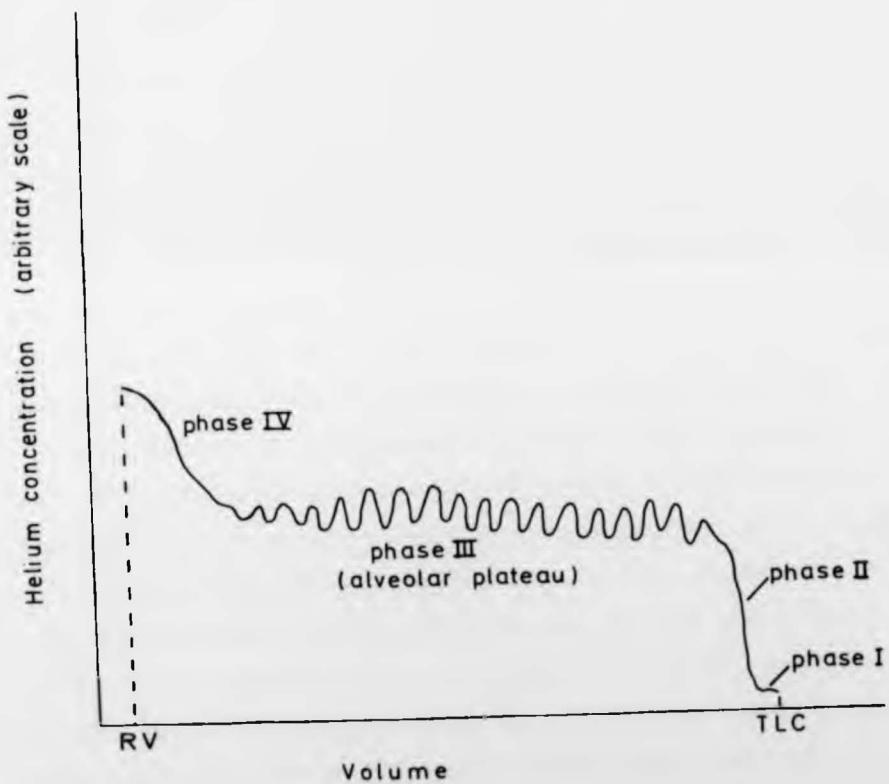
$$\text{ATPS to BTPS factor} = (37 + 273)/(273 + t). \quad \frac{P_B}{P_{\text{H}_2\text{O}}(t)} \cdot \frac{P_B}{P_{\text{H}_2\text{O}}(37)}$$

MEASUREMENT OF CLOSING VOLUME

The underlying principle of the "closing volume" (CV) measurement was first discussed by Dollfus et.al. (1957) and elaborated further by Holland et.al. (1970). It involves the inspiration of a tracer gas bolus, such as radioactive Xenon-133 or Nitrogen-13 or stable argon or helium, followed by measurement of the tracer concentration in the expirate. Alternatively the expirate is analysed for nitrogen, following a vital capacity (VC) inspiration of pure oxygen. The expired volume at which the alveolar gas concentration changes measurably from the alveolar plateau (that is from Phase III to Phase IV) is denoted as the "closing volume". At the "closing volume" airways in the dependent lung regions cease to contribute to gas flow measured at the mouth either because they close (Anthonisen et.al. (1969), Glaister, (1971) or cease being compressed, Hyatt et.al. (1971, 1973).

Typical "closing volume" tracings are shown in figure one following page, similar patterns have been observed by earlier workers (Fowler, (1963); Glaister, (1973) who noted the changes at the terminal portion of the alveolar plateau.

Closing Volume trace showing cardiogenic oscillations.



Earlier measurements of closing volume involved the use of relatively expensive laboratory equipment such as mass spectrometers and radiation detectors, but a simpler technique using an orifice gas analyser was described by Green et.al. (1972). The method used in this Thesis is essentially that of Green (1972) except that a Katspherometer was used instead of the critical orifice analyser. (The equipment is described later).

The manoeuvre begins with the subject, seated and wearing a nose clip, emptying the lungs to residual volume (RV). The breath is held at this volume and the rubber mouthpiece is placed in the mouth. At this point a tap on the mouthpiece is momentarily opened (allowing the 300 ml. bolus of pure helium to enter the inspiratory limb of the apparatus) and a steady vital capacity inspiration is made. At the end of inspiration (subject's lung volume at total lung capacity, (TLC)) a steady expiration back to RV is begun; the manoeuvre ends at RV. During the entire manoeuvre, the subjects are asked to control flowrates aided by the "lamp circuit" and the flow limiters in the limbs of the apparatus. Care is taken to ensure that the bolus inspiration is begun at RV and that no breath holding takes place at TLC. Inspiration of boli at lung volumes other than RV influences the direction of the closing volume transition (junction of Phases III and IV upward or downward and contributes to variability in the measurement of closing volumes. Breathholding at TLC decreases the precision with which the CV point is located (Susskind et.al. 1973).

The flowrate during the entire manoeuvre was monitored on a precalibrated storage oscilloscope and although only the expiratory phase was recorded on the XY plotter, care was taken to ensure that

inspired and expired vital capacities were similar. Tracings were discarded if either of these requirements was not met. Decision to discard was made at the time of testing.

The closing volume was determined from the tracings by drawing the best fit line by "eye" through the alveolar plateau and taking the CV as the volume corresponding to the point where the curve first deviated from the plateau. All curves were read by the author. Other methods of determining the phase III- IV junction were considered (for example the use of CUSUMS) but the extra complexity involved made such methods unacceptable for use in "field surveys". Curves having prominent cardiogenic oscillations were in general easier to read than those with less prominent or irregular oscillations; this was because the oscillations do not persist beyond the end of phase III.

MEASUREMENT OF FLOW-RATE AND VOLUME USING A PNEUMOTACHOGRAPH-PRESSURE TRANSDUCER-AMPLIFIER INTEGRATOR SYSTEM.

PRINCIPLE OF ACTION OF PNEUMATACHOGRAPH

When a gas flows through a tube in which there is a partial obstruction (resistance) the pressure beyond the obstruction is less than the pressure immediately before it. The difference in pressure (the pressure drop across the resistance) is a function of the properties of both tube and gas. For laminar flow, Poiseuille's Law gives the relationship:-

$$p = \left(\frac{8.1 \cdot \eta \cdot v}{\pi \cdot p \cdot r^4} \right)$$

where p is the pressure drop, Pascals (Pa)

v is the length of the tube in metres (m)

η is the viscosity of the gas in poises (a poise = gm. sec⁻¹.cm⁻¹)

\dot{V} is the volume of gas passing through the tube in one second

ρ = density of the gas, gm. cm⁻³; Kg.m⁻³

r is the radius of the tube (m)

To ensure that laminar flow conditions occur at the resistive element in a pneumotachograph, the resistance is placed at the centre of two narrow angled cones. The length and diameter of the cones is arranged to give a maximum in the diameter/length ratio at the resistance. (See diagram for the dimensions of the instrument used in this Thesis

). The pressure drop is measured by connecting a suitable instrument by small bore tubes to either side of the resistance.

The resistance can be of two types:-

- (i) a bundle of fine bore tubes (FLEISCH type or pneumotachograph)
- (ii) a fine mesh screen (screen pneumotachograph)

For a given gas or mixture of gases, the pressure drop across the resistance should be directly proportional to volume flowrate. The range of flowrates over which this condition is met can be satisfied by choice of resistance, diameter of bore at the centre of the pneumotachograph and the entry and exit diameters.

PRINCIPLE OF ACTION OF A PRESSURE TRANSDUCER.

A pressure transducer converts a difference in pressure at its input into an electrical signal at its output. The conversion can be of a variety of types including changes in capacitance or resistance or the generation of a voltage photovoltaically. Whichever type of conversion is used the object is to produce an electrical signal having a magnitude which is directly proportional to the size of the mechanical input.

The active element of the transducer is generally incorporated in a Wheatstone Bridge circuit. The circuit is generally arranged to be balanced when there is no input to the transducer. An input causes the bridge to become unbalanced; the unbalance signal is then detected and amplified electronically.

The pressure transducer used in the apparatus described in this Thesis was of the resistance type giving a direct current output (Statham type P15).

AMPLIFIER/INTEGRATOR.

Many pressure transducers give a voltage output which is typically only of the order of tens of millivolts. Such small signal levels are rarely useful without further amplification. The amplifier used needs to be sensitive, stable in operation and should not introduce spurious outputs ("noise"). The amplification factor (gain) should allow a useful output over a region of the amplifier's characteristics where signal distortion due to 'limiting' is minimum. The frequency response of amplifiers used in respiratory physiology need not in general be greater than from dc (zero frequency) to 20 Hz. Such a response would allow for a change in input signal from zero to maximum in 0.05 seconds.

The flow rate output of the transducer-amplifier can be converted into a volume signal by electronic integration. Integrators need to have a wide bandwidth as they need to respond to signals whose rate of change (dV/dt) cover an immense range.

The transducer amplifier system used in the apparatus described in this Thesis was assembled by the author. Basic general purpose integrated circuit instrumentation amplifiers were purchased and

assembled into an amplifier and integrator. The Philbrick type instrumentation amplifiers are of the operational amplifier type allowing a bandwidth of several KHz in the integrator.

The Amplifier

The Philbrick instrumentation amplifier (type P104) was connected as a differential input operational amplifier having a gain which could be set at any value between 10 and 500. The output impedance was made low (100 ohms) enabling direct connection to chart recorders or oscilloscope separately or in parallel. The amplifier was capable of giving an output of 10 volts without limiting.

The Integrator

The output of the amplifier was directly coupled to an integrator based on the above Philbrick (type P105) instrumentation amplifier. The output impedance of the integrator was low allowing the volume output to be connected to chart recorders, oscilloscope or other ancillary apparatus without the need for an additional 'buffer' amplifier.

The Amplifier/Integrator

The amplifier and integrator described above were housed together in a diecast box. The input and output connectors, gain control, an 'offset' control and a transducer impedance matching switch were mounted on the box. The power supply module was housed in a separate diecast box.

INSPIRATORY AND EXPIRATORY LAMP CIRCUIT

The Closing Volume manoeuvre requires that low (less than 0.5L sec.⁻¹) steady flowrates be maintained throughout. In order to help

subjects to achieve the desired flowrates, a high resistance is placed in series with the inspiratory and expiratory limbs of the CV apparatus. Even with added resistances flowrates could be unsteady. Viewing of the displayed flowrates on the persistence screen of an oscilloscope caused some subjects to have greater difficulty in achieving steady flows; this was because they tended to use the buccal cavity as a reservoir for the expirate. A simple circuit whose output was used to light an inspiratory and an expiratory lamp was devised. The arrangement was such that the brightness of the lamps was proportional to flowrates. The lamp assembly was easily viewed by the subject who found little difficulty in maintaining a steady brilliance in the glow of the appropriate lamp.

The circuit is basically a zero crossing detector. Small hysteresis and rapid response with high sensitivity was achieved by using an operational amplifier. The input to the circuit was provided by a parallel connection to transducer amplifier's output. The gain of the zero crossing detector was adjustable and was set to provide current to the lamps. The magnitude of the current flowing was proportional to flowrates in the range 0.3 to 0.5 L sec^{-1} . Outside this flowrate range the output current was disconnected from the lamps as shunting diodes diverted the current. The subject was instructed to maintain the appropriate lamp glowing with the same brilliance for as long as he could. Reversal in flowrates (from inspiratory to expiratory) was easily achieved without a pause.

The One Second time marker.

The forced expiratory volume in one second can be obtained from the same forced expiration that produces the maximum expiratory flow volume curve if a time marker can be placed on the tracing. The timer should be started as soon as the expiration begins, and the tracing can be marked by a momentary raising of the pen or by grounding the flow signal. Since the FEV_1 is often desirable for comparison with other indices derived from the MEFV curve, subject effort is saved and throughput is increased by the incorporation of such a marker on the flow volume apparatus.

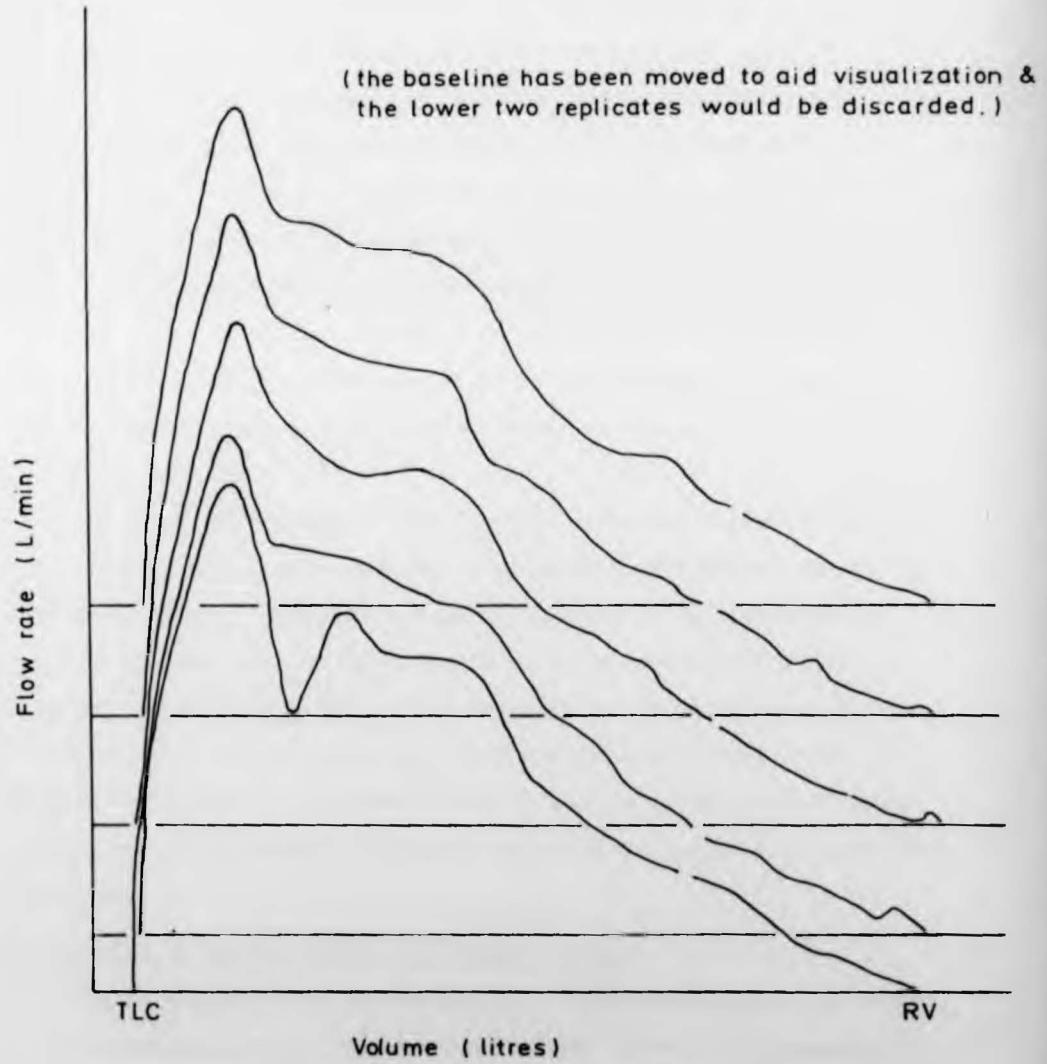
The block diagram of the apparatus shows the time marker (used only on Flow Volume apparatus Mk 11). The timer was initiated by the vel me signal at the output of the Integrator; in this way flowrate dependence of the 1sec. mark was eliminated. The trigger point could be accurately set between 0.05L and 0.1L; in this range flowrate independence was maintained. The timer was accurate to better than 1 part in 1000 (1 msec). The output of the marker actuated the penlift mechanism (P) of the XY plotter for about a millisecond. An example of a marked MEFV tracing is shown.

PRINCIPLE OF ACTION OF THE KATAPHEROMETER.

The Katapherometer is an apparatus for continuous gas analysis. It is based on the thermal conductivity principle. A low pressure in the analysing chamber is used to give a response time suitable for making dynamic measurements of helium and carbon dioxide in respiratory gases.

In the Godart apparatus, (used for this Thesis), the measuring head consists of two cells provided with platinum filaments.

Example of Maximal Expiratory Flow Volume tracings,
showing FEV₁ (indicated as a break in the tracing)



The cells form part of a "Wheatstone Bridge".

Measurement of unbalance in the bridge is facilitated since it is fed by an oscillator which maintains a constant current of about 100 milliamperes. Unbalance of the bridge produces an alternating current (A C) signal which is amplified and demodulated to provide a deflection on a meter calibrated in helium concentration.

The thermal conductivity of a gas mixture depends on the concentrations of the components. The gas passes through one pair of analysing cells; these cells have platinum wire stretched along their axes. The gas to be analysed flows past the filament in one cell, but the other cell is surrounded by a reference gas and sealed. The temperature of the two filaments depend on heat loss to the surroundings, and their resistance is proportional to the absolute temperature. If the concentration of a component gas in the mixture being analysed changes, the cell's resistance changes and causes an unbalanced bridge.

A vacuum pump draws the gas into the measuring head at about 0.3 litres min.⁻¹ and also maintains a pressure in the cell of about 50 mm of mercury. The cell volume is stated to be 1.2 ml and the volume flowrate through it at the 50 mm Hg. pressure in the cell is said to be 15 litres min.⁻¹. The cell volume is replenished in about 5 msec. The thermal inertia of the filament may be of the order of 40 msec. The response time of the instrument could be about 2.5 times this limiting response if delay due to feedline is taken into account.

PRINCIPLE OF HELIUM-OXYGEN FLOW VOLUME CURVES.

Flow volume curves produced whilst a mixture of helium and oxygen is breathed are called helium-oxygen curves, (He-O₂). The mixture

usually consists of 80% Helium and 20% Oxygen but the proportions need not be accurately known.

Helium is less dense but more viscous than air. Calculations of the density and viscosity of mixtures of gases can be made from a knowledge of the relative proportions of the component gases. At a temperature of 20°C and pressure of 760 mm of mercury, the 80:20 He-O₂ mixture has a density which is 0.44 that of moist air and a viscosity which is 6.9% greater. The kinematic viscosity (viscosity/density) of the mixture was calculated to be 2.43 that of moist air. Since viscosity and density have dissimilar temperature coefficients and reference values are not available at all temperatures and pressures, the calculated values serve only as a guide.

Laminar flow is independent of density but dependent on viscosity, whereas in turbulent or disturbed flow the converse is true. Flow in smaller peripheral airways are more likely to be laminar than flows in the larger central airways (Mead, 1967). Changing the respiratory gas will therefore affect the flow volume curve directly but such a change will also affect the MEFV by altering the equal pressure point (EPP) (Mead, 1960).

At lung volumes close to total lung capacity (TLC), when the EPP is in central airways (Cotes, 1975) and the flowrate are therefore predominantly density dependent, He-O₂ flowrates may exceed those on air. The ratio:- $MEF50(\text{He-O}_2) - MEF50(\text{Air}) / MEF50(\text{Air})$ may then be positive (greater than unity). As lung volume decreases, the EPP moves towards the lung periphery into airways of smaller internal diameter.

Flow through the airways then becomes more laminar and therefore less density dependent, being primarily governed by the resistance of the

airways between the alveoli and the EPP.

If measurements of flowrates at lung volumes close to residual volume reflect small airway calibre, then the ratio:-

$$(\text{MEF25}(\text{He-O}_2) - \text{MEF25}(\text{Air})) / \text{MEF25}(\text{Air})$$

could be negative or equal to unity. Further since lung volume and hence the vital capacity (VC) is largely independent of the physical properties of the gas breathed, a point should exist in the VC range of most subjects where He-O₂ and Air flowrates are of equal magnitude. The lung volume at which this equality in flowrates is achieved has been called the isoflow volume point (IFVP) by Bode, 1975 and the point of identical flow (PIF) by Despujol.

The He-O₂ and Air MEF25 curves are usually compared by direct superimposition of the two tracings (BODE, 1975). The IFVP may however be calculated from measurements of flowrates at least two points in the VC range. In this Thesis calculation of IFVP was made from measurements of He-O₂ and Air flowrates at 50% and 25% of VC (RV taken as 0% vVC).

The position of the isoflow volume point in healthy and diseased airways

In young healthy airways, the rise in intrapleural pressure caused by contraction of the respiratory muscles initially assists active expiration. The intrapleural pressure also tends to collapse the airway as the opposing force due to the traction of the lung tissue and the intraluminal pressure decrease progressively with decreasing lung volume. The equal pressure point could therefore be very close to or even at RV. (In young healthy persons RV may be determined by the elasticity of the thoracic cage (Cotes, 1975). If EPP is taken as dividing the airways into an upstream (viscosity dependent) and

downstream (density dependent) segment then the IFVP may occur at or close to RV.

As the airways age RV is determined by the elastic recoil pressure of the lung and the calibre of the small airways (Cotes, 1975), the IFPP may therefore be located nearer to the mouth of an older healthy subject than in a younger healthy one. Consequently the position of the IFVP may be found at a higher percentage of the VC in the older subject.

In obstructive lung disease obstructed small airways may be decreased in calibre and require less pressure to cause them to close with the consequence that RV may also be increased (Cotes, 1975). The IFVP could then be expected to be found at progressively higher percentages of the VC. However, diminished elastic recoil pressure (as for example in emphysema) may also cause RV to be higher than in health since the additional pressure needed to open these airways may not be available from the musculature. The proportion of the airway tree which exhibits density dependence may thus be similar in both the obstructive and restrictive types of airway disease. If the restriction is due to a process which increases elastic recoil pressure (as in pulmonary fibrosis) the RV may be diminished compared to health and the proportion of the airway tree in which density dependent flow occurs may be increased. In such a case IFVP may be expected to be at a lower percentage of the VC.

In summary, an IFVP can be expected to be found somewhere in the vital capacity range. The position of IFVP as a percentage of VC is higher in older than in younger subjects. However in diseases which decrease airway calibre the IFVP should occur at a higher percentage of VC. In diseases which cause a relative increase in

downstream (density dependent) segment then the IFVP may occur at or close to RV.

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In summary, an IFVP can be expected to be found somewhere in the vital capacity range. The position of IFVP as a percentage of VC is higher in older than in younger subjects. However in diseases which decrease airway calibre the IFVP should occur at a higher percentage of VC. In diseases which cause a relative increase in

airway calibre the IFV_F occurs at a lower percentage of the VC than would occur in health.

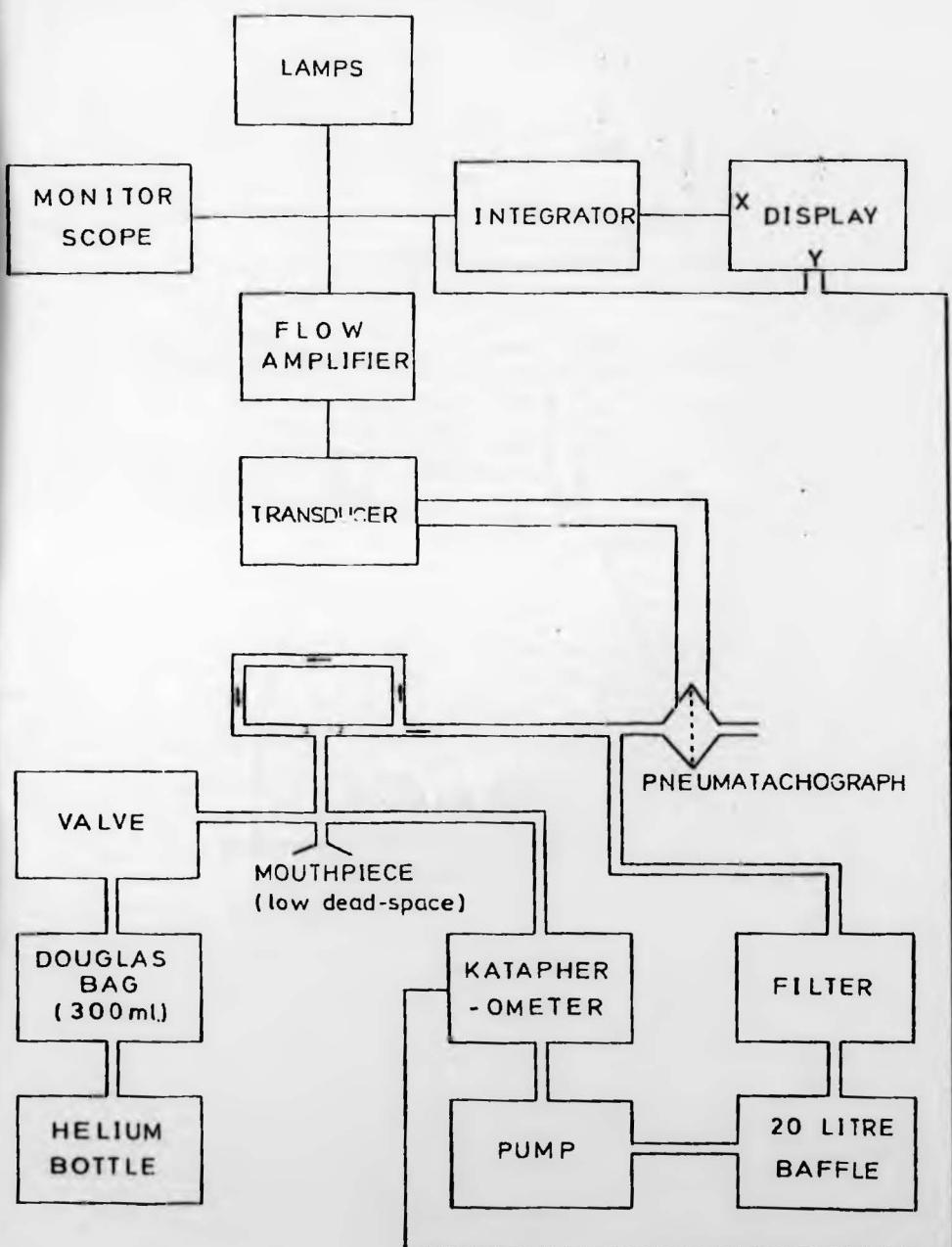
APPARATUS FOR THE MEASUREMENT OF CLOSING VOLUME AND MAXIMAL EXPIRATORY FLOW VOLUME CURVES.

The measurement of Closing Volume (CV) and the tracing of maximal expiratory flow volume (MEFV) curves require that gas flow at the mouth be measured. For the CV expiratory volume is required to be plotted on the abscissa against marker gas concentration on the ordinate. An MEFV curve is a plot of forced expiratory flowrates on the ordinate against lung volume on the abscissa. These requirements allow for the use of components which are common to both determinations. In the apparatus being described here maximum use was made of common components.

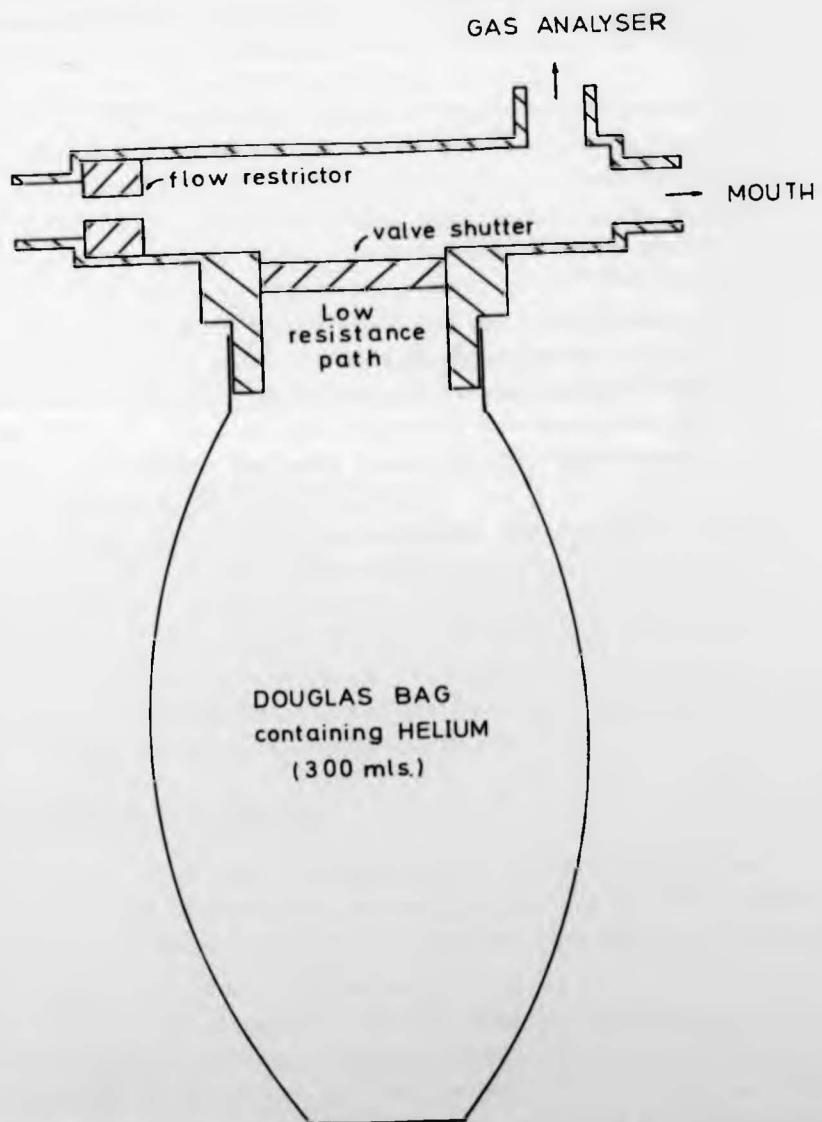
Gas flowrate was detected by a common pneumotachygraph pressure transducer amplifier/integrator system. Volume was obtained as the integral with respect to time of the instantaneous flowrate. The underlying principles of the technique has been discussed above. The common display system was an X Y plotter, but a persistence screen oscilloscope was also shared.

Apparatus having appropriate frequency response, stability and linearity was designed and assembled by the author in consultation with Mr.E.Reeves, an electronic engineer from the Medical Research Council's Environmental Physiology Unit in this School. Special features of the apparatus are the simple 'lamp circuit' (used to help subjects maintain steady flowrates on the CV manoeuvre) and the one second marker which gave the FEV₁ on the MEFV curve. A special low dead space mouthpiece for the CV breathing port was made by the workshop staff.

A schematic diagram of the Closing Volume apparatus is shown. Marker gas, (helium) was admitted to the inspiratory limb via a low resistance port when the mouthpiece valve was opened. A diagram of the specially designed low dead space mouthpiece assembly is shown. The volume of marker gas was therefore inhaled as a bolus followed by room air. On exhalation the expirate was sampled at a constant rate and the helium concentration measured by the Katsapherometer before being returned to the expiratory arm via the pump, baffle and filter. The pneumotachograph



Block diagram of Closing Volume apparatus



Low dead space mouthpiece assembly: closing volume apparatus.

senses both inspiratory and expiratory flow rates and the flow signal operates the lamp circuit. Microswitches mounted on the actuating arm of the mouthpiece valve controlled the penlift mechanism of the X Y plotter so that the inspiratory phase of the manoeuvre was not recorded. A storage oscilloscope monitored the flowrate during the entire manoeuvre.

The block diagram of Flow volume apparatus (Mk 2.) is shown.

Both the inspiratory and expiratory flow rates are sensed thus enabling a complete flow volume loop to be drawn if desired.

By observing inspired volume, submaximal inspired volumes can be reproducibly obtained. This feature is particularly useful when partial expiratory flow volume (PEFV) curves are being plotted.

The earlier flow volume equipment (Mk 1.) excluded the one second marker. The penlift output from the marker circuit could be overridden when not desired as for example on PEFV curves or on MEFV curves when a gas other than air was being breathed. When only air is breathed the He/O₂ Reservoir, CO₂ absorbed and rebreathing circuit is not used.

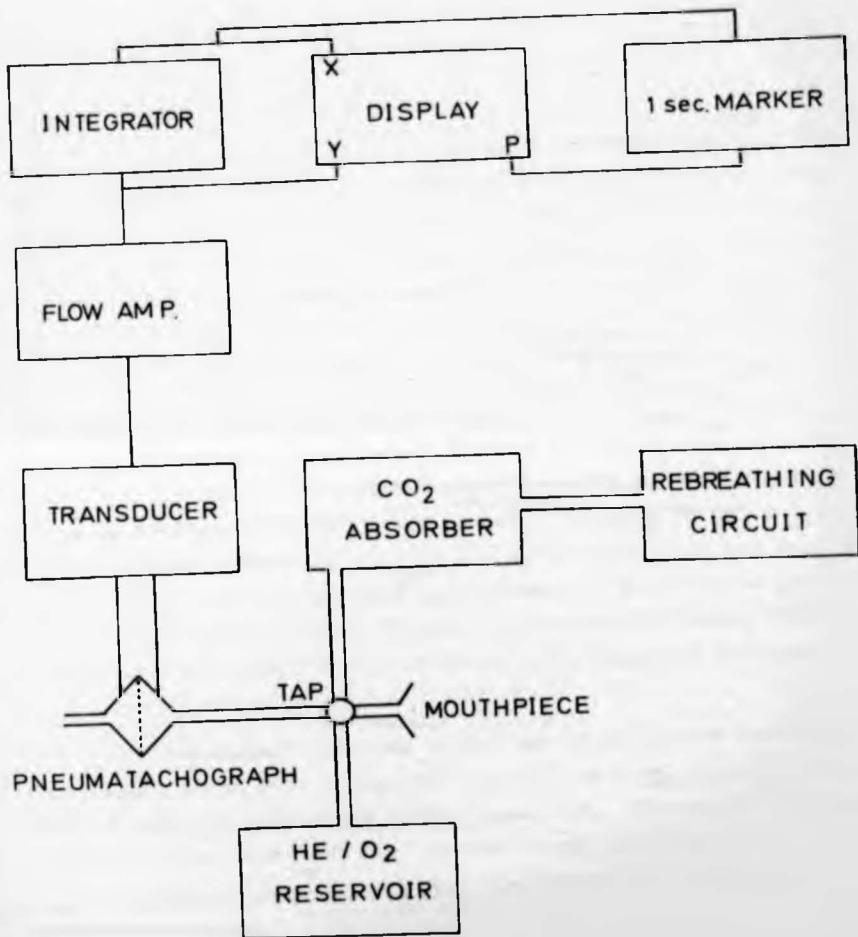
For the generation of helium-oxygen flow volume curves, the full circuit of the flow volume apparatus was used to provide a rebreathing circuit and a reservoir of the correct helium-oxygen concentration. The purpose of the rebreathing circuit was to economise in gases. In addition, the gain of the transducer amplifier is reduced (by switched attenuator) to compensate for the lowered density of the mixture compared to air.

CALIBRATION OF THE APPARATUS.

The calibration of the apparatus can be divided into two parts; (1) the response and linearity of the gas analyser, (2) the stability and accuracy with which volume and flow rate can be measured.

(1) Calibration of the katapherometer.

Because a fixed volume of helium would be used (300 ml) as the marker gas the final height attained by the helium concentration will be dependent on the apical residual volume of the volunteers. For example a person having half the residual volume in the independent



Block diagram of Flow Volume apparatus (Mk.II)

lung regions as another, would demonstrate a final peak concentration which would be twice that of the person with the larger apical residual volume. The linearity of the analyser to different helium gas/air concentrations was therefore checked. The result is shown as Fig.1. The dilutions used were checked on a recently calibrated helium analyser (Cambridge Analyser).

The response time of the Katapherometer was measured. The X Y plotter was operated with a time base signal (variable speed settings cm sec^{-1}) fed to the X axis and the helium concentration on the Y axis. The time was set running before a tap connecting the helium gas to the analyser was opened. The response time was the time taken for the analyser to record 90% of its steady state value. This 0 - 90% response time was 440 milliseconds.

(2) Volume and flow rate calibration.

The CV manoeuvre takes place at very low flow rates. It is therefore important to measure volumes accurately, and be aware of leaks or electronic drift in the system. The sampled gas was returned to the measuring circuit and both inspiration and expiration was passed through the same pneumotachograph. Differences between inspired and expired fixed volumes could be due to leaks, drift or difference in pressure flow relationship for opposite direction of flow in the pneumotachograph.

The volumetric response of the apparatus was determined by discharging gases (air or helium-oxygen) from a calibrated syringe (1.5 L) into the mouthpiece of the apparatus. The amplifier/integrator was then adjusted so that the correct volume was displayed on the X Y recorder. Both rapid and slow discharges into the mouthpiece was made.

The flow rate response was determined by connecting a rotameter in series with the mouthpiece. The flow rate was assessed at both high and low flowrates.

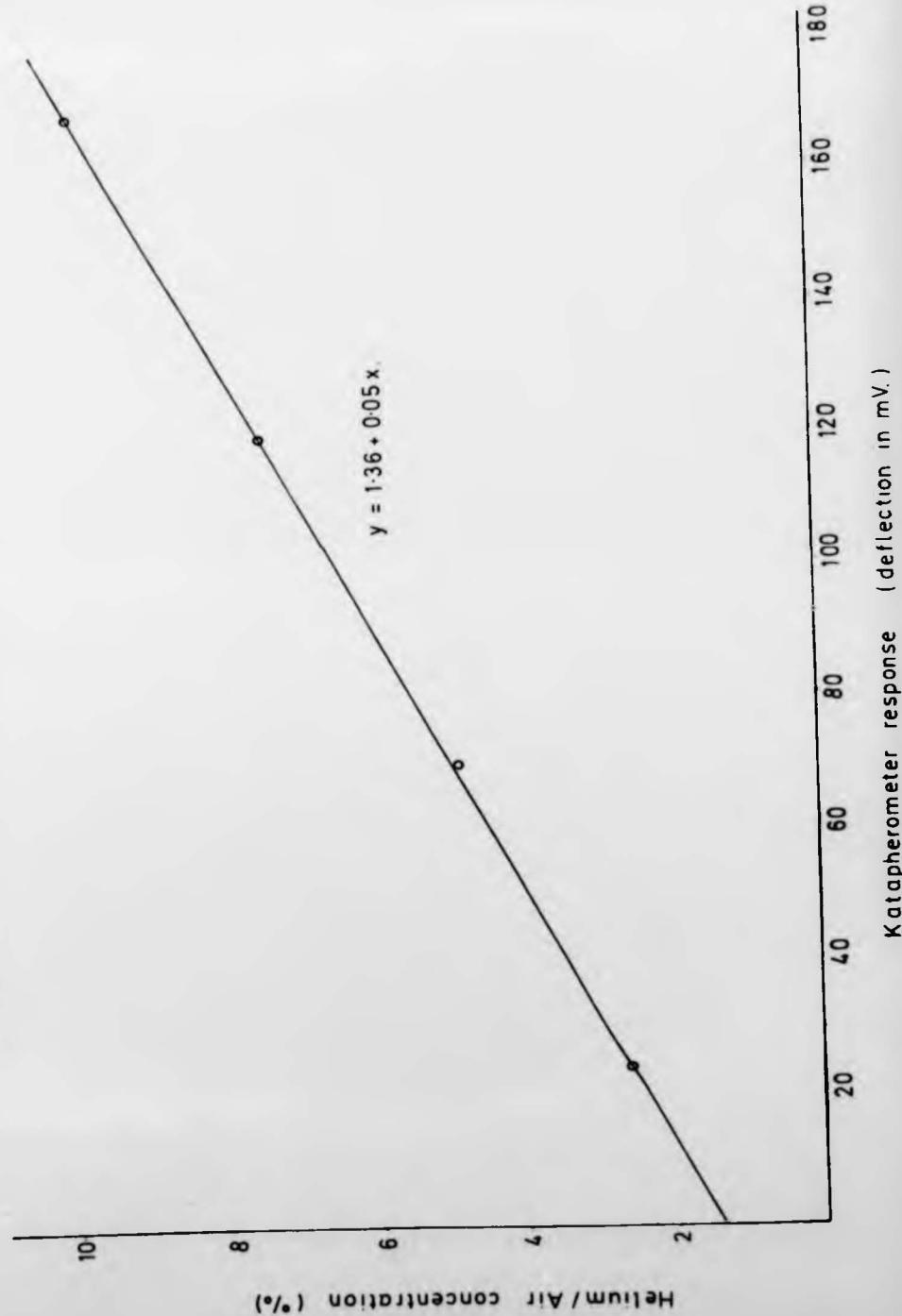
The calibration graphs are shown.

Dynamic calibrations of the flow volume apparatus was carried

out by direct comparison of volunteers performance on the apparatus with those on other apparatus which had been previously been calibrated.

An account of both the short-term (one day) and long-term (several months) behavior of the apparatus is given below.

FIG. 1 Katapherometer response to Helium in Air.

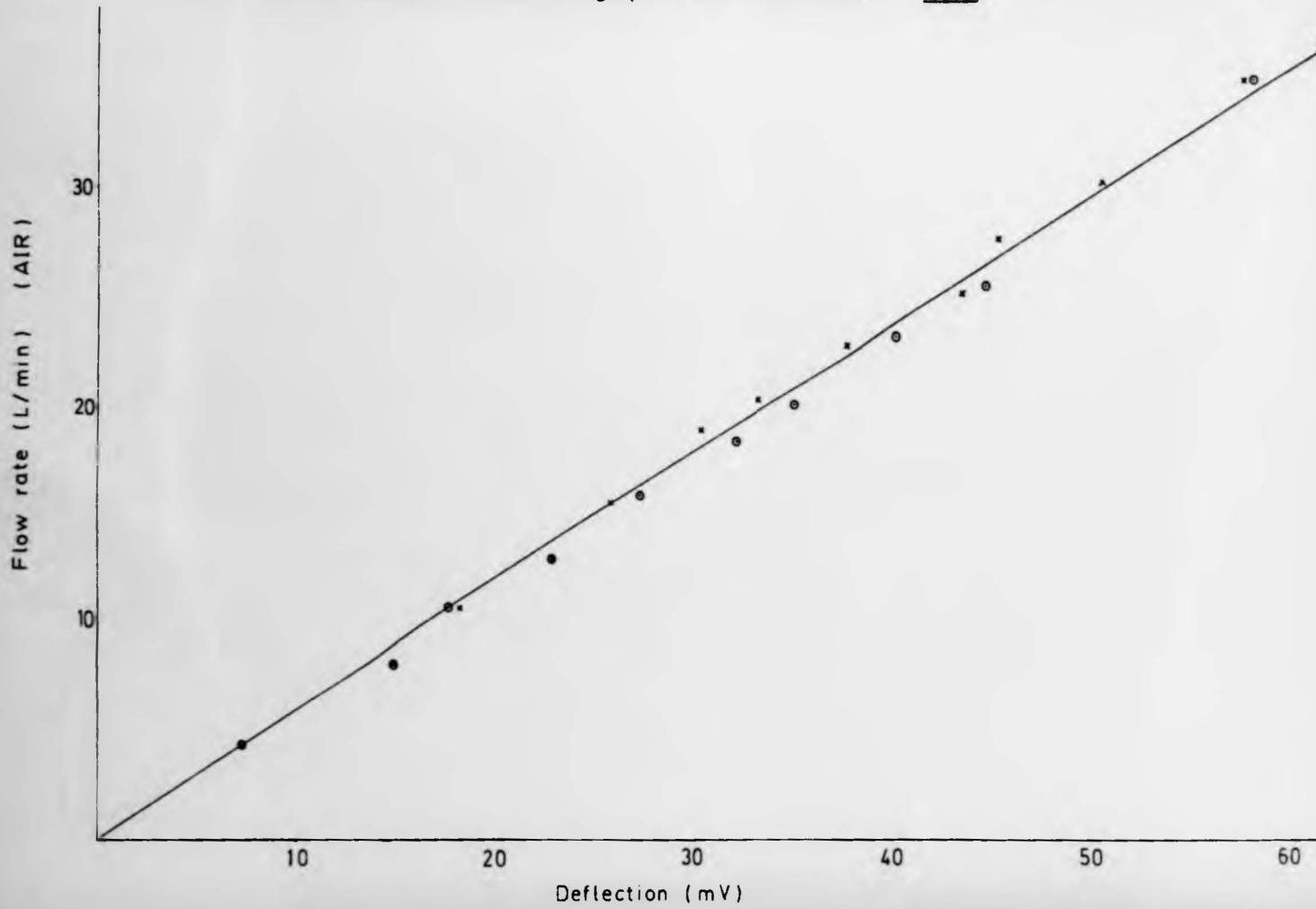


HeLiUm / Air concentration (%)

Katapherometer response (deflection in mV)

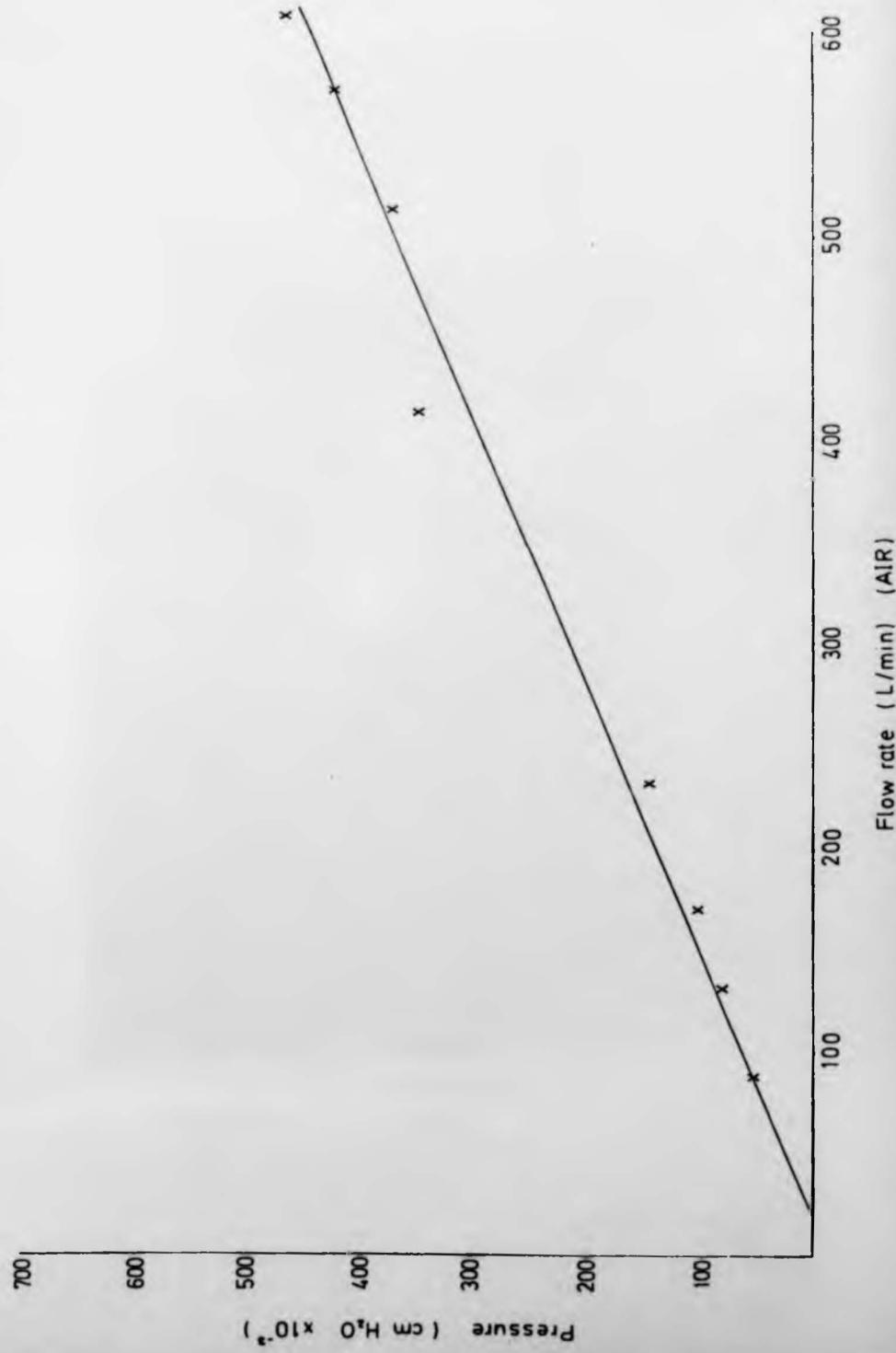
Calibration of Heated Pneumatachograph at low flow rates

FIG. 2.



Calibration of Heated Pneumotachograph

FIG. 3.



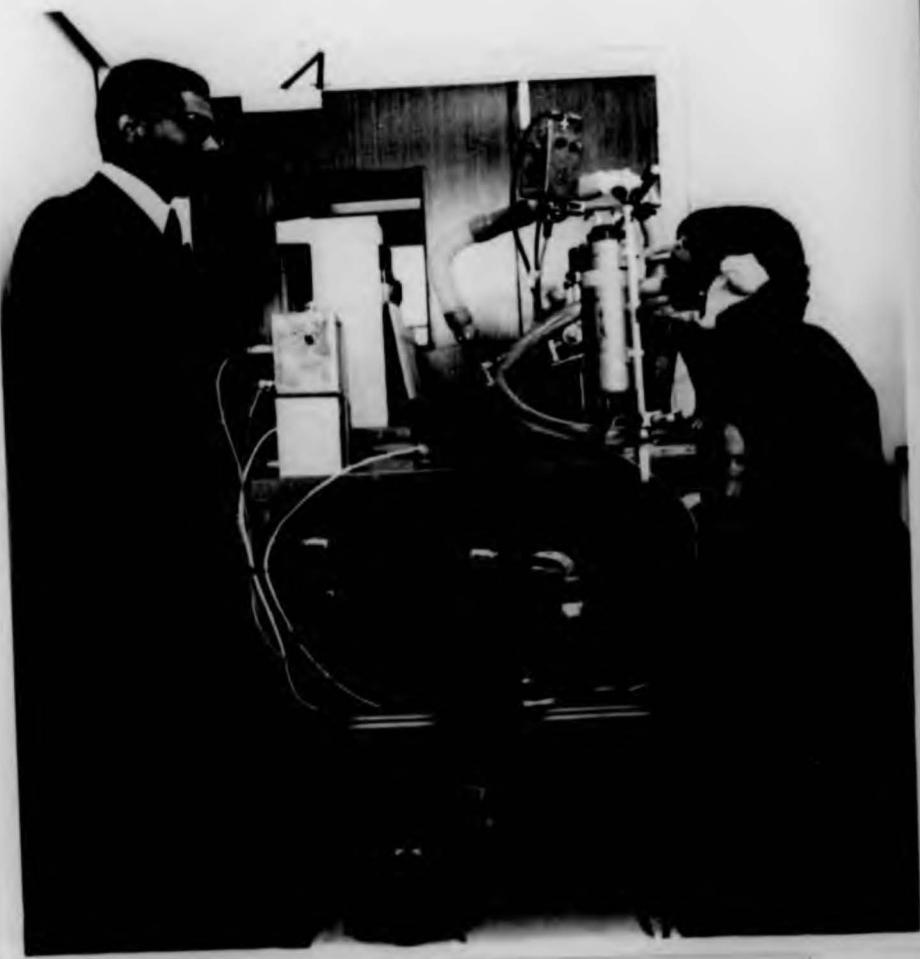


PHOTO.1.

Showing the relationship between volunteer
and observer during the tests.

The gas analyser head and lamp circuit is
seen near to the volunteer.

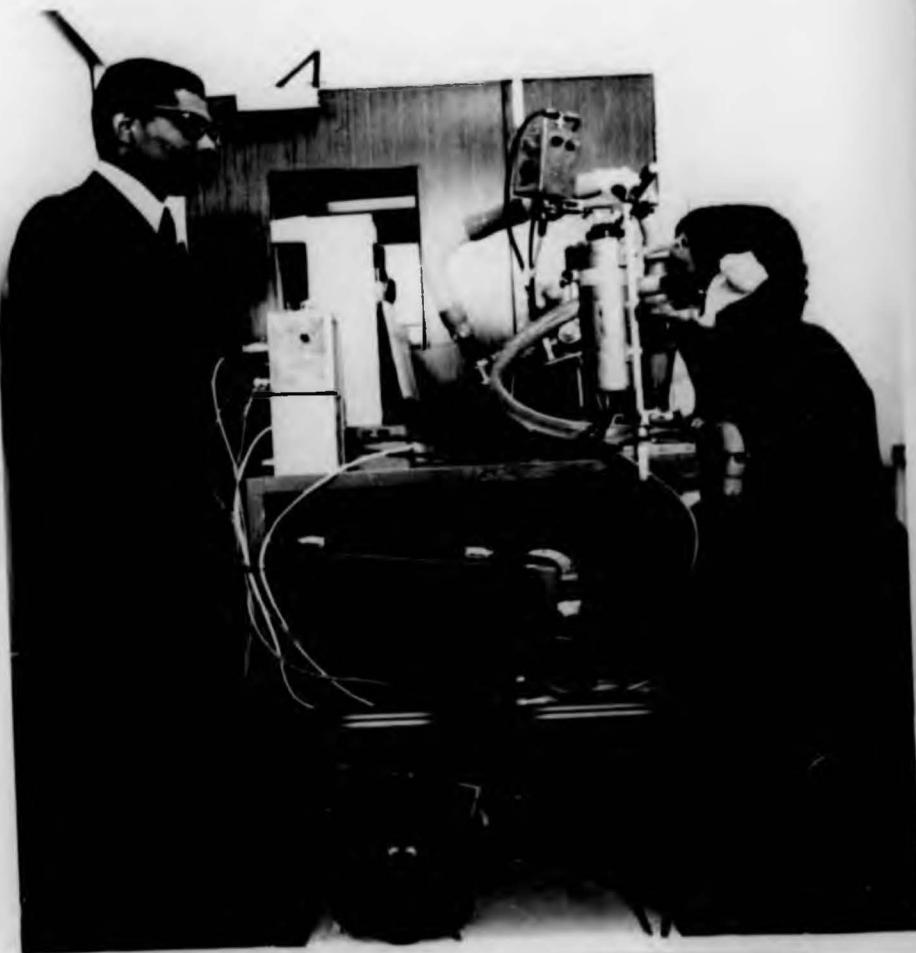


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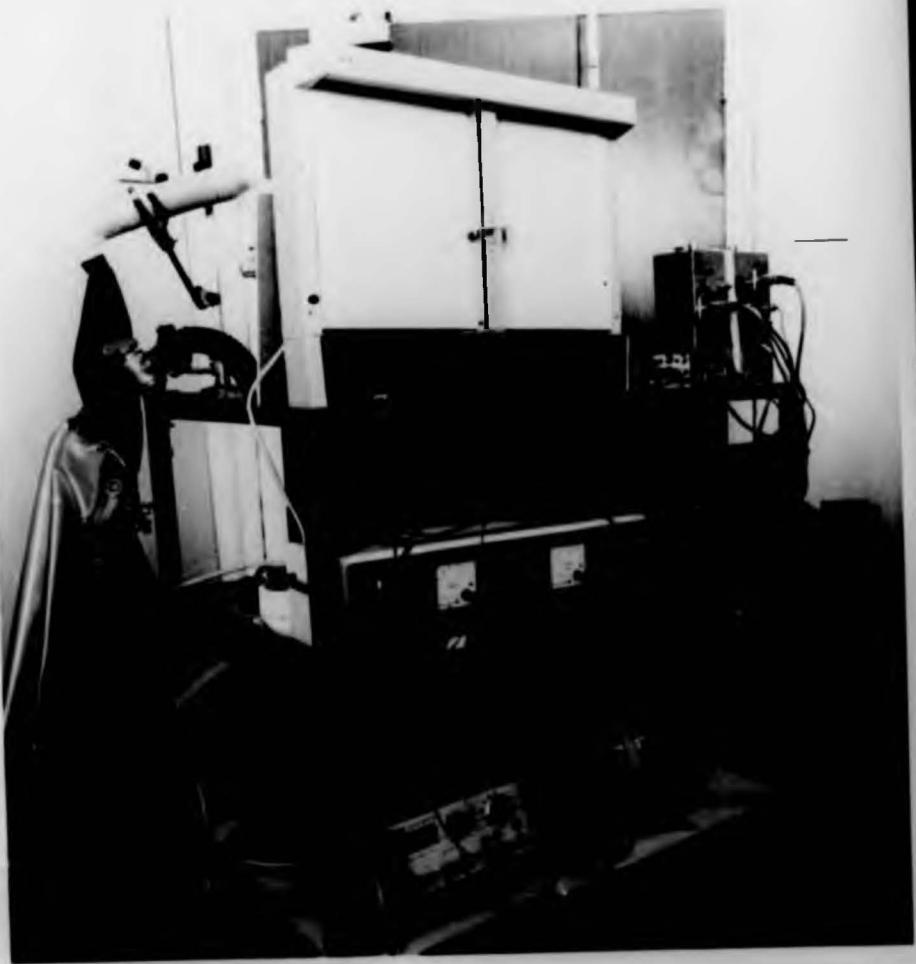


PHOTO.2.

View of equipment showing:-

- (1) Top shelf:- Display (XY) plotter and amplifier/integrator modules.
- (2) Middle shelf:- Monitor oscilloscope and gas analyser amplifier.
- (3) Bottom shelf:- Baffle and electronic calibrator module.



PHOTO .2.

View of equipment showing:-

- (1) Top shelf:- Display (XY) plotter and amplifier/integrator modules.
- (2) Middle shelf:- Monitor oscilloscope and gas analyser amplifier.
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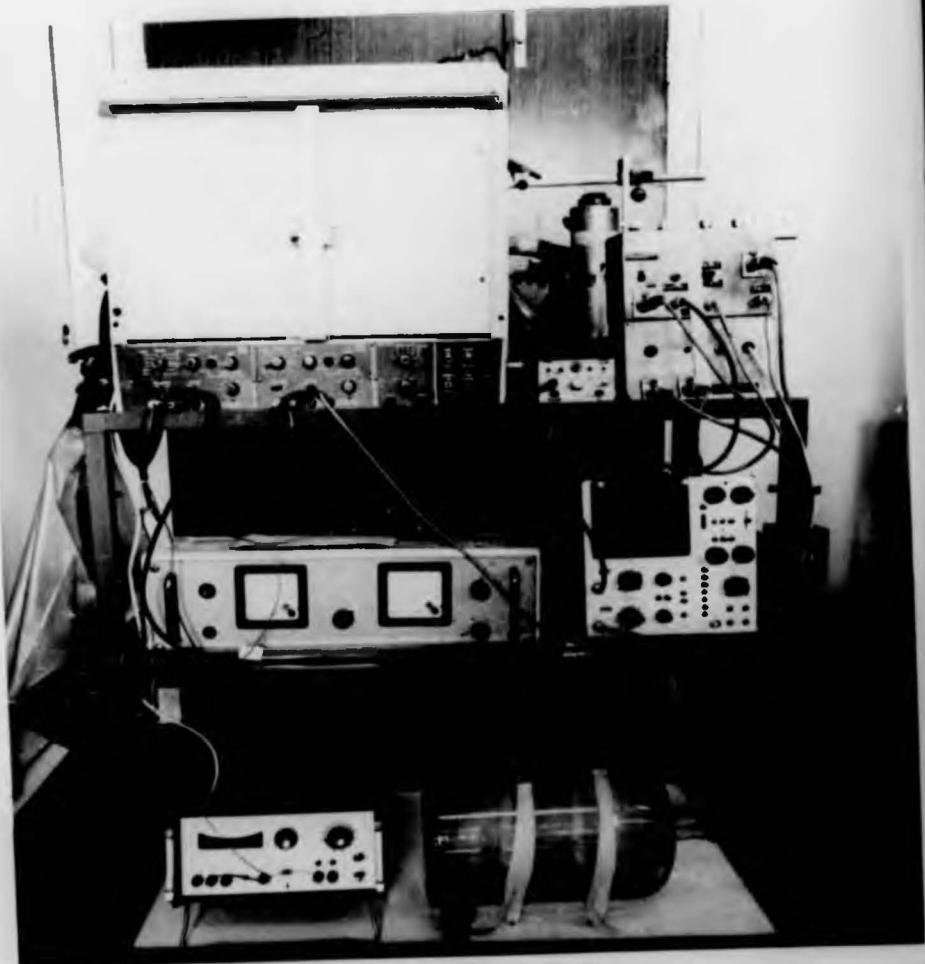


PHOTO 3.

Shows operator view of the apparatus.

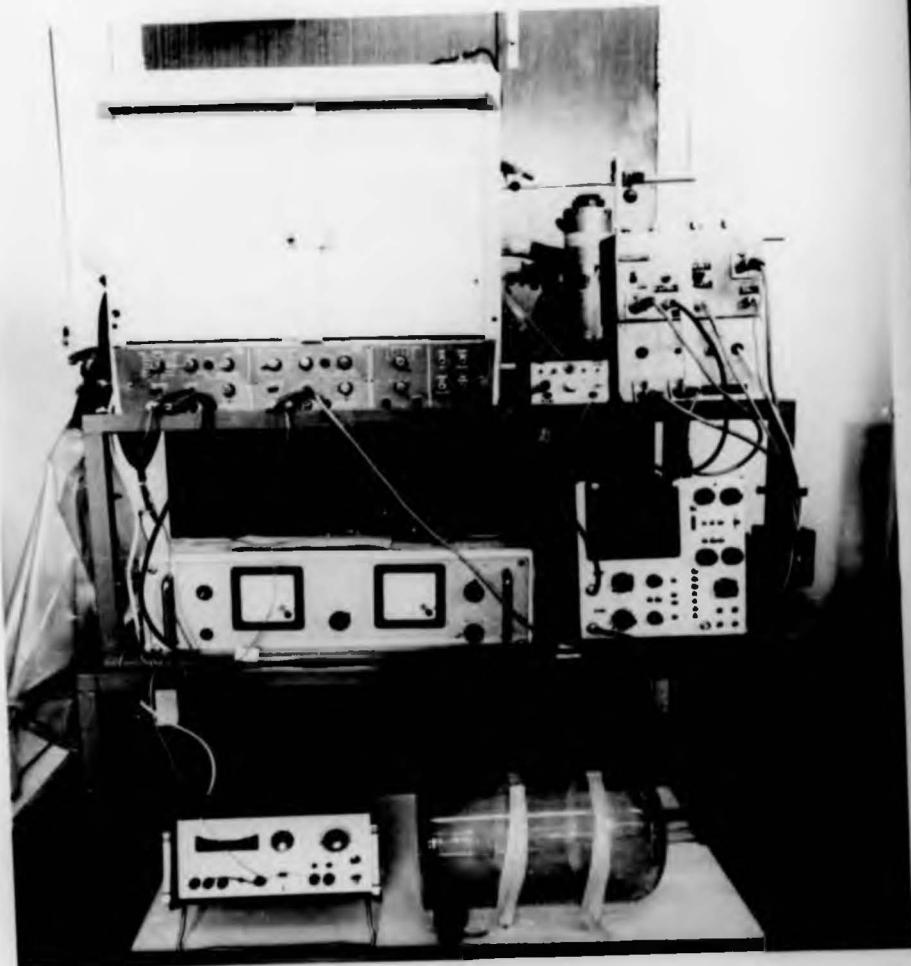


PHOTO 3.

Shows operator view of the apparatus.

CHAPTER 2

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MEASUREMENTS MADE ON FLOW VOLUME APPARATUS COMPARED WITH THOSE ONDRY SPIROMETER AND PEAK FLOW METER. (Short-term).

On the same day, results from the following three recently calibrated instruments (1,2 and 3) were compared by testing seven asymptomatic adults in randomised order. As usual readings were corrected to BTPS.

1. A "vitalograph" wedge spirometer was used and from the tracings FEV₁, FVC and the maximal midexpiratory flow rate (MMF) were obtained.
2. The peak expiratory flow rate (PEFR) was measured on a Wright peak flow meter.
3. MEFV curves were made on the Mk.2 flow volume apparatus (previously described) and the indices:- FEV₁, FVC, MEF50, MEF40, MEF25 and PEFR determined.

The mean, standard deviation (SD), standard error of the mean ($SEM = SD \times n^{-\frac{1}{2}}$) and the coefficient of variation (COV) based on the last three of five "blows" ($n = 3$) are given in the table together with the subjects' sex and age. Analysis of variance (ANOVA) and Regression were the statistical techniques used in analysis of the results.

Results.

The overall mean (mean for the seven subjects) shows that measurements of the same index on different instruments did not differ greatly: the vitalograph's mean FEV₁ was slightly larger than that on the flow volume apparatus (difference 0.08L), flow volume FVC was larger than vitalograph (by 0.2L) and the flow volume's PEFR was higher than the Wright's Peak flow meter by $29L \text{ min}^{-1}$. The overall

- 2 -

COV (calculated as the overall SD as a percentage of the overall mean) reflects the relative stability of the instruments. Again identical indices give COVs which are not very different; for example FEV_1 19.6% on vitalograph compared with 17% on the flow volume apparatus. By ANOVA, Snedecor's F values of 0.4, 0.26, 4.07 and 1.06 were found for FEV_1 , FVC, $FEV_1/FVC(\%)$ and PEFR and in no case were these values significant at the 5% level.

The MMF (converted to $L \text{ min}^{-1}$) was also not significantly different from the MEF50 ($F = 0.38$). The between subject variations in MEF⁴⁰ and MEF25 (measured by the COV), on the same instrument were however statistically different ($F = 7.11$, $P < 0.05$).

The seven mean values were used in a linear regression analysis to find equations relating the various indices. Under these circumstances differences in the subjects' physical characteristics do not influence the regressions since each subject was tested on all the instruments. The following equations were found:-

$$(i) \quad MEF^{40} (L\text{min}^{-1}) = 0.75 MEF50 (L\text{min}^{-1}) + 4.37;$$

$r = 0.992$, t slope = 39.3 $P < 0.001$.

$$(ii) \quad MEF50 (L\text{min}^{-1}) = 1.31 MEF^{40} (L\text{min}^{-1}) + 9.61;$$

$r = 0.992$ t slope = 39.3 $P < 0.001$.

$$(iii) \quad MEF25 (L\text{min}^{-1}) = 21.32 MMF (L\text{sec}^{-1}) + 4.04;$$

$r = 0.73$, $t = 5.35$ $P < 0.005$.

$$(iv) \quad MMF (L\text{sec}^{-1}) = 0.025 MEF25 (L\text{min}^{-1}) + 1.58;$$

$r = 0.73$, $t = 5.35$, $P < 0.005$.

$$(v) \text{ MEF50 (Lmin}^{-1}\text{)} = 61.97 \text{ MMF (Lsec}^{-1}\text{)} + 5.43;$$

r = 0.87; t = 8.86 P<0.005.

$$(vi) \text{ MMF (Lsec}^{-1}\text{)} = 0.012 \text{ MEF50 (Lmin}^{-1}\text{)} + 0.806$$

r = 0.87; t = 8.86 P<0.005.

Conclusions

The results indicate as expected that the various indices gave the same value when measured on recently calibrated instruments. This finding though expected, should never be assumed to hold since systematic errors can be larger than intersubject variations.

It is emphasized that although the subjects here were members of staff of this School they were not "trained subjects".

The regression relationships though based on a small sample show significant correlations between the indices as would be expected. The equations should be useful as a guide in relating for instance the MMF and the MEF50. The relationship between MEF50 and MEF40 in normal subjects will be valuable in transforming measurements made at these percentages of the vital capacity.

TABLE 1.

Subject	VITALOGRAPH			FLOW		VOLUME		CURVES		PEAK FLOW METER	
	FEV ₁ L BTPS	FVC L BTPS	MMF LSec ⁻¹	FEV ₁ L BTPS	FVC L BTPS	V50 LMin ⁻¹	V40 LMin ⁻¹	V25 LMin ⁻¹	PEFR LMin ⁻¹	PEFR LMin ⁻¹	
1/M/36/S											
Mean	4.28	5.21	4.39	3.90	5.48	293	218	87	711	647	
SD	0.08	0.10	0.20	0.13	0.05	13	7	8	10	12	
SE	0.05	0.06	0.11	0.07	0.03	8	4	4	6	7	
COV	1.9	1.2	2.5	1.8	0.6	4.4	3.2	8.6	1.4	1.9	
2/M/36/XS											
Mean	3.24	4.31	3.75	3.21	4.84	159	107	42	562	533	
SD	0.03	0.03	0.14	0.06	0.05	3.8	7.5	3.8	27	46	
SE	0.02	0.02	0.08	0.03	0.03	2.2	4.3	2.2	16	27	
COV	0.9	0.7	3.7	1.9	1.0	2.4	7	9	44.8	8.6	
3/F/23/NS											
Mean	2.37	2.71	2.62	2.51	2.89	165	126	66	475	383	
SD	0.05	0.03	0.06	0.22	0.05	7.5	9.9	6.5	13	15	
SE	0.03	0.01	0.03	0.13	0.03	4.3	5.7	3.8	7.5	8.7	
COV	1.3	1.1	1.2	8.8	1.0	4.5	7.8	9.9	2.7	3.9	
4/M/25/NS											
Mean	4.48	6.16	3.39	4.36	6.13	235	174	96	577	587	
SD	0.05	0.04	0.02	0.03	0.05	13	7.5	7.5	4.0	2.9	
SE	0.03	0.02	0.01	0.02	0.03	7.5	4.3	4.3	2.3	1.7	
COV	1.1	0.7	0.6	0.7	0.8	5.5	4.3	7.8	0.7	0.5	
5/M/39/NS											
Mean	3.99	5.22	3.37	3.79	5.02	252	172	77	664	655	
SD	0.13	0.17	0.06	0.09	0.08	7.5	9.9	7.5	8.1	13.2	
SE	0.08	0.10	0.03	0.05	0.04	4.3	5.7	4.3	4.7	7.6	
COV	3.3	3.3	1.8	2.4	1.6	3.0	5.8	9.8	1.2	2.0	

Table 1 (continued)

6/11/30/NS

Mean	3.91	4.41	5.47	3.92	4.65	351	263	139	678	661
SD	0.02	0.03	0.16	0.08	0.03	11.3	9.9	9.9	4.0	1.2
SE	0.01	0.02	0.09	0.05	0.02	6.5	5.7	5.7	2.3	0.7
COV	0.5	0.7	2.9	2.0	0.7	3.2	3.8	7.1	0.6	0.2

7/M/41/NS

Mean	3.47	5.23	2.33	3.36	5.62	152	116	61	525	527
SD	0.03	0.03	0.03	0.03	0.05	7.5	9.9	3.8	14	6.4
SE	0.02	0.02	0.02	0.01	0.03	4.3	5.7	2.2	8	3.7
COV	0.9	0.6	1.3	0.9	0.9	4.9	8.6	6.1	3	1.2

Overall Mean	3.68	4.75	3.62	3.58	4.95	230	168	81	599	570
Overall SD	0.72	1.09	1.07	0.61	1.04	76	57	31	87	100
Overall SE	0.27	0.41	0.40	0.23	0.39	29	22	12	33	38
Overall COV(%)	19.6	22.9	29.6	17.0	21.0	33	34	38	15	18

*Key:- Subject Number/Sex/Age/Smoking category

V50 Maximum expiratory flow rate at 50% of vital capacity.

V40 Maximum expiratory flow rate at 40% of vital capacity.

V25 Maximum expiratory flow rate at 25% of vital capacity.

NS Non-smoker

XS Ex-smoker

S Smoker

CHAPTER 3

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SCATTER DIAGRAM SHOWING TEMPERATURE RELATIONSHIP OF FEV ₁	Fig	5

COMPARATIVE MEASUREMENTS OF THE FORCED EXPIRATORY VOLUME IN ONE SECOND
(FEV₁) ON A BEILOR'S SPIROMETER (VITALOGRAPH) AND THE FLOW VOLUME
APPARATUS FITTED WITH A ONE SECOND MARKER. (Long term).

Over a period of several months a total of 105 persons were tested on each of two instruments to determine their forced expiratory volume in one second (FEV₁). The subjects blew into the instruments in succession on the same occasion, care being taken to ensure that approximately half of the subjects were tested first on different instruments. Usually no more than five persons were tested on a single day and the subjects were comprised of members of staff, students and visitors to this school. For this comparison, the volunteers did not complete a questionnaire on respiratory symptoms, and only their name was taken for reference in order to avoid counting the results from the same person twice. There were five blows into each instrument and the mean of the last three blows was taken. As usual, volumes were corrected to BTPS. Results were logged at the time of testing.

Results

The table gives the results obtained in the 105 subjects. The overall mean FEV₁ measured on the flow volume apparatus was 3.49 ± 0.91 L BTPS and that on the vitalograph 3.60 ± 0.98 L BTPS and the mean difference of 106 - 316 ml was statistically significant at the 1% level.

The regression line obtained from the 105 means is shown in Fig. 1 together with the correlation coefficient and the standard error of the mean. The overall coefficient of variation (COV) indicates that the spread of results on each instrument were almost identical (COV 26% and 27% for flow volume apparatus and vitalograph respectively).

Figs. 2 and 3 show the results from the first 23 subjects. Both regression lines are given in Fig. 2 and ATPS values are given for both instruments. In Fig. 3 the readings are converted to BTPS and the scattergram shown with the line of identity. The mean difference in FEV_1 for these 23 subjects was 0.162 ± 0.08 L BTPS, and the difference was significant at the 0.1% level ($t = 9.6$). The difference in COV was 1.42% and this difference was not statistically significant at the 5% level.

Figs. 4 and 5 deal with the results from the next 14 subjects namely Nos. 24 to 37. For these the mean difference in FEV_1 was 0.365 ± 0.34 L BTPS and the difference was statistically significant at the 0.1% level ($t = 11.07$). Again the difference in COV was small and insignificant (0.98%, $t = 0.92$). The scattergram for these 14 subjects consisting of 9 males and 5 females is given in Fig. 4, and Fig. 5 shows the vitalograph results plotted against the calibrated volumes on the flow volume apparatus.

COMPARATIVE MEASUREMENTS OF THE FORCED EXPIRATORY VOLUME IN ONE
SECOND (FEV₁) ON A BELLOWS SPIROMETER (VITALOGRAPH) AND THE FLOW
VOLUME APPARATUS FITTED WITH A ONE SECOND MARKER.

SUBJECT	FLOW VOLUME	SPIROMETER	DIFFERENCE (FLOW VOLUME - SPIROMETER)
NUMBER	MEAN FEV ₁	MEAN FEV ₁	
1	3.69	3.51	0.18
2	3.22	3.30	-0.08
3	3.46	3.78	-0.32
4	3.55	3.71	-0.19
5	4.48	4.52	-0.04
6	4.63	4.60	0.03
7	5.69	5.69	0.00
8	2.70	3.00	-0.30
9	2.51	2.55	-0.04
10	4.00	4.08	-0.08
11	3.47	3.21	0.26
12	4.02	4.37	-0.35
13	4.51	5.00	-0.49
14	4.91	5.03	-0.09
15	1.56	1.47	0.09
16	3.27	4.12	-0.85
17	3.98	4.06	-0.08
18	2.70	2.48	0.22
19	2.46	2.57	-0.11
20	4.83	5.21	-0.38
21	3.81	3.93	-0.12
22	4.56	4.67	-0.11
23	4.38	5.21	-0.83
24	2.41	2.65	-0.24
25	2.46	1.84	0.62

26	2.53	3.00	-0.47
27	2.69	3.00	-0.31
28	2.97	3.41	-0.44
29	3.58	3.99	-0.41
30	4.18	5.00	-0.82
31	2.96	3.29	-0.33
32	3.59	4.35	-0.76
33	2.31	2.40	-0.09
34	4.02	4.55	-0.52
35	3.49	4.05	-0.56
36	3.61	3.97	-0.36
37	3.10	3.52	-0.12
38	4.68	4.55	0.13
39	4.53	4.49	0.04
40	3.60	3.71	-0.11
41	3.49	3.75	-0.26
42	3.69	3.48	0.21
43	3.26	3.27	-0.01
44	2.60	2.89	-0.29
45	2.94	2.84	0.10
46	2.26	2.37	-0.11
47	2.95	3.12	-0.17
48	3.64	3.76	-0.12
49	4.86	4.84	0.02
50	3.83	3.88	-0.05
51	3.40	3.34	0.06
52	4.13	3.95	0.18
53	4.08	4.54	-0.46
54	5.67	5.55	0.12
55	3.05	3.02	0.03

56	2.71	2.92	-0.21
57	3.98	4.16	-0.18
58	2.06	1.93	0.08
59	2.54	2.57	-0.03
60	2.61	2.54	0.07
61	3.01	2.74	0.27
62	2.33	2.32	0.01
63	2.91	2.97	-0.06
64	3.23	2.82	0.41
65	2.54	2.79	-0.25
66	3.36	3.18	0.18
67	4.42	4.12	0.30
68	2.05	2.17	-0.12
69	4.12	3.95	0.17
70	2.70	3.03	-0.33
71	2.83	2.79	0.04
72	4.92	5.15	-0.23
73	2.50	2.54	-0.04
74	4.11	4.12	-0.01
75	2.73	2.47	0.26
76	3.58	3.76	-0.18
77	2.04	1.50	0.99
78	4.14	4.11	0.03
79	4.44	5.18	-0.74
80	4.61	4.62	-0.01
81	3.90	3.94	-0.04
82	4.04	4.00	0.04
83	3.26	4.07	-0.87
84	1.57	1.42	0.15
85	4.62	4.97	-0.35

86	2.66	3.72	-0.06
87	1.38	4.69	-0.31
88	1.91	4.67	0.24
89	5.34	4.99	0.35
90	3.08	3.26	-0.18
91	3.76	3.52	0.24
92	3.47	3.37	0.10
93	2.73	2.58	0.15
94	2.83	2.91	-0.08
95	2.59	2.64	-0.05
96	2.73	2.59	0.14
97	2.44	2.46	-0.02
98	1.59	1.94	-0.35
99	3.36	3.20	0.16
100	3.36	3.20	0.16
101	4.06	4.04	0.02
102	2.54	2.52	0.02
103	3.60	3.38	0.22
104	2.73	2.96	-0.23
105	5.68	5.63	0.05
Mean	3.49 (0.91)	3.60 (0.98)	-0.106 (0.316)
SE(M)	0.09	0.10	0.031
t	-	-	3.44
P	-	-	<0.01
COV	26	27	-

Vitalograph $FEV_1 = 1.02$ Flow volume $FEV_1 + 0.04$
 $r = 0.95$; $t = 30.14$; $P < 0.01$

FIG.1 Regression line for converting FEV_1 (L,BTPS)
on Flow Volume apparatus to FEV_1 ()
on Vitalograph spirometer.

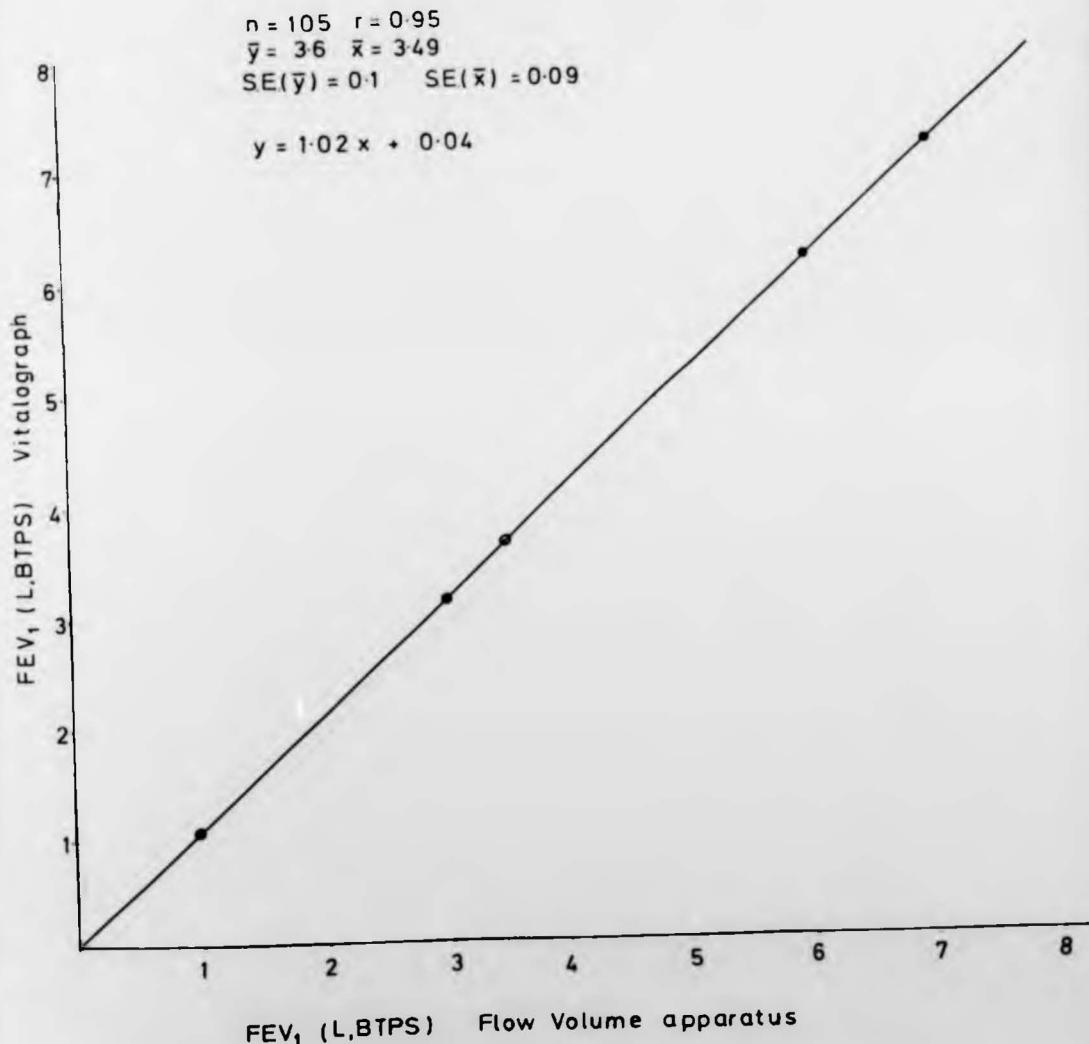


FIG.2 Regression lines of FEV₁ measured on
Flow Volume apparatus & Vitalograph for
23 subjects.

solid line - $y = 0.84x + 0.27$, SD(y) = 1.02
dashed .. $x = 1.09y - 0.02$, SD(x) = 0.89

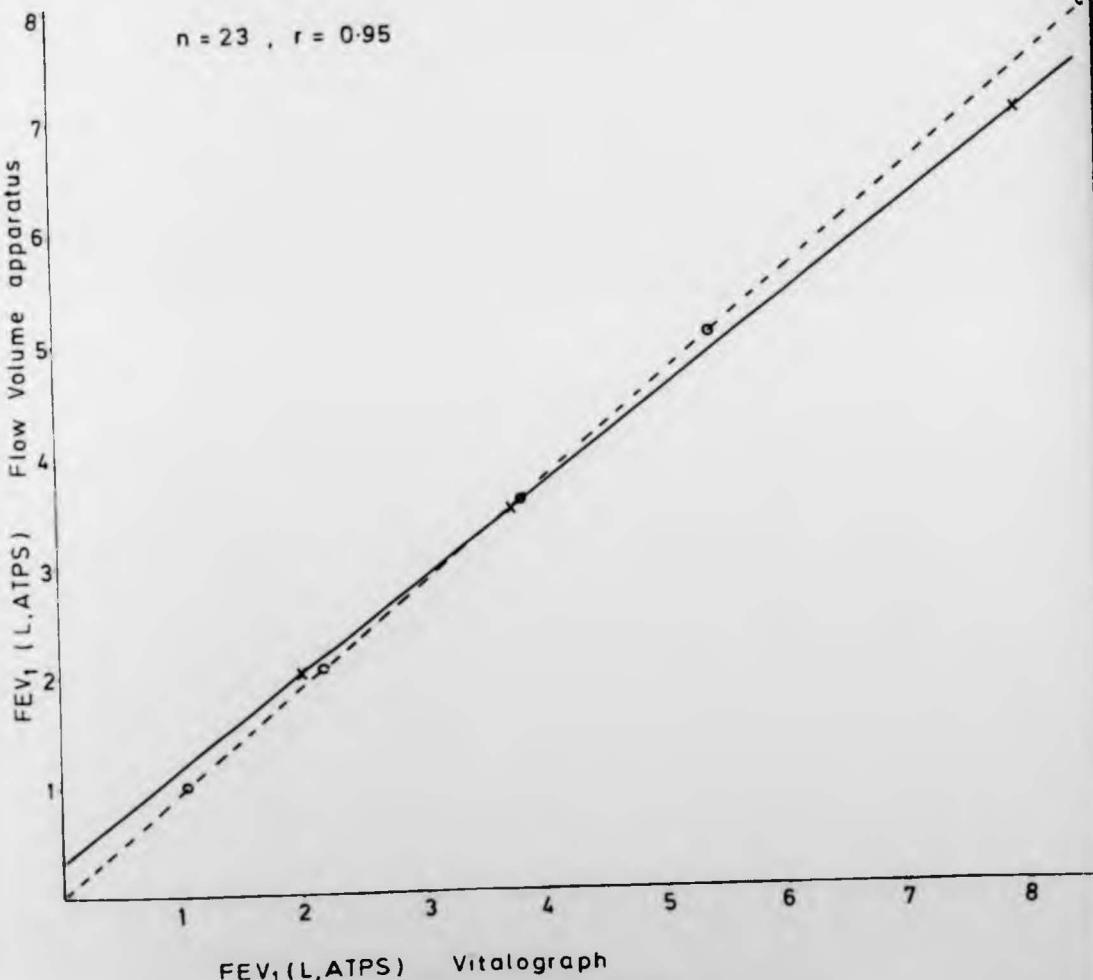


FIG 3 Comparison of FEV₁ in 23 subjects measured on
Flow Volume apparatus & bellows spirometer

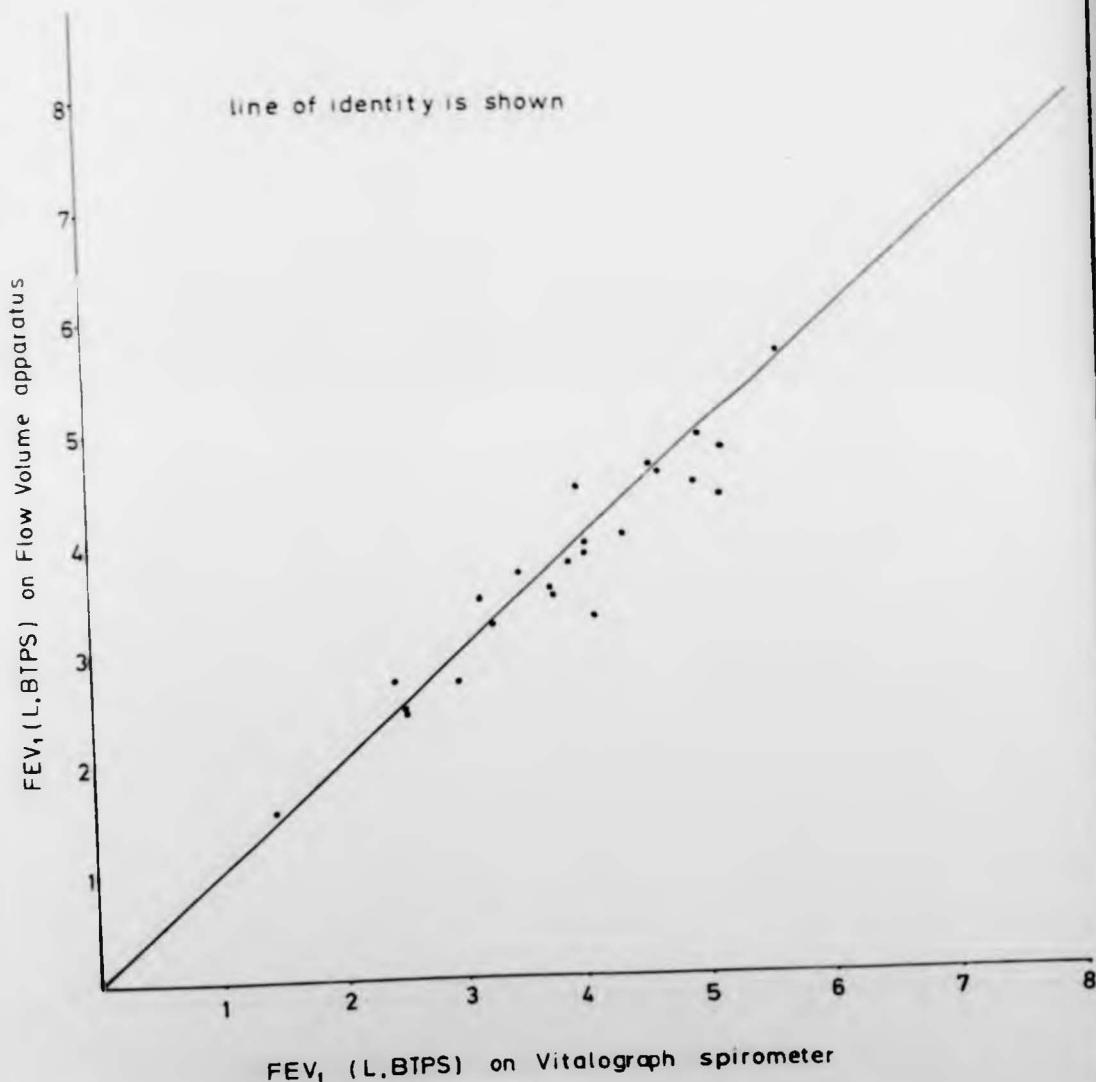
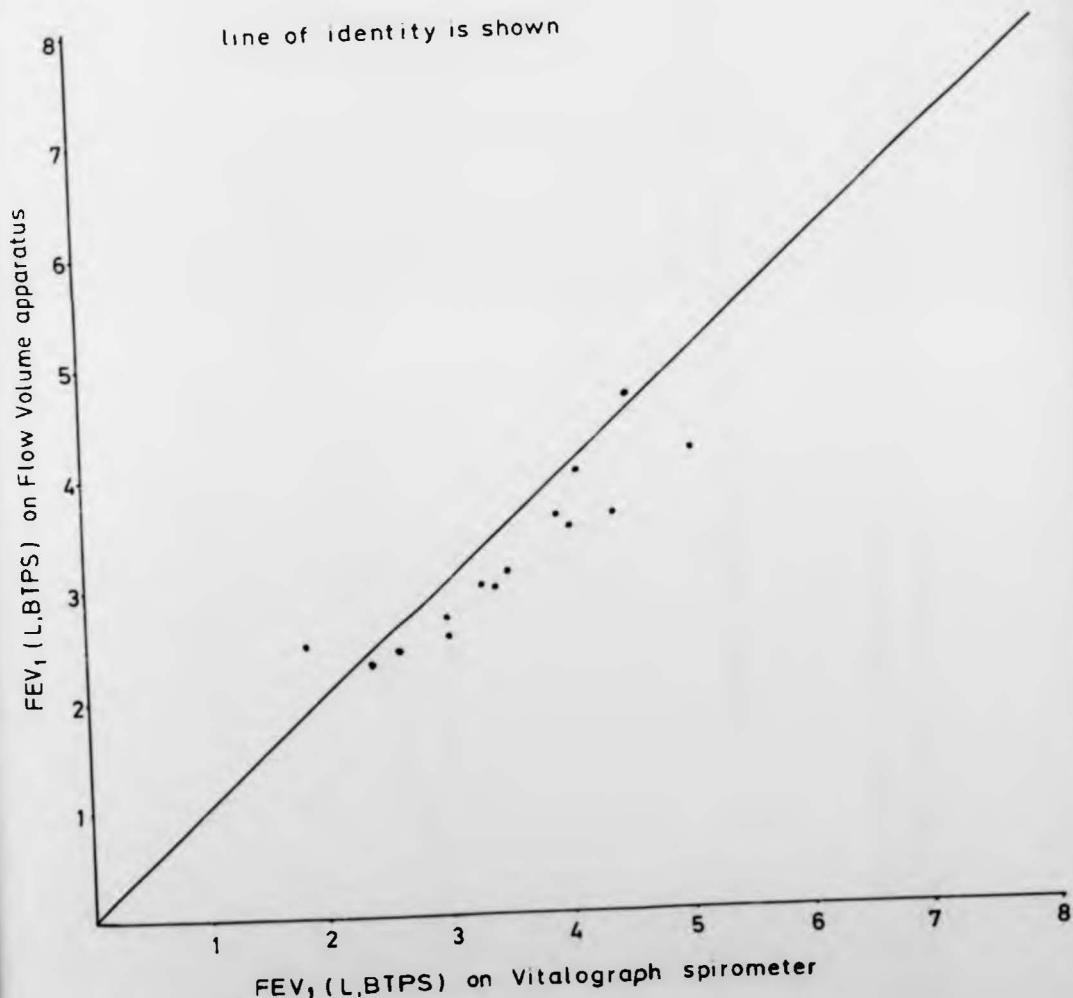


FIG.4 Comparison of FEV₁ in 14 subjects (nos. 24-37)
measured on Flow Volume apparatus & bellows spirometer.



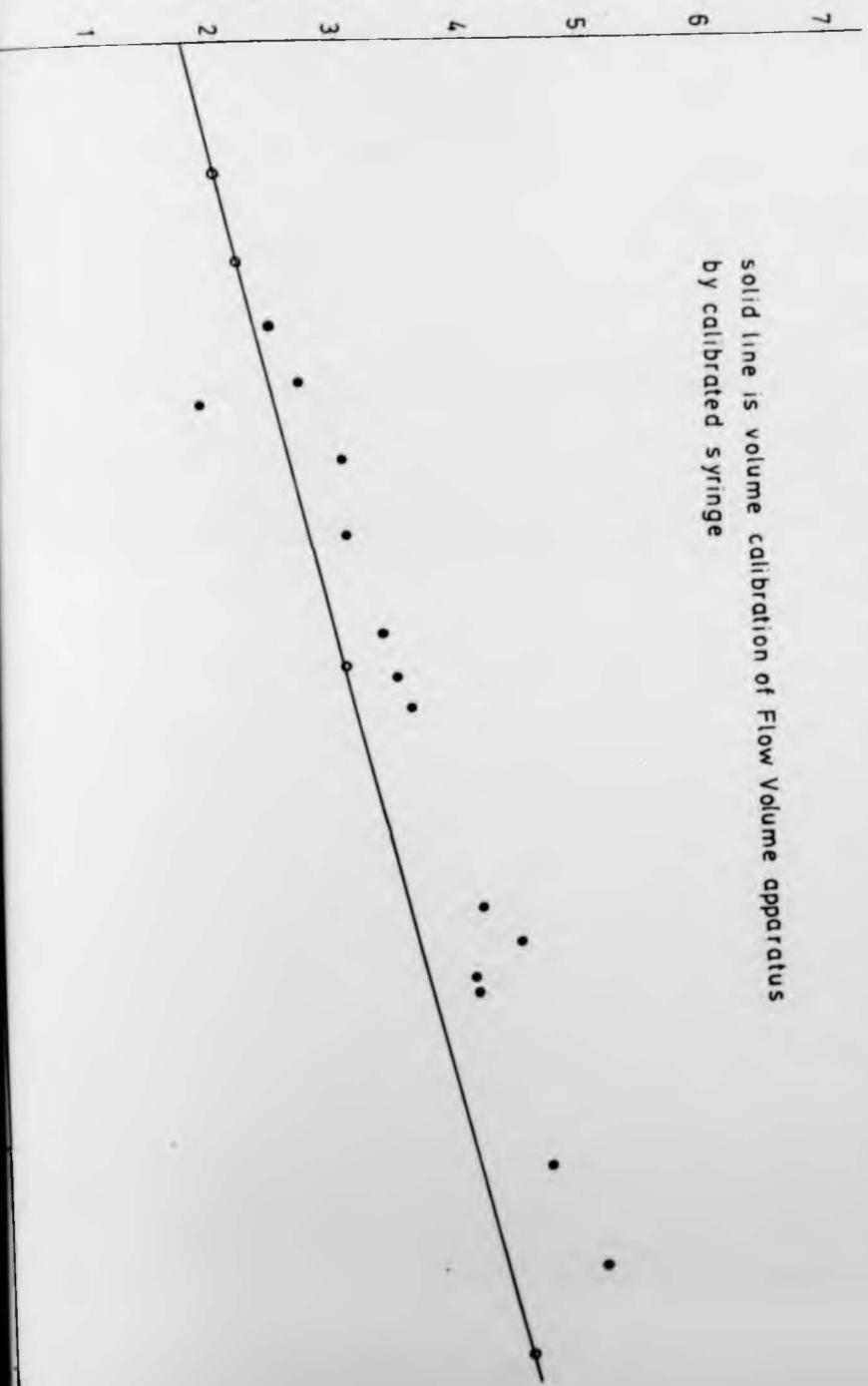
FEV₁ (L,BTPS) Vitalograph

FIG 5 FEV₁, measured on Vitalograph Vs. deflection (cm.) on Flow Volume apparatus in 14 subjects (9 males & 5 females), over temperature range 23°C – 27°C

solid line is volume calibration of Flow Volume apparatus
by calibrated syringe

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AN INVESTIGATION OF THE CHARACTERISTIC "NOTCH" ON FLOW VOLUME CURVES
IN HEALTHY PERSONS

Introduction

In forced expiration the flowrate is determined by at least two different mechanisms; the first portion of the flow volume curve being effort dependent and the second portion largely independent of effort (Try and Hyatt, 1960; Dayman, 1967; Bouhuys, 1974). On flow volume curves generated outside a body plethysmograph, a characteristic "notch" separates the two portions of the curve (Tammeling, 1966; Clarke, 1969). The suggestion that the notch is due to an equipment artefact (Tammeling, 1966) has been disproved by Clarke (1969). The last named author investigated the behaviour of the notch in healthy persons and patients comparing the results qualitatively. The present study was carried out to extend the analysis of the notch's characteristic in apparently healthy persons who may nevertheless have overt small airway defect.

Subjects and Methods

The apparatus (Mk. 1 flow volume) has been previously described (Chapter 1.). The subjects were all medical practitioners attending a course in respiration physiology in this department. Of the 28 volunteers, two were Asians and the rest Caucasians. Ten of the 22 males were smokers (MRC questionnaire on respiratory symptoms, 1966). Only one of the females smoked. A shortened questionnaire was used to determine the subjects' smoking category and to establish whether a history of respiratory malfunction existed.

Three basic variables were measured: peak flow (PEFR), forced vital capacity (FVC) and the instantaneous flow rate after two thirds of the vital capacity was expelled (MEF₅₀).

The notch was closely examined and the instantaneous flowrate at the notch (V_N or MEF(N)) together with the position on the volume axis below TLC (V_N) noted.

Results

The volume V_N was standardised by expressing it as a fraction (V_N/FVC) of the vital capacity. V_N was similarly standardised as the fractions MEF(N)/PEFR and MEF33/MEF(N). The observed FVC was also expressed as a percentage of predicted value (Cotes, 1975). Only one subject (No. 1), a smoker, had an observed FVC below 65% of predicted. Table 1 gives details of the physical characteristics, smoking category and the measured and calculated pulmonary function indices. For males, the mean height and age were similar in smokers and non-smokers, and the mean values of the various indices were not statistically different in either group. The females were on average 17 cm shorter and one year older than the males. The pulmonary function indices were generally lower in the females: this is in keeping with the known sexual dependence of these indices.

The notch was located at a mean volume which was 2% (below TLC) of the observed vital capacity. (28% in males and 31% in females). The mean instantaneous flowrate at the notch was 57% PEFR (55% in males, 62% in females). MEF33 was on average 36% of the flowrate at the notch and was again slightly higher in females (61%) compared to males (55%). Neither the position (FVC below TLC) nor the height of the curve at the notch (MEF(N)/PEFR) was correlated to the physical variables height and age thus making internal standardisation of these variables unnecessary. Fig. 1 is the scattergram for notch position and age and Fig. 2 that for notch height and PEFR. The weak negative correlation of notch height and PEFR accounts for only about 8% of the total

variability in the data.

Notch position was negatively correlated with observed FVC (Fig. 3) and also with FVC expressed as% predicted (Fig. 4), the correlation was significant at the 1% probability level. Statistically significant positive correlations were found for the notch height with PEFR (Fig. 5) and MEF33 (Fig. 6).

Table 2 summarised the results of regression analysis for all the subjects grouped as well as for male smokers and male non-smokers separately. As expected MEF33 and FVC gave negative age coefficients in both smokers and non-smokers. Fig. 7 is a scattergram relating the instantaneous flowrate at the notch to flows in both the dependent and independent portions of the flow volume curve and illustrates that 8 out of the 11 smokers (72.7%) but only 2 of the 17 non-smokers had notch flowrates which were more than 60% of the PEFR.

Discussion

The possibility that the notch phenomenon was no more than an equipment artefact was considered. Clearly the ability of the output device to handle high frequency transients can influence the shape of the MEFV curve traced. Fig. 8 shows the output traced on an oscilloscope: the "light" pen (electron beam) is not subject to mechanical drag or inertia. Fig. 9 (a) and (b) shows the curve when traced by mechanical XY plotters. In (a) an MEFV was recorded on an FM magnetic tape recorder (Racal Store 4) at 76.2 cm sec^{-1} and subsequently played back onto a Bryans XY plotter (Model 2400) at 76.2 cm sec^{-1} and 38.1 cm sec^{-1} . At 38.1 cm sec^{-1} the plotted curve resembles that obtained on the oscilloscope. At the higher speed, the plotter is unable to follow the curve and distorts it. In (b) the three MEFVs were performed by

the same subject as in (a); in this case the XY plotter (Hewlett Packard model 7047A) was operated with input filters in various signal paths. It is seen that when the transients are filtered out (filters in both the X and Y signal paths), the tracing is smoothest but the maximum height of the curve is the lowest. The effect is therefore to reduce PEFR and to "smooth" out the notch. Figs. 8 and 9 are copies of actual traces and the axes were scaled such that the Y amplification was twice that of X. The effect of the distortion of the output device appears to be confined to the upper two thirds of the vital capacity. It therefore seems unlikely that the equipment produces the notch; rather the evidence points to equipment being used in such a manner as to obscure the notch. This finding is in agreement with that of Clarke (1969).

If physiological factors determine notch formation then physical or environmental factors would be expected to influence it in some way. The position of the notch has indeed been shown to vary in health and disease and also with the initial lung volume at which forced expiration begins (Clarke, 1969). The finding in the present study that in the "less healthy" (lower % predicted FVC) the notch occurred later in expiration also agrees with Clarke's finding. It is emphasized that these subjects were healthy and that, other than notch position, even the smokers showed little evidence of ventilatory deficiency. However it is seen (Fig. 7) that a larger measure of separation between smokers and non-smokers can be achieved using notch characteristic. Since the separation is greater on the axis representing events in the effort dependent zone of the MEFV, (MEF(5)/PEFR), the respiratory muscles could be involved. The observation that flowrate of the notch has a significant positive correlation with both PEFR and MEF35 (Table 2),

suggests dynamic compression of airways as a possible mechanism of notch formation. If this is so, the position of the notch in health and disease may be explained on Mead's equal pressure point (EPP) theory; the notch being formed when the EPP becomes trapped in a compressible airway segment.

It is concluded that the notch is unlikely to be an artefact, but rather than its formation has a physiological basis. It is also possible that notch characteristic may be useful in early detection of small airway abnormality. This latter point is, on these observations, only speculative and needs to be confirmed by studying a large number of subjects.

TABLE 1. PHYSICAL CHARACTERISTICS AND PULMONARY FUNCTION INDICES OF 28 HEALTHY ADULTS

TABLE 1. PHYSICAL CHARACTERISTICS												
Sub.No.	Age (Yr)	Ht (m)	S* or NS	FVC (L)	PEFR L/min ⁻¹	MIF33 L/min ⁻¹	V _N (L)	V/FVC %	MIF(H) L/min ⁻¹	MIF(H) PLFR ⁻¹	MIF33/HIF(H) %	% Pred FVC
Sex												
1	M	31	1.78	S	3.25	72	99	1.53	67	292	62	34 65
2	M	30	1.80	S	6.17	75	56	1.48	24	266	36	21 121
3	M	27	1.78	S	6.59	828	179	1.58	24	673	82	26 130
4	M	23	1.82	S	6.65	65	152	1.09	16	483	75	32 127
5	M	34	1.75	S	5.83	57	12	1.76	30	392	68	36 123
6	M	31	1.81	S	5.79	86	191	1.80	31	506	60	38 113
7	M	29	1.67	S	6.92	561	12	1.81	26	259	16	40 156
8	M	38	1.76	S	6.07	687	110	1.27	21	337	49	33 129
9	M	28	1.88	S	5.93	855	179	1.81	30	518	61	35 108
10	M	36	1.83	S	6.27	723	139	1.90	30	115	62	31 116
MEAN		31.2	1.79	S	5.95	693	139	1.60	27.9	17.6	60.1	32.6 119
SD		3.7	0.06		1.02	131	51	0.27	8.2	13.1	13.6	5.6 23
11	M	26	1.78	NS	5.20	715	172	1.74	35	379	53	45 102
12	M	32	1.72	NS	6.04	570	190	1.18	20	449	79	42 130
13	M	27	1.74	NS	5.43	638	129	1.11	21	379	59	34 112

14	M	37	1.80	NS	5.57	725	132	1.63	29	119	58	32	113
15	E	37	1.67	NS	3.44 ^a	568	72	1.26	37	232	41	31	93
16	M	40	1.95	NS	5.75	941	112	2.00	35	291	31	39	102
17	E	27	1.79	NS	5.89	619	129	1.13	19	360	53	36	115
18	E	26	1.83	NS	6.20	708	142	1.63	26	336	49	37	116
19	M	29	1.73	NS	6.01	856	110	1.85	31	317	37	35	126
20	M	27	1.85	NS	6.15	577	80	1.04	17	291	50	28	113
21	M	31	1.72	NS	3.90	658	110	1.36	35	214	48	35	87
22	M	37	1.73	NS	4.98	723	155	1.49	30	389	53	40	109
MEAN		31.3	1.73	NS	5.38	699	128	1.45	27.8	350.5	51.3	36.2	110
SD		5.1	0.08		0.89	118	35	0.32	7.0	62.1	12.3	4.8	12
MEAN		31.3	1.73	MALES	5.64 ^b	696	133	1.52	37.8	311.0	55.3	34.5	114
SD		4.4	0.07		0.97	121	37	0.30	7.1	104.4	13.4	5.4	18

23	F	27	1.65	S	3.9 ^b	562	126	1.03	26	359	63	35	106
24	F	35	1.5 ^b	NS	2.83	256	89	1.45	50	136	53	65	95
25	F	38	1.66	NS	1.31	432	106	1.16	27	359	83	30	120
26	F	26	1.62	NS	3.53	437	103	1.00	28	263	56	43	100
27	F	34	1.55	NS	5.11	691	129	1.30	26	102	58	32	165
28	F	35	1.65	NS	3.53	264	129	1.56	15	321	59	40	98
MEAN		32.5	1.61	FE	3.89	407	114	1.25	33.7	303	62	40.8	114
SD		6.8	0.05	MALE	0.76	149	16.8	0.23	10.9	97.9	10.8	12.8	27
MEAN		31.5	1.75	ALL	5.27	652	129	1.46	29.1	301	56.8	35.9	114
SD		6.5	0.10		1.17	152	34.6	0.30	8.4	106.3	13.0	7.7	20

TABLE 2 .

Regression analysis(a) Multiple regression in male smokers and non-smokers

	Const	Age coefft.	Height coefft.	Number
MEF33	30.31	- 3.03	112.40	10 smokers
	202.5	- 1.73	- 11.76	12 non-smokers
FVC	8.93	- 0.05	- 0.82	10 smokers
	- 3.01	- 0.32	6.14	12 non-smokers
MEF(N)	- 1030.4	- 13.16	1036.6	10 smokers
	495.5	- 3.48	- 21.4	12 non-smokers
PEFR	- 1238.0	- 7.73	1212.0	10 smokers
	- 1047.0	4.16	536.2	12 non-smokers
V _N	0.09	0.003	0.35	10 smokers
	- 2.05	0.02	1.65	12 non-smokers

(b) Simple regressions

All subjects:- n = 23

(a) $V_N = 0.078 FVC + 1.05 \quad r = 0.701 \quad P > 0.05, COV_x = 22.2\%$
 $COV_y = 20.8\%$

(b) $MEF(N) = 0.387 PEFR + 111.7 \quad r = 0.554 \quad P < 0.01, COV_x = 23.3\%$
 $COV_y = 29.2\%$

(c) $MEF(II) = 2.42 MEF33 + 52.6 \quad r = 0.706 \quad P < 0.01, COV_x = 26.8\%$
 $COV_y = 29.2\%$

(d) $V_N/FVC\% = -0.34 FVC + 51.5 \quad r = -0.640 \quad P < 0.01, COV_x = 20.6\%$
 $COV_y = 26.3\%$

(e) $\frac{MEF(N)}{PEFR}\% = -0.118 \frac{MEF33}{MEF(II)} + 61.0 \quad r = -0.07 \quad P > 0.05, COV_x = 21.5\%$

$COV_y = 22.9\%$

(f) $\frac{MEF(N)}{PEFR}\% = -0.024 PEFR + 72.3 \quad r = -0.279 \quad P > 0.05, COV_x = 23.4\%$

$COV_y = 22.9\%$

(g) $V_N/FVC = -0.265 FVC + 59.3 \quad r = -0.62 \quad P < 0.01, COV_x = 26.6\%$
 $COV_y = 26.3\%$

(h) $V_N/FVC = -24.06 \text{ Height} + 71.1 \quad r = -0.277 \quad P > 0.05, COV_x = 5.5\%$

$COV_y = 26.3\%$

FIG. 1

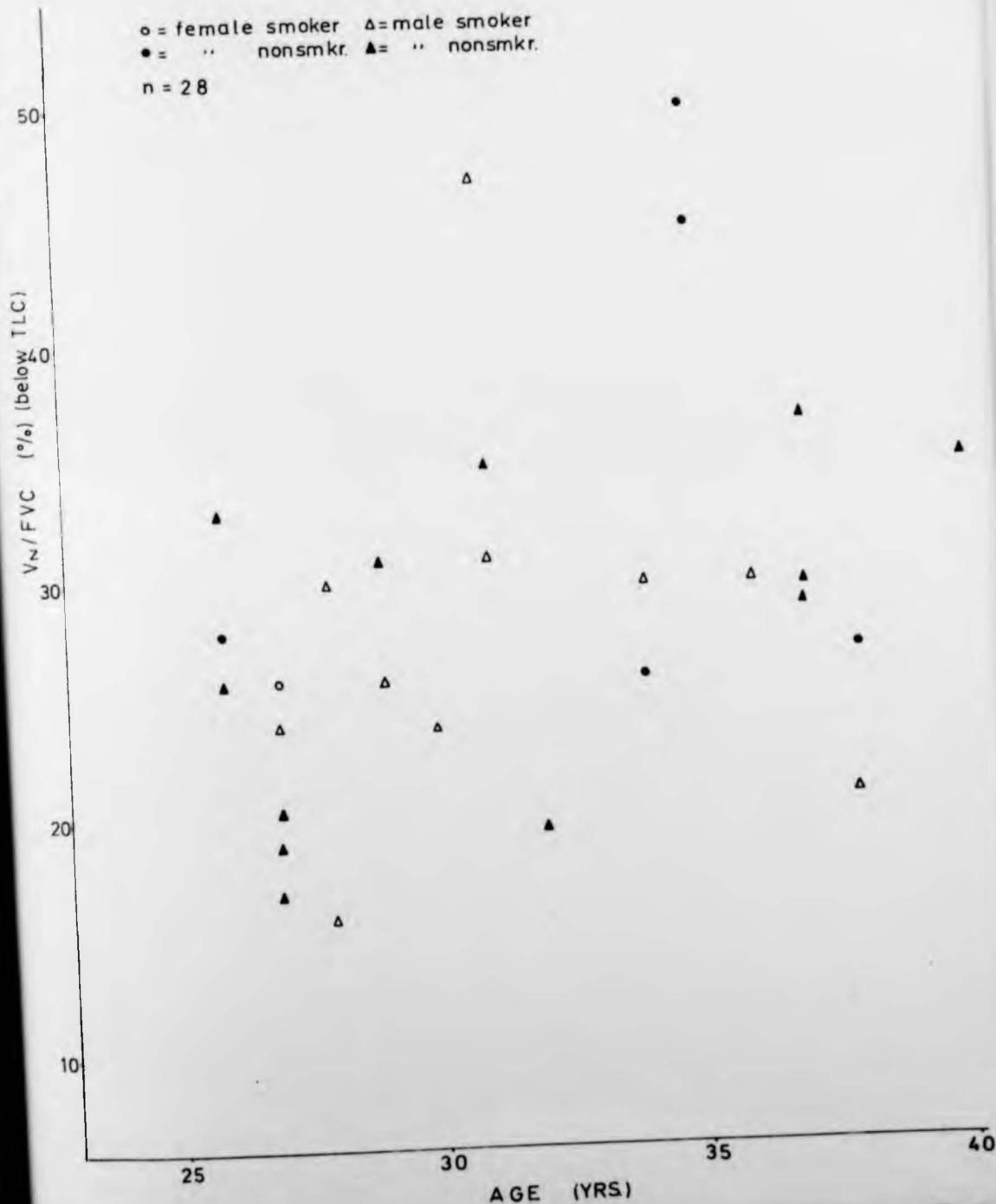


FIG. 2

Ch. 4.

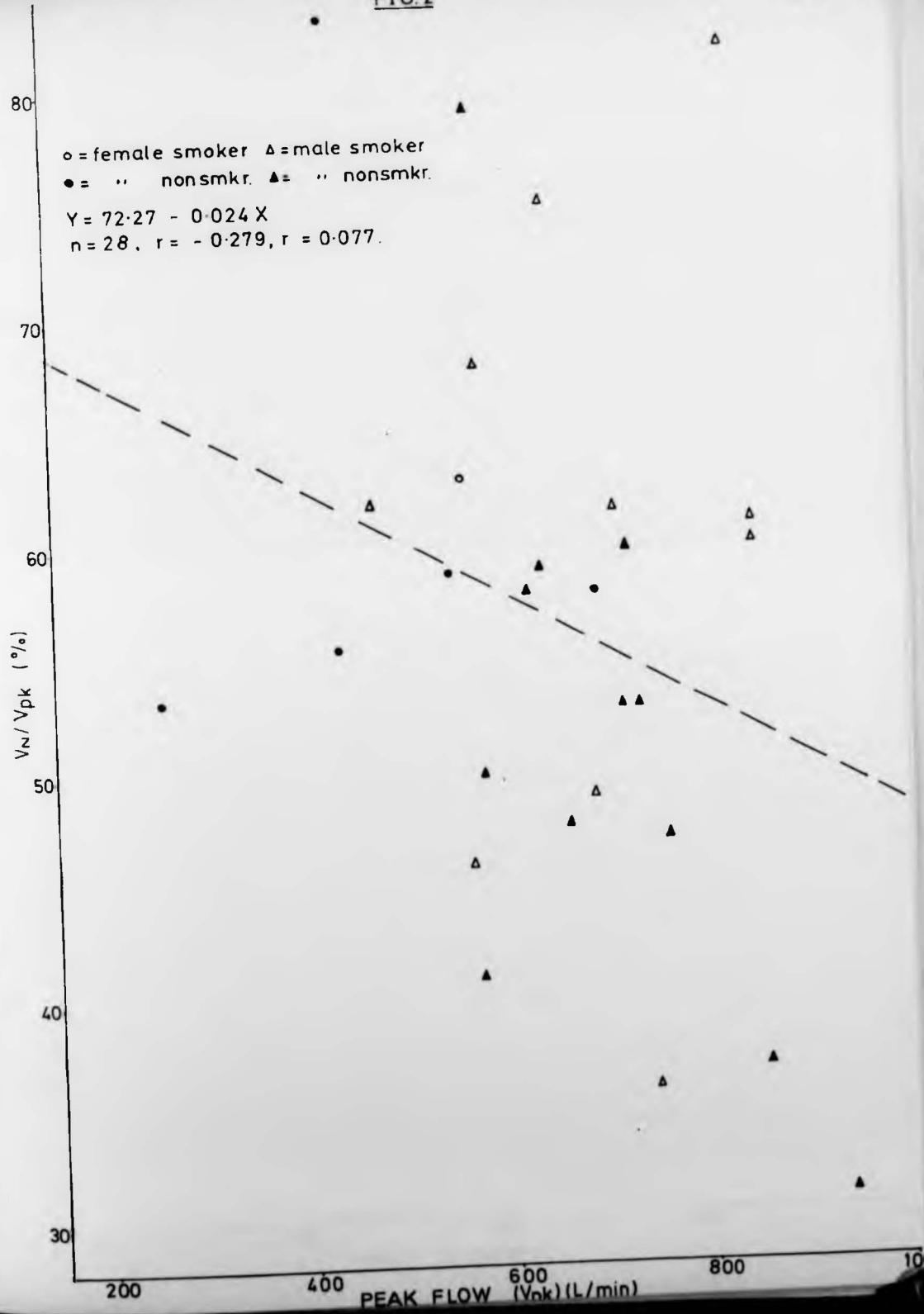


FIG. 3

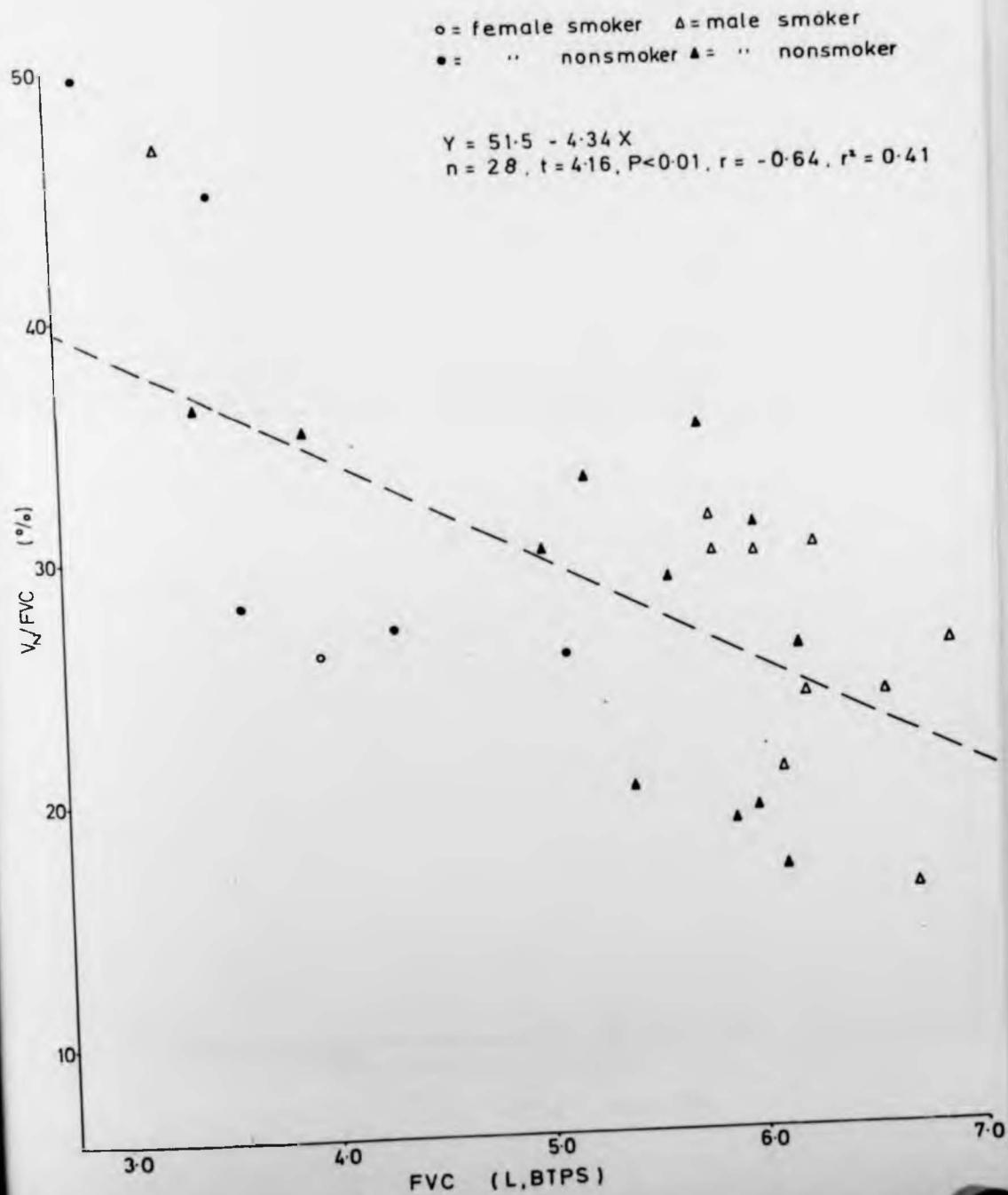


FIG. 4

○ = female smoker △ = male smoker
 ● = " ▲ = nonsmoker ▲ = .. nonsmoker

$$Y = 59.26 + 0.265X$$

$$n = 28, t = 4.03, P < 0.01, r = -0.62, r^2 = 0.384$$

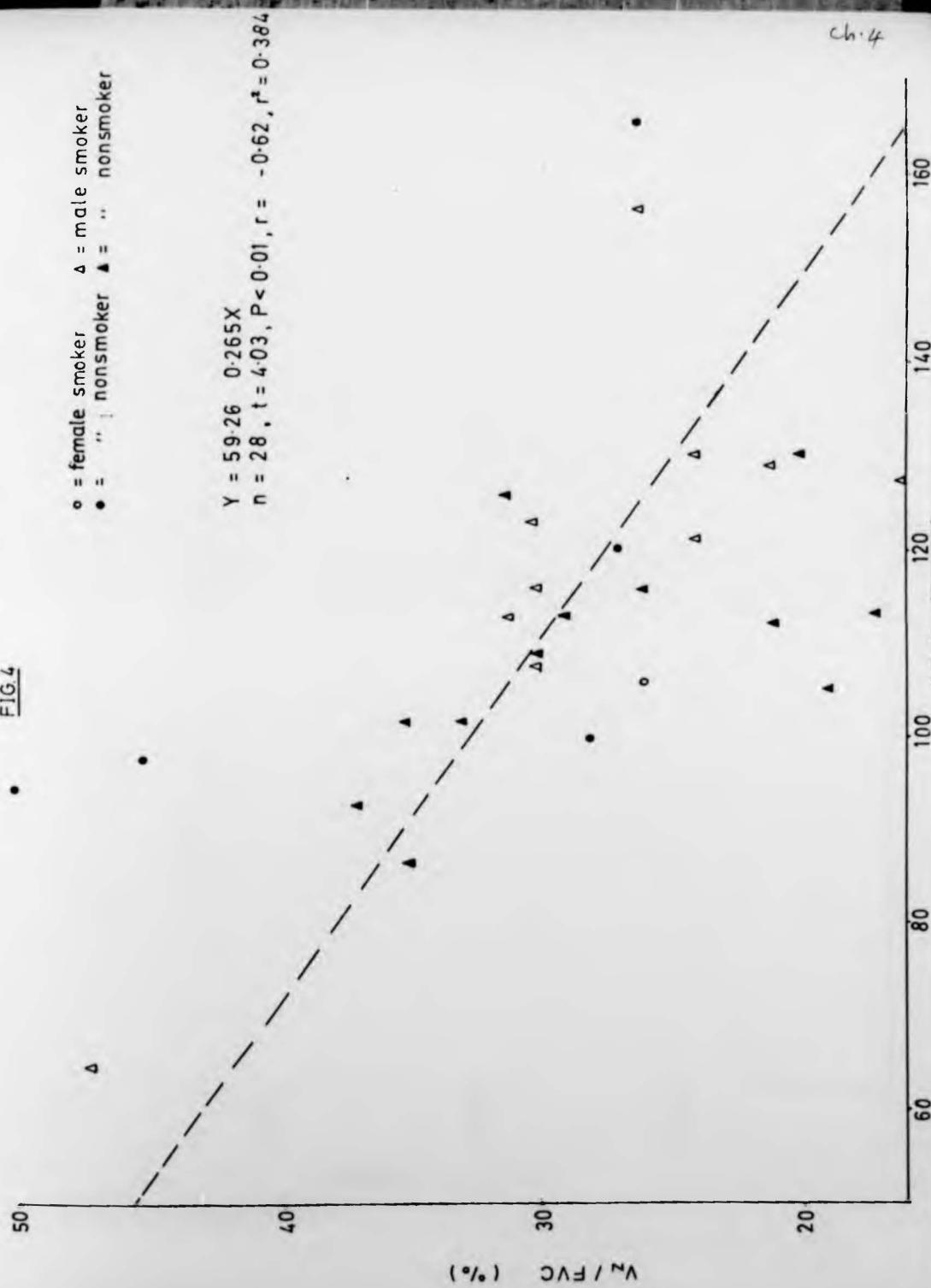


FIG. 5

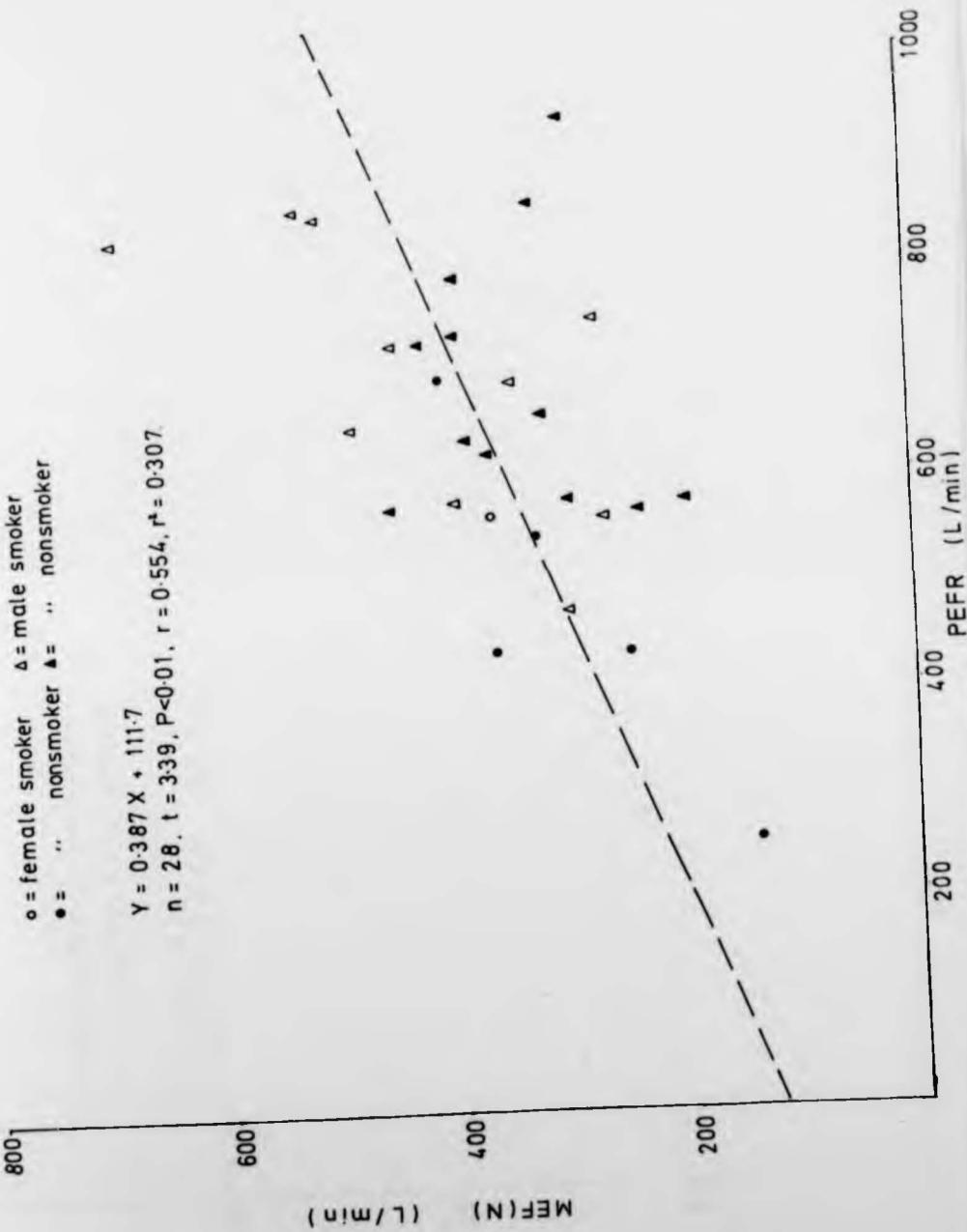


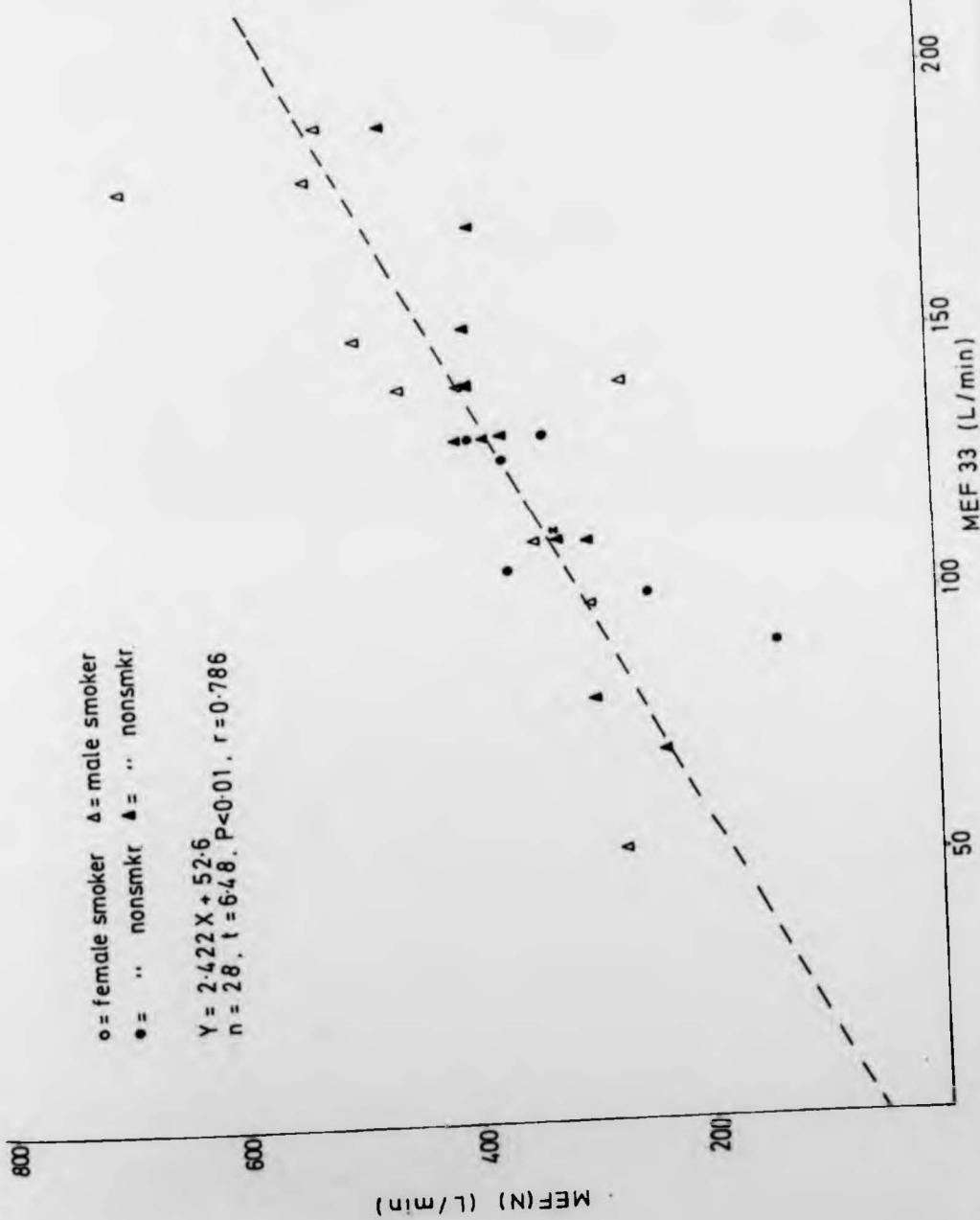
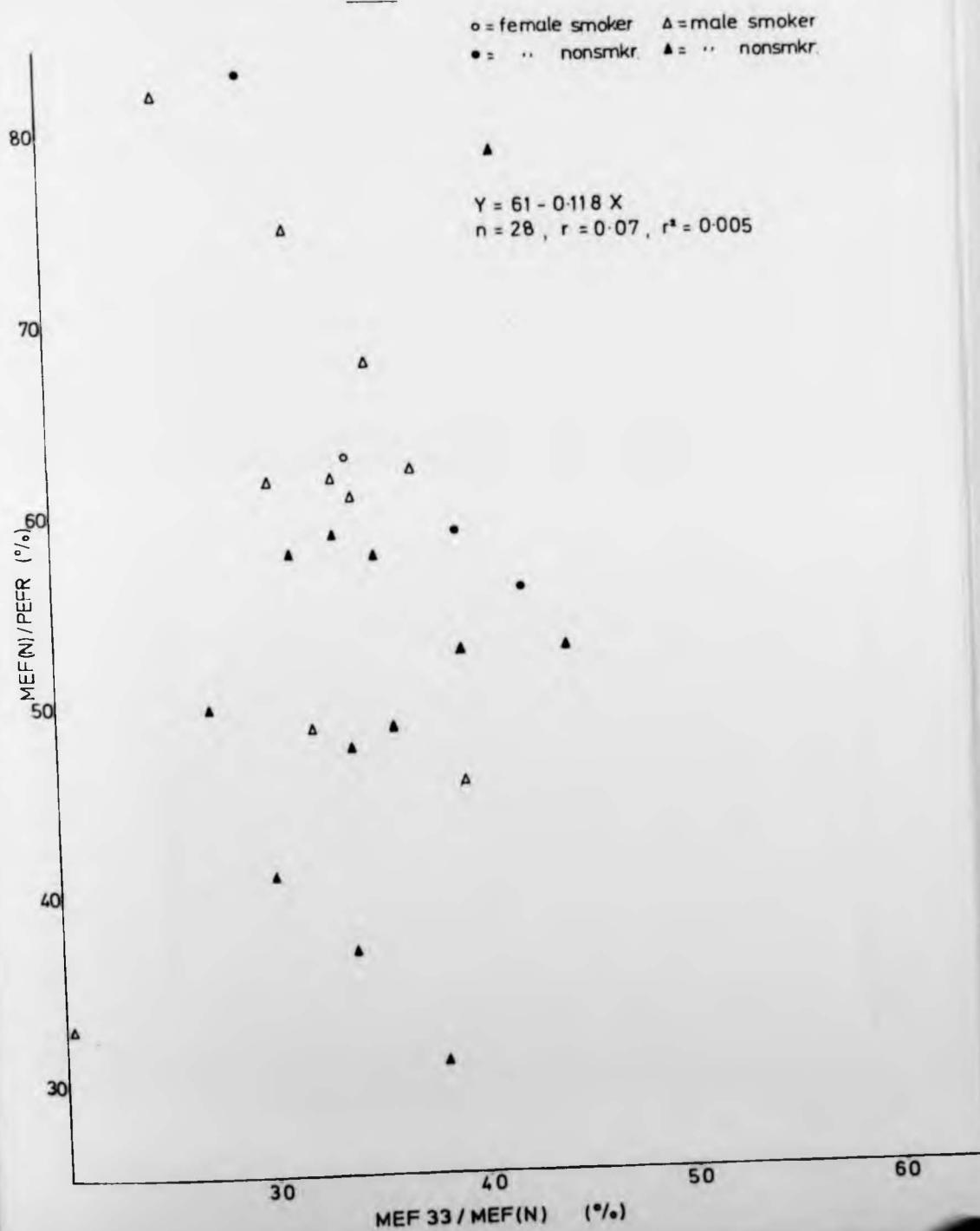
FIG. 6

FIG. 7



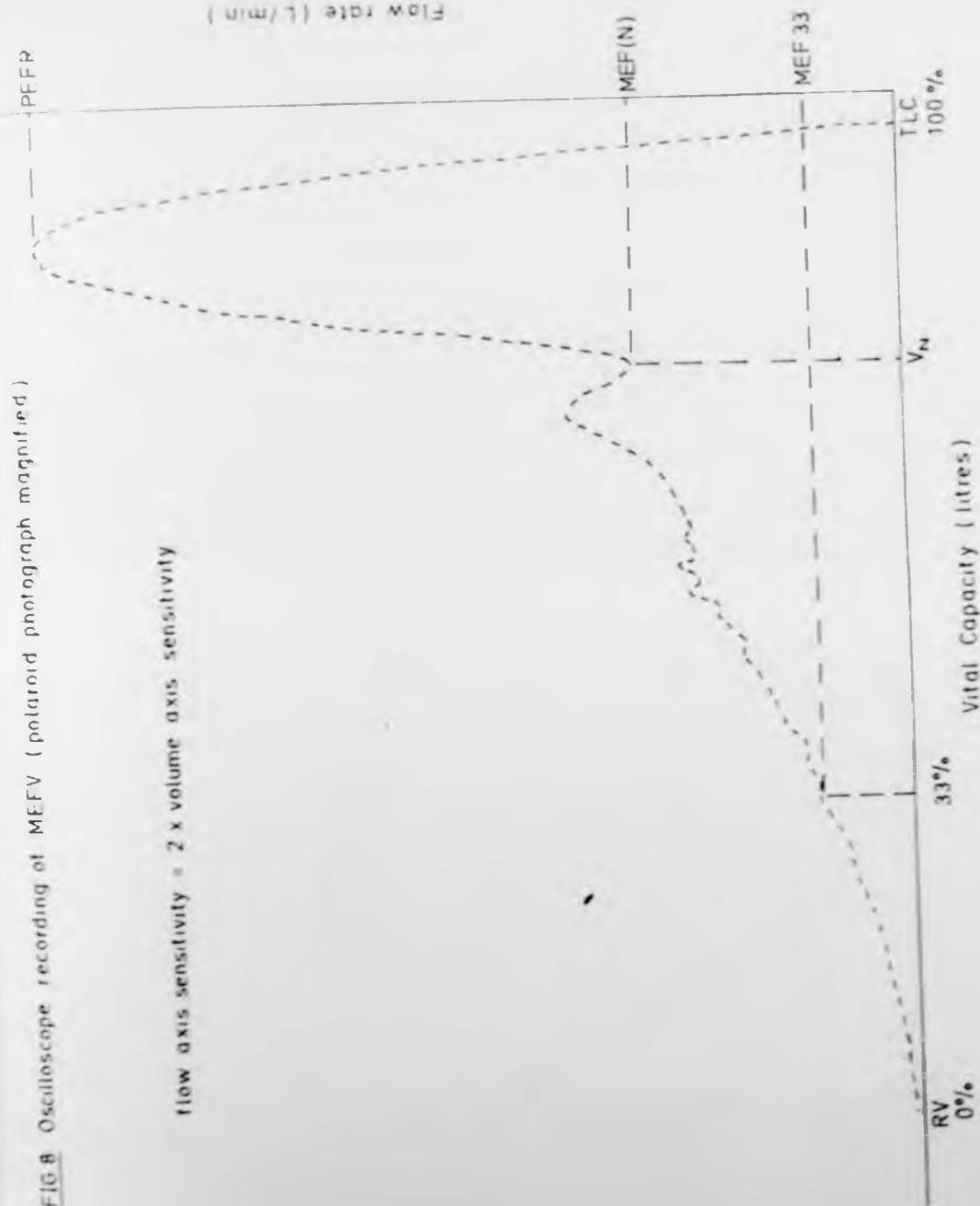
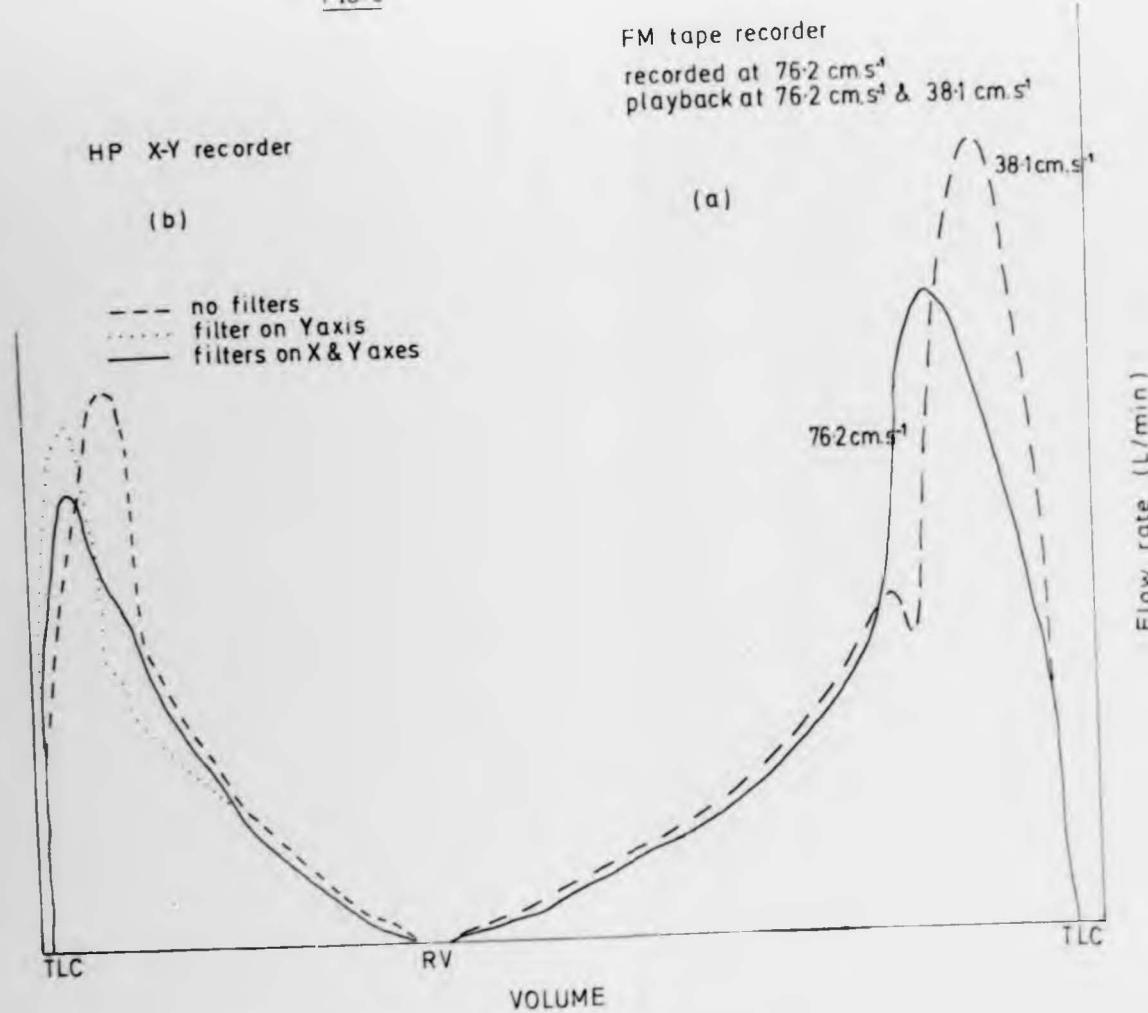


FIG 9



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THE RELATIONSHIP BETWEEN PARTIAL AND MAXIMAL EXPIRATORY FLOWRATES
IN HEALTHY SUBJECTS

Introduction

The expiratory resistance of lung airways changes with lung volume, being lower in the full (TLC) than in the deflated lung (RV). Expiratory flowrates are highest close to TLC where they depend on effort and the large airway resistance and become zero at RV. Below a certain lung volume the height of the flow volume curve is independent of effort but depends inter alia on the smooth muscle tone and also on the elastic retraction of the small airways which at low volumes become flow limiting.

When flow volume curves are generated outside a body plethysmograph an absolute lung volume scale is not available on the volume axis. If vital capacity changes, it is not known whether TLC, RV or both have moved relative to an absolute lung volume scale. If the flow volume curves are to be compared under these circumstances an arbitrary decision must be taken regarding the reference point. It has been stated (Comroe,

1955) that RV is a more stable reference than TLC, however, whilst flow volume curves are generally similar in shape near to TLC, they exhibit a variety of shapes (concave, convex and linear with abrupt cut off) as RV is approached.

In this study maximal (MEFV) and partial (PEFV) curves were aligned at TLC and the instantaneous flowrates were compared at 50% (MEF₅₀) and 75% (MEF₂₅) of the vital capacity.

Subjects and methods

The apparatus and technique have been described previously (Chapter 4.). The subjects were male medical practitioners undergoing a training course in anaesthetics. Their physical characteristics,

tobacco consumption and simple ventilatory indices are given in Table 1.

Results

The three smokers in the group had a maximum tobacco consumption of only 0.75 pack years (20 cigarettes per day for 1 year). These smokers revealed no evidence of overt chest disease either by questionnaire or in their observed FEV, FVC and PEFR.

The results for the whole group and for the non-smokers only were analysed separately. Regression equations between the partial and maximal flowrates were determined and the least square lines plotted together with the lines of identity on the scattergrams (Figs. 1,2,3 and 4).

The association between MEF50 and the physical variables height and age were tested by calculating simple and multiple regressions. The multiple regression relationship is given by the equation:-

$$\text{MEF50} = 217.7 + 0.036 \text{ Age} + 81.77 \text{ Height}$$

where age is in years and height in metres.

The association accounts for just over one half of one percent of the observed variation in MEF50 ($R^2 = 0.0057$) implying that the physical variables of these subjects did not contribute significantly to the observed variation. The simple correlation coefficients (r) for MEF50 and age and height were 0.464 and 0.108 respectively for the total group. The corresponding r values for the non-smokers were 0.413 and 0.261. Thus by including the smokers the total variability in MEF50 increased and this is as would be expected. None of the correlation coefficients reached the 5% level of significance, but this may be due in part to the small number of subjects.

On average in the non-smokers the MEF50(P) was higher than the MEF50 (Slope 1.43) but the MEF25(P) was lower than MEF25 (Slope 0.9).

The average coefficient of variation for MEF50(P) was 28.4% and for MEF25(P) 38.3%. As expected, these variations were larger than those for MEF50 (COV 17.2%) and MEF25 (COV 29.6%). The nearly constant difference of 10% to 12% between the COV of the flowrates at the higher lung volumes and those at lower lung volumes is perhaps indicative of systematic errors possibly related to the height of the ordinates on the flow volume curves at 50% and 75% of vital capacity. Since the curve is higher at 50% vital capacity (VC) than at 75% VC, reading errors in MEF25 will be a larger proportion of the true instantaneous value than similar errors in MEF50. Since the same reference point (TLC on the maximal curve) was used for both MEFV and PEFV indices variations in TLC would not contribute to the systematic errors being described here.

Discussion

It is known that some pulmonary function indices show a systematic difference between sex as well as between ethnic groups (Cotes, 1975). Only one sex (male) but three ethnic groups (Caucasians, Negroes and Asians) comprised the present volunteers. However, their predicted ventilatory indices (FEV_1/FVC and $FEV_1/FVC(\%)$) were calculated from prediction equations appropriate to each ethnic group. The observed pulmonary function of all the subjects (including the smokers) fell within the predicted normal range. If it is assumed that variations of airway smooth muscle tone are subject to the same fundamental factors in all the ethnic groups, then the unexpected positive correlation coefficient (r) of MEF50 and Age found in this study may be reflecting systematic ethnic differences in MEF50.

The underlying physiology of the airways is different when maximal and partial indices are measured. The MEFV curve is traced when tone in the airways is reduced or absent because the preceding inspiration to TLC has almost abolished airway smooth muscle tone (Vincent, 1970). By contrast, the inspiration preceding a PEFV curve should not abolish airway tone as it is terminated at about 60 - 70% TLC. For the PEFV curve the airways may therefore be influenced by the normal tonal state.

Airway smooth muscle is arranged differently according to airway generation. For those large airway generations which have cartilaginous support in the form of almost complete rings, the smooth muscle is arranged as longitudinal bands capable of stiffening the airway walls by pulling the rings closer to each other. This arrangement makes these airways less susceptible to dynamic compression during forced expiration. Where, as in the smallest airways, cartilage is scanty and occurs as irregular plates, the smooth muscle is arranged circumferentially. An increase in tone (shortening of the muscle fibres) will thus tend to decrease the bore of these more peripheral airways. In this study, MEF50(P) was higher than MEF50 and MEF25(P) lower than MEF25. These observations could be explained by tonal changes if it is assumed that at 50% of vital capacity the equal pressure point (EPP) resides in airways where there is full cartilaginous support. The upstream segment (whose resistance determines maximal flow) could be stiffened to resist collapse from dynamic compression by the increase in tone. Under these conditions, resistance to air flow would be less than when there was an absence of tone, MEF50(P) would therefore be larger than MEF50. By the time 75% of the VC was expelled, the EPP might well have moved peripherally to reside in the smaller airways.

An increase in tone in this region would be accompanied by an increase in upstream resistance and consequently a fall in maximal flowrates. Thus it would be predicted that MEF_{25(P)} should be less than MEF₂₅. The anatomical location of the EPP is not exactly known, but it is reasonable to assume that it would 'move' peripherally as lung volume decreases until it becomes "trapped" (MERRD, 1960).

It is therefore possible that in normal subjects flow volume curves generated outside a body plethysmograph can respond to changes in airway smooth muscle tone and that such tonal changes explain some of the variability associated with MEFV indices.

TABLE I PHYSICAL CHARACTERISTICS, TOBACCO CONSUMPTION, VENTILATORY CAPACITIES AND PEAK EXPIRATORY FLOW RATE IN

HEALTHY MALES

Sub. No. & RACE	Age (Yr)	Height (m)	Smoking (Pk-Yr)	FEV ₁ Obs. (L) ¹	FEV ₁ Pred. %	FVC Obs. (L)	FVC Pred. %	FEV ₁ /FVC (%) Obs. (%) ¹	FEV ₁ /FVC (%) Pred. (%)	PEFR L/min
1 C	27	1.80	-	4.48	105.5	5.40	106.0	85.1	103.6	796
2 C	30	1.68	-	4.45	119.3	4.73	105.2	93.4	115.0	678
3 C	31	1.80	-	4.18	101.0	4.92	97.1	85.1	106.0	916
4 C	37	1.83	0.375	4.23	103.6	5.98	117.9	71.0	91.0	599
5 C	31	1.66	-	4.00	110.0	6.06	116.0	79.1	98.6	925
6 C	36	1.88	0.75	4.82	147.5	6.03	111.5	79.9	102.0	982
7 A	34	1.78	0.75	3.83	96.5	3.91	79.6	98.0	124.0	774
8 A	29	1.77	-	3.68	89.6	4.66	93.6	79.0	97.8	784
9 N	27	1.87	-	4.28	95.0	5.14	93.0	83.3	102.5	789
10 N	31	1.76	-	3.56	89.5	3.85	79.0	92.5	115.3	784
MEAN *	32	1.78		4.36	114.5	5.35	109.0	81.9	102.7	816.8
SD *	3.8	0.09		0.29	17.4	0.55	7.7	7.4	8.0	153.4
MEAN **	29.4	1.76		4.09	101.4	4.82	98.6	85.1	105.5	810.3
SD **	1.8	0.07		0.36	11.1	0.50	11.9	5.8	7.1	85.5
MEAN	31.3	1.78		4.15	105.8	4.97	99.9	84.4	105.6	803.2
SD	3.4	0.07		0.39	17.3	0.74	13.9	8.1	9.8	115.7
C Caucasian, A Asian and N Negro			*	Caucasians only		**	Non-smokers only			

TABLE 2. Expiratory flow rates at two lung volumes in healthy males.

Sub. No.	MEF50 (L/min)		MEF25 (L/min)	
	Maximal	Partial	Maximal	Partial
1	277	298	110	115
2	317	356	178	145
3	370	430	166	186
4	438	496	159	185
5	262	232	99	84
6	297	223	137	53
7	668	611	235	189
8	272	206	81	56
9	323	269	99	99
10	413	426	153	96
M *	326.8	339.2	141.5	128.0
SD *	66.2	109.4	31.8	54.1
M **	320.1	316.7	127.0	111.6
SD **	54.9	89.8	37.6	42.7
M	364.4	354.7	142.0	120.8
SD	122.0	133.9	46.0	52.6

* Caucasians only

** Non-smokers only

Regression Equations(1) All n = 10

(a) $MEF50(P) = 1.006 MEF50 - 11.814$

$\bar{y} = 354.7 \pm 133.9 \quad COV_y = 37.7\%$

$\bar{x} = 364.4 \pm 121.98 \quad COV_x = 33.5\%$

$r = 0.916, \quad r^2 = 0.839, \quad t = 6.454 \quad P < 0.001$

(b) $MEF25(P) = 0.861 MEF25 - 1.498$

$\bar{y} = 120.8 \pm 52.58; \quad COV_y = 43.5\%$

$\bar{x} = 142.0 \pm 46.0 \quad COV_x = 32.4\%$

$r = 0.753, \quad r^2 = 0.568, \quad t = 5.862, \quad P < 0.001$

(2) Non-smokers n = 7

(a) $MEF50(P) = 1.43 MEF50 - 141$

$\bar{y} = 316.7 \pm 89.8 \quad COV_y = 28.4\%$

$\bar{x} = 320.1 \pm 54.9 \quad COV_x = 17.2\%$

$r = 0.874, \quad r^2 = \quad t = 4.02 \quad P < 0.01$

(b) $MEF25(P) = 0.90 MEF25 - 2.79$

$\bar{y} = 111.6 \pm 42.7 \quad COV_y = 38.5\%$

$\bar{x} = 127.0 \pm 37.6 \quad COV_x = 29.6\%$

$r = 0.795, \quad r^2 = 0.632, \quad t = 2.93 \quad P < 0.05$

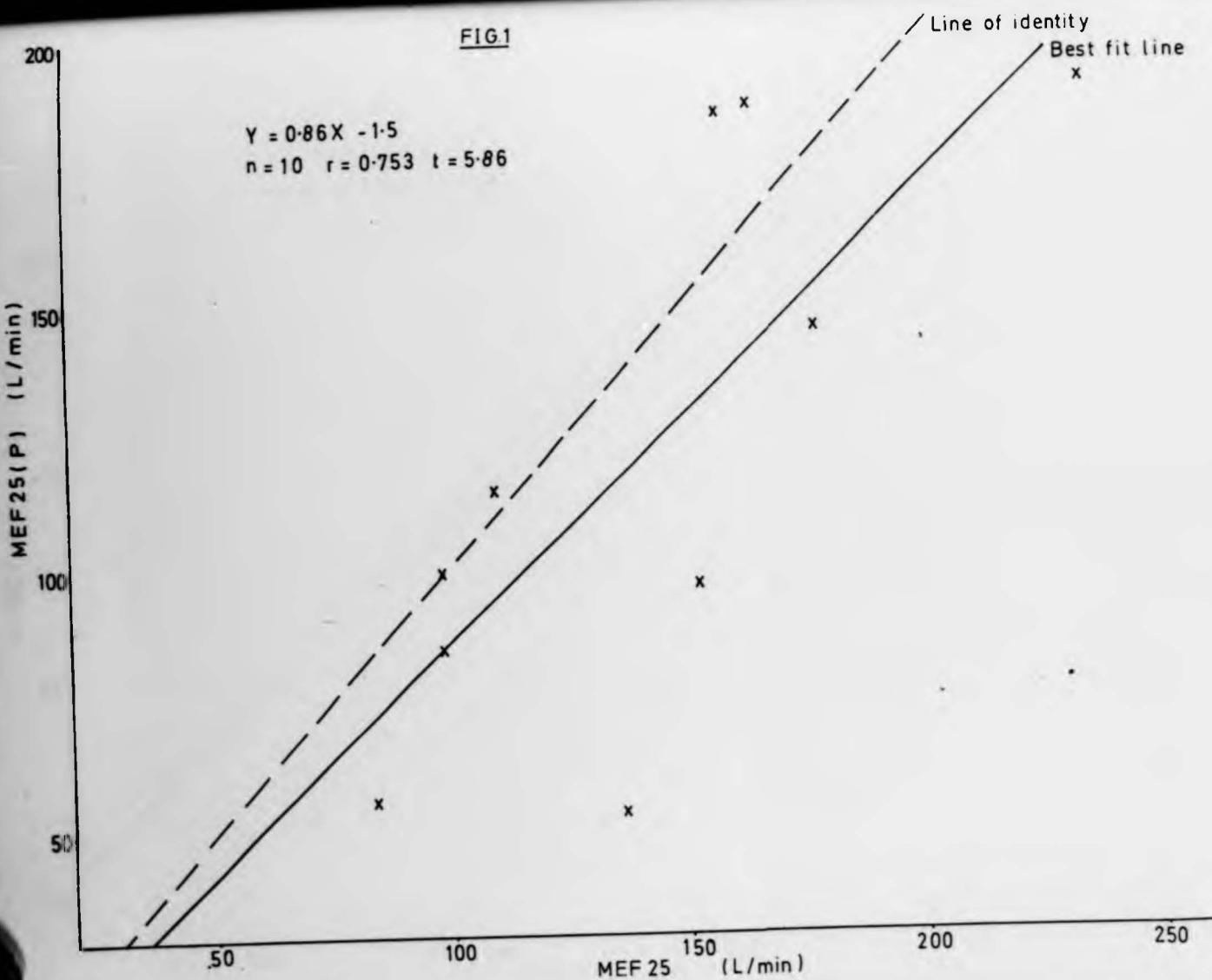


FIG 2

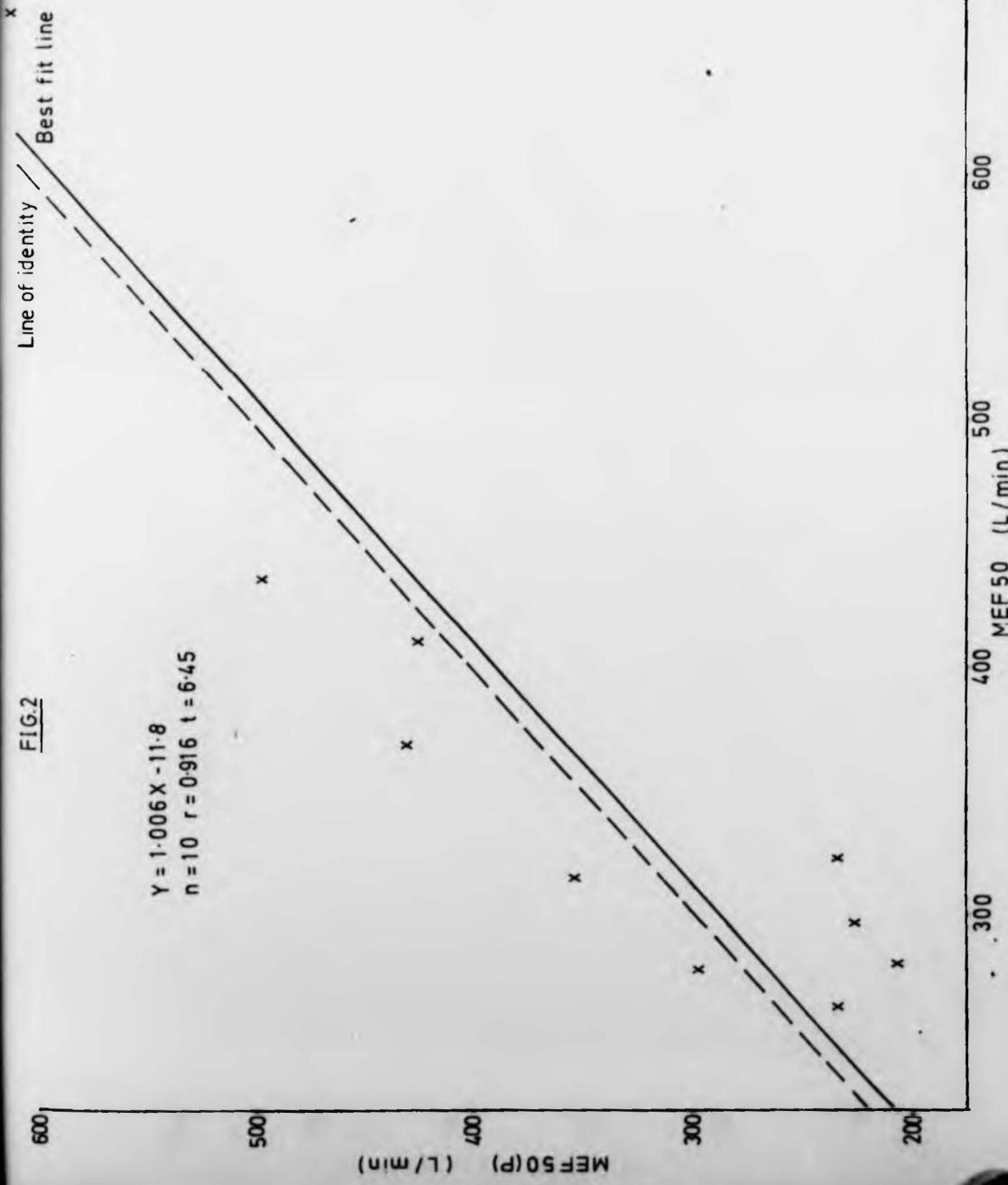


FIG 3 NONSMOKERS

25

$$\begin{aligned}Y &= 0.9X - 2.79 \\n &= 7 \quad r = 0.795 \quad t = 2.93\end{aligned}$$

MEF 25(P) (L/min)

200
175150
125100
75

125
100 MEF 25 (L/min) 125
150 175

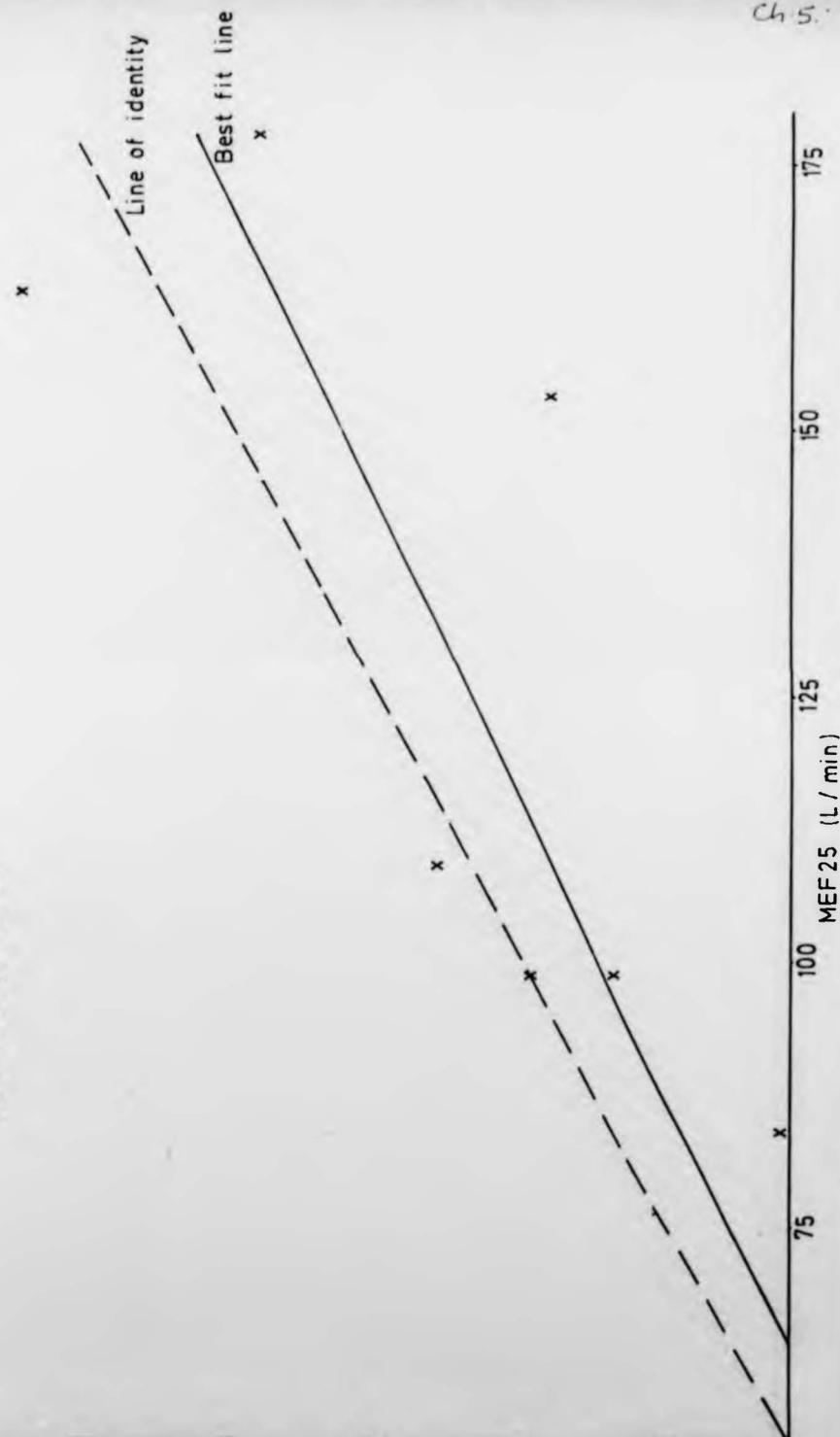


FIG. 3

NONSMOKERS

25

200

175

150

125

100

75

MEF 25(P) (L/min)

$$Y = 0.9X - 2.79$$
$$n = 7 \quad r = 0.795 \quad t=2.93$$

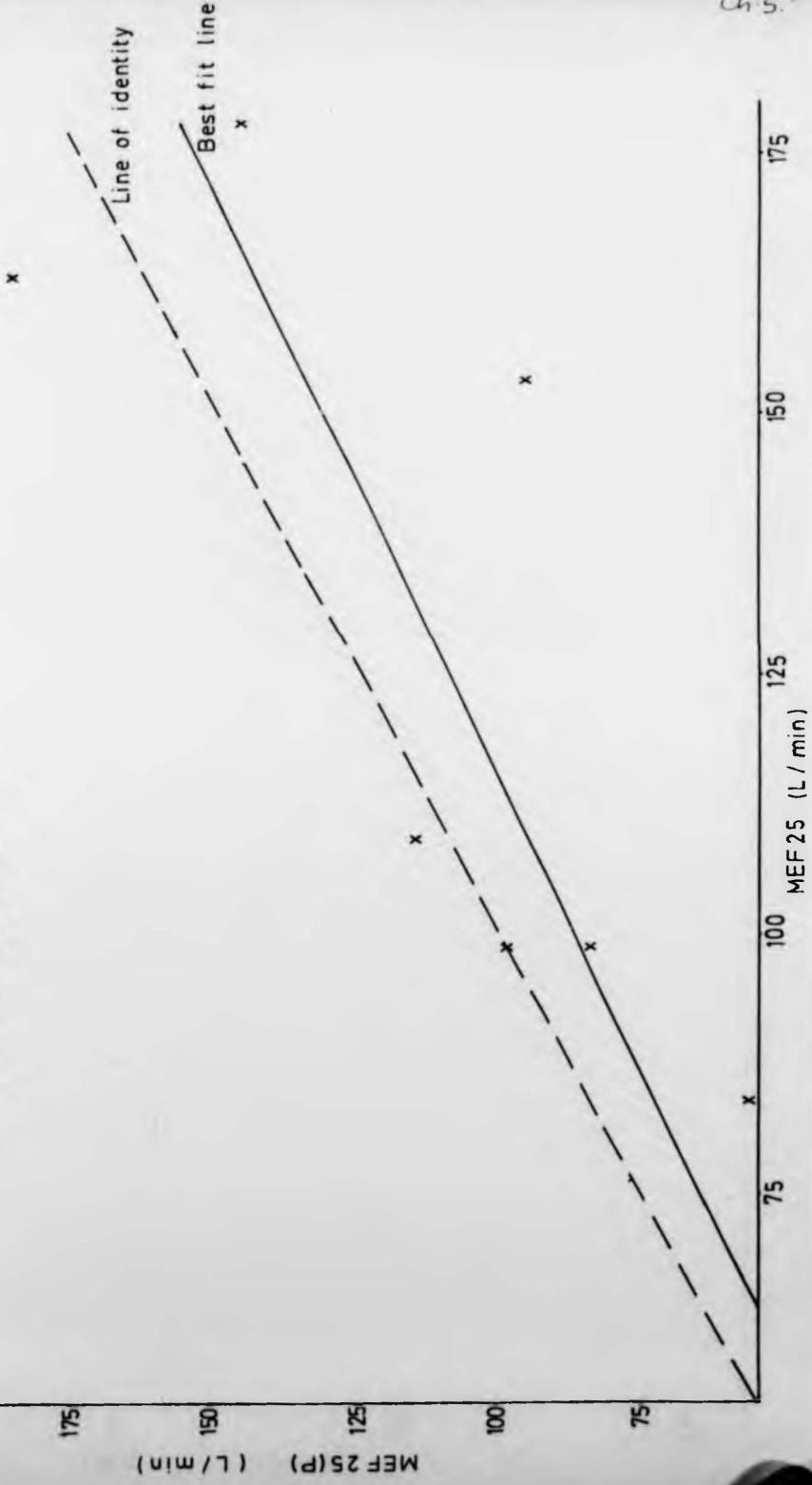
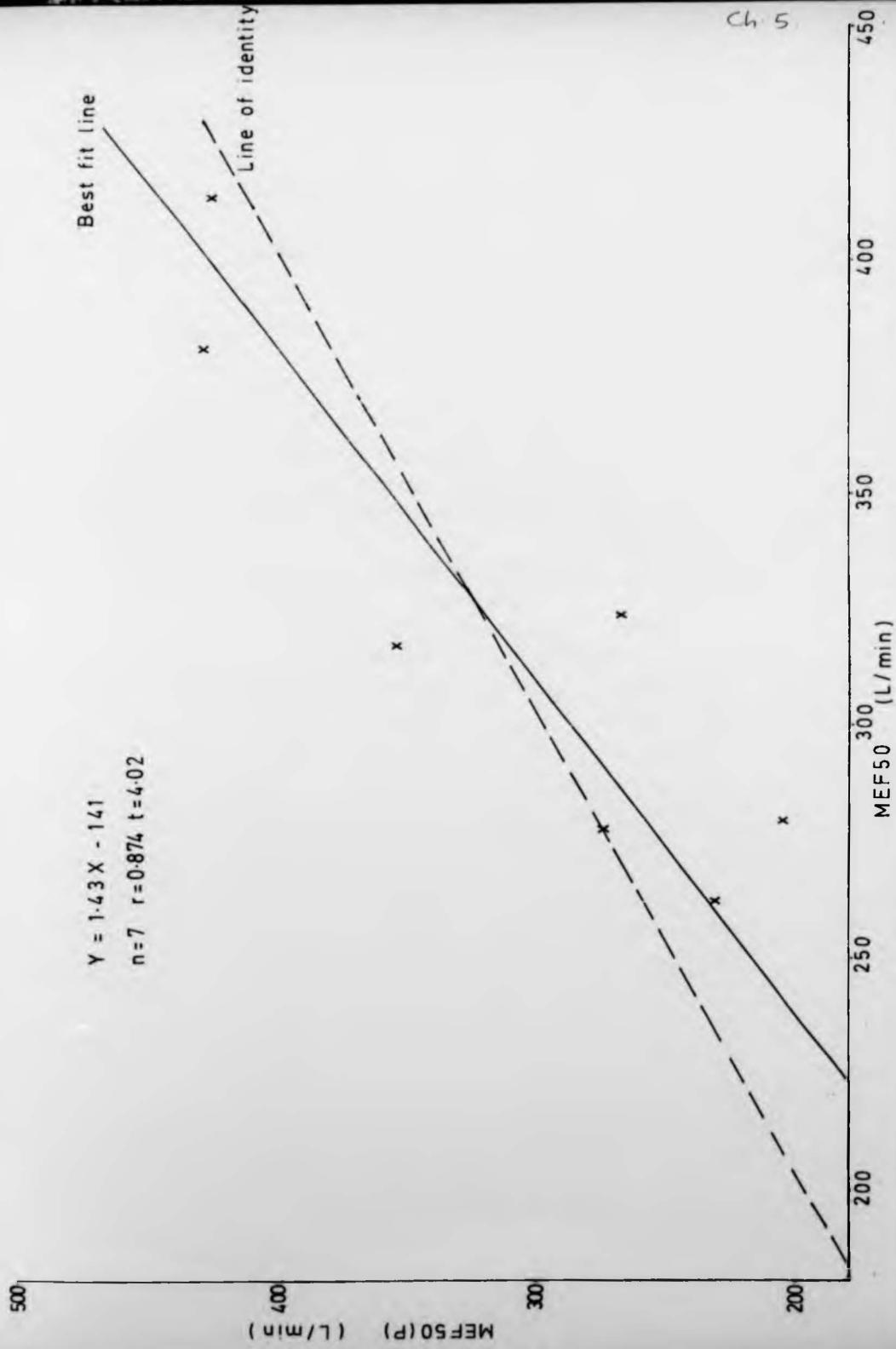


FIG4 NONSMOKERS



C H A P T E R 6

C O N T E N T S

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THE EFFECTS OF INHALED PROSTAGLANDIN E₂ (PGE₂) ON THE AIRWAYS OF
YOUNG HEALTHY ADULTS.

INTRODUCTION

The prostaglandins are a group of similar compounds derived from prostanoic acid. They are normal constituents of all animal tissues and are implicated in a wide variety of biological actions, including the relaxation of bronchial smooth muscle. (Bergström, 1968; Weeks, 1972; Main, 1964). Prostaglandins are rapidly and efficiently metabolised both near their site of production and in the general circulation. (Samuelsson, 1971). The prostaglandins are present in tissue in only small amounts: PGE₂ being in a relatively higher concentration in bronchial tissue than other prostaglandins, for example PGF_{2α} (Smith, 1972, 1975). PGE₂ has been shown to relax isolated bronchial smooth muscle (Main, 1964), and to dilate bronchi in animals and man (Sweatman, 1968; Large, 1969; Rosenthal, 1970; Cuthbert, 1969, 1971; Smith, 1972, 1975). The metabolites show much reduced potency in their action on smooth muscle. (Anggard, 1971).

Because of the reported rapid metabolism and the reduced potency of the metabolites on smooth muscle, PGE₂ seemed eminently suitable for producing short duration bronchodilation of the human lung airways. A possible disadvantage however was its reported irritancy and therefore bronchoconstrictive effect when given as aerosol from pressurised canisters. (Smith, 1975). The present study was carried out to investigate the magnitude of the bronchomotor response in healthy airways, and in particular to determine whether the response could be easily detected by methods other than by plethysmography. To achieve this doses were chosen which were larger than previously reported

(Smith 1975), and the aerosol given without propellants. A single dose of solvent (ethanol) was given as placebo in one of the experiments. Airway resistance, static lung volumes and flowrates on expiratory flow-volume curves were measured.

SUBJECTS AND METHODS

Subjects

Twenty healthy medical students of normal ventilatory indices aged 19 to 22 years were studied. The three smokers, two males and one female, were light smokers according to Medical Research Council criteria (less than $\frac{1}{4}$ gm of tobacco per day). After giving informed consent, all subjects were familiarised with the required breathing manoeuvres. At the time of testing all subjects were free from respiratory symptoms, had not smoked for at least one hour or taken aspirin within four hours. The physical characteristics and ventilatory indices of the subjects are given in Table 1.

Methods

Before and after aerosol inhalation (treatment) changes in airway resistance (R_{aw}) were measured by body plethysmography, flowrate changes by maximal and partial flow-volume curves (MEFV and PEFV), residual volume change by a closed circuit helium dilution technique and other lung volume changes by dry spirometry. FEV_1 and FVC were also measured on the MEFV. (Mk. II apparatus described in Chapter 1).

Flow volume curves

MEFV and PEFV curves were generated whilst the subjects breathed air. The partial curves (submaximal inspiration) were always done before maximal curves so that the reference point (TLC) was that of the

immediately succeeding maximal manoeuvre. Flowrates were measured at 50%VC (MEF50 or more briefly V50) and after 75% of the VC was expelled (V25). Measurements made on PEFV are distinguished by "P", for example V25(P). Results were taken as the mean of the last three of five "blows".

Body plethysmography

R_{aw} and thoracic gas volume (V_{tg}) were measured in a constant volume body plethysmograph using the method described by Dubois (1956). The principle of the method has been given in Chapter . The mean of five replicates was taken and specific airways resistance SR_{aw} , calculated by multiplication of R_{aw} by the V_{tg} at which it was measured.

Closed circuit helium dilution for measuring residual volume

The method used is described in Chapter 4 . The mean of two replicate manoeuvres was taken.

Dry spirometry

A bellows type dry spirometer, (Vitalograph), was used. In addition to measuring FEV_1 and FVC the maximal midexpiratory flow (MMEF) was obtained. MMEF was calculated as the slope of the line joining the 25% and 75% volume points on the Vitalograph tracing. Five forced expirations were done and indices calculated as the mean of the last three. All volumes were corrected to BTPS, but allowance was not made for the fact that in a dry spirometer the gas is not fully saturated even after 100 blows when it is said that water can be poured out of the spirometer. The error involved in this procedure is small and present in all the spirometer measurements.

Aerosol administration

Both drug (PGE_2) and placebo (ethanol) were given as aerosols produced by an ultrasonic nebuliser (Mistogen Nebuliser). The dose was calculated from the known concentration of the solution together with the loss of weight from the nebuliser and its connecting tubes. The PGE_2 was supplied as a sterile solution in ethanol (10 mg/ml "Prostolin E₂" Upjohn Ltd., and diluted with distilled water to give the desired concentration needed to get the doses in 10 breaths. The placebo was ethanol (20% v/v distilled water). Subjects inhaled a mixture of air and nebulised particles through a mouthpiece whilst wearing a noseclip. Total inhaled volume was monitored by a Wright's respirometer. Each breath (of approximately 1 litre) was followed by a one second breath-hold.

Particle sizing

The particles were sized by two methods. A sample collected by thermal precipitation was viewed under oil by light microscopy. The range found was 0.5- 2.5 μm , about 10% of the sample (by frequency) was larger than 2 μm . The mean aerodynamic diameter (by Schaefer cell) was $0.6 \mu\text{m} \pm 0.31 \mu\text{m}$.

PROCEDURE

The experiments were carried out at three morning sessions extending over a period of ten days. At each session pretreatment (baseline) measurement, treatment and post treatment measurements were done sequentially. Post treatment determinations on body plethysmograph and by closed circuit helium technique were timed at 5 to 15 minutes after inhalation, those on flow volume and dry spirometer were done

between 15 and 30 minutes post inhalation. This procedure was chosen in an attempt to ensure that the less sensitive tests were done when the drug effect should have been maximal.

The medical student volunteers were in two groups: the first (8 persons) attended for two sessions which were three days apart; the second (12 persons) attended for a single session. There were three experiments. The subjects for the first were the first group of volunteers. This group of volunteers were allocated to either of two doses of drug and one of placebo in randomised order subject to the constraint that half the volunteers received different drug doses. At the first session, only six students were present, but the group of eight was made up by substituting two very much older subjects. The blinded nature of this experiment was thus preserved, with the result that only six of the group received drug, but all eight were given placebo.

The second group of students were allocated to experiments two and three which were performed at the same session. These volunteers all had the same drug dose but were randomly allocated to be tested by plethysmography and flow volume curves or by helium dilution residual volume and dry spirometry. Again randomisation was constrained to allow an even split between test methods.

All subjects recorded their subjective post - treatment symptoms.

RESULTS

The mean difference between baseline and post-treatment values was calculated and expressed as a percentage of the baseline value, (post-treatment mean less pretreatment mean as a percent of pretreatment mean).

Statistical analysis was by Wilcoxon's signed rank sum test (SRST). The SRST is particularly suited to an investigation of this type since the sample size was too small to establish the extent of non-normality of distribution (Armitage, 1971). To calculate the SRST statistic "T", the observations are put in ascending order of magnitude, ignoring the sign, and given ranks 1 to n (zero values are ignored and tied values allotted the mean rank. The ranks take the sign of the original observations. T is calculated as the sum of the ranks of the positive or negative values.

Table 1 gives details of the subjects taking part in each experiment together with type of treatment and dose, type of tests and their baseline FEV₁, expressed as percentage of their predicted values. The mean FEV₁% predicted was not significantly different for the different experiments.

Table 2 details the percentage change in the various indices relating the changes to the dose administered. The probable overall effect is given by the grand mean for the six subjects. The effect of the placebo is also listed in Table 2. The results of experiment 3 is given in Table 3.

In experiment 1, a PGE₂ doses of 372 µg (given to three subjects) gave a mean rise of 2.3% (SE 1.4%) in FEV₁, a mean fall in FVC of 2.5% (1.1%), mean increase in MEF50 (10.8 (9.5)% MEF25 fell by 7.7 (10.4)% whilst the mean decreases in SR_{aw} and V_{tg} were 14% (9%) and 5% (5%) respectively. A further three subjects who received a dose of 184 µg of PGE₂ and gave a mean fall in FEV₁ of 7% (2%), decrease of 10% (2%) in FVC, increases of 9% (4%) and 5% (7%) in MEF50 and MEF25 respectively,

and also falls of 12% (5%) and 1% (2%) in SR_{aw} and V_{tg} respectively. The placebo given to all of the six above subjects (and two additional ones) gave a small but significant increase in FVC (mean 1% (2%), $P<0.05$ on SRST). The probable overall effect of a dose of PGE_2 greater than 180 μg in six subjects is given by the mean for $n = 6$ in Tables 1 and 2. Only FVC and R_{aw} changed significantly.

In experiments 2 and 3 where all twelve subjects were given a dose of 45 μg of PGE_2 and half of them tested on the plethysmograph and flow-volume apparatus and the other half on the spirometer and helium dilution apparatus, regarded as a single group (Subjects 9 to 20), 45 μg of PGE_2 gave changes (significant at 5% level by SRST) in the following indices:-

- (i) R_{aw} and SR_{aw} decrease (respective (Table 2) means of 20% (6%) and 21% (5%))
- (ii) RV decreases (Table 3)(mean 14% (13%)).

The change in FVC was equivocal; six subjects (9-14) showed a small but significant increase and the other six showed a larger but nonsignificant decrease. The FEV_1 change in both groups (9-14 and 15-20) were nonsignificant although there was a mean fall (3%) in the first and a slight increase in the second (0.3%).

Subjective comments of the subjects

All of the eight subjects who had drug and placebo (six had both, two had placebo only) reported onset of cough which they said was caused by throat irritation. The incidence of coughing was similar after inhalation of either drug or placebo but whilst they all reported increased phlegm after drug, only one person did so after placebo.

The remaining twelve subjects reported throat irritation resulting in cough. For four of these twelve, that was the only comment; two reported increased phlegm in addition to cough, four said that chest tightness was experienced. The tightness began shortly after inhalation and persisted for approximately 30 minutes. One person wheezed and another said that heartrate was increased.

DISCUSSION

The results reported here were obtained on a study which was not double blind because it was impractical to achieve and did not thwart the primary purpose of the study. It was blinded to the extent compatible with the safety of the subjects (young medical students) whilst allowing valid comparisons to be made. The subjects, although aware of the object of the experiments were not given details of doses or of when drug or placebo were being administered. The measurements were made by observers who were unaware of the type of treatment or dose received by the subjects. At all times during the study the subjects were under the supervision of an observer, (the author), who was aware of type of treatment and dose received.

Aspirin is known to inhibit prostaglandin synthesis and release (Vane, 1971); none of the subjects studied here had taken aspirin for at least four hours before their airways were challenged.

In the present study SR_{aw} fell significantly after all the doses of PGE2, confirming dilation of large airways. Since there was a rise in SRaw following placebo, the measured effect is attributed to the active constituents of the drug. The larger mean fall on the smaller doses is suggestive of an increase in airway compressibility

with increasing dose even at low flow rates. The compressibility phenomenon could explain the equivocal changes in FEV_1 and FVC.

Evidence of small airway dilation is less direct than that for large airways. The indices that monitor small airway function are dependent on forced expiration and will thus be influenced by large airway compressibility. Larger mean changes in MEF50 and MEF25 resulted from higher doses of drug than for smaller doses. V_{tg} fell after treatment with drug (all doses), and placebo. This fall reflects change in the FRC. An FRC change can result from changes in either RV, ERV or both. The body plethysmograph data suggests that constriction followed placebo (increase in SRaw), so that an increase in RV may have resulted. An increase in ERV was found following low dose of drug; this, combined with the observation of dilation supports a fall in RV. Such a fall was detected by the closed circuit helium dilution technique. The change in RV is likely to have been underestimated because of the fall in FVC. Since RV may be defined as "the volume of gas in the chest at the end of a forced expiratory effort when the combined effects of the closure of the small airways and the elasticity of the thoracic cage prevent further expiration", (Cotes; 1975), an increase in small airway calibre is a likely explanation for the observed fall in RV.

Clinical symptoms associated with bronchoconstriction namely chest tightness and wheezing were described by the volunteers. The wheezing might be explained by the increased compressibility already mentioned since wheezing during a forced manoeuvre has been partly attributed to this cause by Bouhuys (1974). The irritation may have produced reflex bronchoconstriction (WIDDICOMBE, 1975)

but this would have been diminished or reduced by the full inspiration needed for the forced manoeuvres (Vincent, 1970). The irritation and phlegm production would reduce subject acceptance of the drug and may thus be a disadvantage of PGE₂ as a bronchodilator.

Smith et.al. (1975) obtained increased specific airway conductance (SG_{aw}) in five healthy males aged between 27 and 42 years following the inhalation of 55 μ g of PGE₂ delivered from pressurised canisters. The metered aerosol doses which they used contained the drug in ethanol solution with dichlorotetrafluoroethane and dichlorodifluoromethane as propellants. A mean increase of 17.6% in SG_{aw} (maximum effect) was found in their subjects 15 minutes after inhalation, the effect being complete at 50 minutes.

A slightly larger bronchodilation (20% decrease in SR_{aw}) was found in the present study despite a smaller dose (45 μ g) and collecting the plethysmograph data between 5 and 15 minutes post inhalation. Some of this possible difference may be due to less irritation from the smaller dose. But another explanation might be due to differences in the deposition of the aerosols used. Deposition of aerosols is known to be affected by particle size and mode of inhalation (PAVIA, 1977). The particle size used by Smith et.al. (1975) was not stated but in some instances the size distribution of particles generated from pressurized cans may be similar to that made by ultrasonic nebulisers in spite of differences in the principle of operation of these two types of nebulisers (Raabe, 1975). For ultrasonic nebulisers, the size of particles carried out of the generator by the airstream is highly dependent on rate of removal from the site of formation as well as on the hygroscopicity of the nebulised particles. Since particle

sizing could not be carried out under identical conditions to those prevailing during inhalation by the subjects, the sizes reported in the study can only be regarded as a guide.

It is felt that the size distribution of the particles in the present study were such that deposition would have occurred in large and small airways. If the measured sizes are representative of those particles entering the respiratory tract and no change in size occurred in transit through the tract, the breath holding pause after each breath would have increased the chance of small airway deposition. If on the other hand the particles increased in size during transit of the tract, then the "growth" factor might be the determining factor governing small airway deposition. Dautrebande (1961) estimated the growth factor of NaCl microcrystals (maximum initial size of 1 μm) to be 7. It is felt that a growth factor much different to that of NaCl is unlikely to apply to PGE₂ although the larger particles ($>1\mu\text{m}$) would probably not increase as much and with growth factors of this order the particles could still be deposited in the small airways.

In another study, Smith (1974), found that intravenous infusion of PGE₂ (5 to 10 $\mu\text{g}/\text{min}$) produced only slight or variable effects on FEV₁. This observation lends weight to the proposition that large changes in FEV₁ were not measured in the present study because PGE₂ relaxes airway smooth muscle and in so doing increases the compressibility of the airways and limits the maximum airflow during forced expiration.

It is concluded that PGE₂ aerosol inhalation dilated the airways of the healthy volunteers in this study, but that the effective

dilation is not easily measured by tests which use a forced expiration. The irritation reported subjectively was probably due to the drug rather than the solvent and this together with increased phlegm would militate against the use of PGE₂ as a bronchodilator in the response testing of human lung airways.

TABLE 1 THE PHYSICAL CHARACTERISTICS, INITIAL VENTILATORY CAPACITIES, TYPES OF TESTS, TREATMENT, DOSES AND CHANGE IN VENTILATORY CAPACITIES AS PERCENTAGE OF INITIAL VALUES FOR 20 VOLUNTEERS.

Sub. No.	SEX	HEIGHT (m)	AGE (Yr)	FEV ₁		FVC		FEV ₁ /FVC%		TESTS **	TREATMENT ***	PGE ₂ DOSE (mg)	%Change Placebo FEV ₁ FVC		%Change Drug FEV ₁ FVC	
				Obs.	%Pred.	Obs.	%Pred.	Obs.	%Pred.				FEV ₁	FVC		
1	M	1.78	20	4.57	104	5.46	105	84	99	B and FV	D and P	369	-1.2	5.2	2.2	-4.2
2	M	1.90	20	4.72	97	5.49	94	86	98	"	"	369	1.7	3.2	4.8	-0.4
*3	M	1.85	20	3.70	79	4.93	88	76	90	"	"	378	1.1	0.2	0.0	-2.9
n =	3															
MEAN		1.84	20	4.33	93.3	5.29	95.7	82	95.7			372	0.53	2.9	2.3	-2.5
SD (SE)		0.06	(0)	0.32	(12.9)	(0.18)	(5.0)	(3.1)	(2.9)			(3)	0.88	(1.5)	(1.4)	(1.1)
4	M	1.79	21	3.68	83	5.32	101	69	82	B and FV	D and P	184	6.4	2.8	-10.2	-13.5
*5	M	1.79	20	4.50	101	4.67	89	96	114	"	"	184	1.3	3.3	-7.5	-7.2
+6	M	1.92	20	4.31	88	5.79	91	74	88	"	"	184	-5.0	-6.9	-3.8	-9.0
n =	3															
MEAN		1.83	20.3	4.16	90.7	5.26	93.7	79.7	94.7			184	0.9	-0.3	-7.2	-9.9
SD (SE)		0.08	0.58	(0.25)	(5.4)	(0.32)	(3.7)	(8.3)	(9.8)			(0)	(3.3)	(3.3)	(1.9)	(1.9)
n =	6															
MEAN		1.84	20.2	4.25	92.0	5.28	94.7	80.8	95.2			278	0.72	1.3	-2.4	-6.1
SD (SE)		0.06	0.4	(0.18)	(4.1)	(0.17)	(2.8)	(4.0)	(4.6)			(42)	(1.5)	(1.8)	(2.4)	(2.0)

47	F	1.75	19.0	2.79	77.0	3.28	69	85	98	B and FV	P only	-	-8.2	-7.8	-	-
8	M	1.82	21	3.73	82	5.07	91	71	88	"	"	-	0.9	5.1	-	-
<i>n = 8</i>																
MEAN		1.83	20.1	4.00	88.8	5.00	91.4	80.5	94.6				-0.38	0.64	-	-
SD (SE)		0.06	0.6	(0.23)	(3.7)	(0.28)	(3.8)	(3.1)	(3.5)				(1.6)	(1.8)	-	-
9	M	1.73	21	3.72	89	4.15	81	90	78	B and FV	D only	"5		-43.6	6.8	
10	F	1.64	20	2.92	89	3.70	97	79	91	"	"	"6		-7.5	5.6	
11	M	1.83	21	4.85	106	5.38	99	90	107	"	"	"5		3.6	4.8	
12	M	1.80	22	4.37	99	4.66	88	91	112	"	"	"6		4.6	3.9	
13	M	1.64	22	2.67	72	3.73	88	72	86	"	"	"5		18.5	1.9	
14	F	1.78	21	3.51	96	3.55	79	99	114	"	"	"5		7.2	-11.1	
<i>n = 6</i>																
MEAN		1.74	21.2	3.67	91.8	4.2	89.2	87.3	98			"5.3		-2.9	1.97	
SD (SE)		0.08	0.75	(0.34)	(4.8)	(0.29)	(3.1)	(4.1)	(6.1)			(0.2)		(8.8)	(2.7)	
15	M	1.76	20	4.23	97	5.17	101	82	97	HD and S	D only	"5		-1.2	0.2	
16	F	1.70	21	3.18	93	3.50	85	91	105	"	"	"7		-4.0	-3.7	
17	M	1.72	20	3.76	94	5.18	106	73	81	"	"	"4		9.9	-3.9	
18	F	1.65	20	3.45	101	3.78	98	91	105	"	"	"6		-8.1	-6.1	
19	F	1.67	20	3.55	105	3.90	98	91	105	"	"	"6		-6.3	-7.1	
20	M	1.70	19	4.98	120	5.95	105	81	99	"	"	"5		11.7	-6.6	
<i>n = 6</i>																
MEAN		1.7	20	3.86	102.2	4.58	98.8	85.3	99.2			"5.5		0.33-4.5		
SD (SE)		0.04	0.63	(0.27)	(4.1)	(0.40)	(3.1)	(3.0)	(3.4)			(0.4)		(3.4)	(1.1)	

* indicates smoker; + last onset of cold/flu 4 weeks previously; # : ~~last onset of cold/flu 2 weeks previously~~

** : ~~last onset of cold/flu 2 weeks previously~~

*** : ~~last onset of cold/flu 2 weeks previously~~

**** : ~~last onset of cold/flu 2 weeks previously~~

TABLE 2 CHANGES (AS PERCENTAGE OF INITIAL VALUES) IN FLOWRATES, AIRWAY RESISTANCE AND THORACIC
GAS VOLUME AFTER INHALATION OF DIFFERENT DOSES* OF PGE₂ AND PLACEBO

Sub. No.	MEF50	Drug	Placebo									
			MEF25	MEF25P	R _{aw}	SR _{aw}	V _{tg}	MEF50	MEF25	R _{aw}	SR _{aw}	V _{tg}
1	-0 -9	14.8			-16.1	-30.0	-14.3	-0.5	-10.4	15.8	2.5	-11.4
2	29.2	-20.5			-0.7	1.1	1.9	3.0	7.6	5.5	9.7	2.3
3	1.1	3.4			-10.3	12.6	-2.5	0.7	0.0	11.2	4.5	-5.9
n = 3												
MEAN (SE)	10.8 (9.3)	-7.7 (10.4)			-8.8 (4.7)	-13.8 (9.0)	-5.0 (4.8)	1.07 (1.02)	-0.9 (5.2)	10.8 (3.0)	5.6 (2.1)	-5.0 (4.0)
4	6.9	6.3			-9.6	-14.1	-5.0	15.2	26.3	3.2	-2.9	-5.8
5	2.2	7.0			-0.2	-18.8	1.0	0.8	33.2	-3.7	-6.8	-2.8
6	16.4	-16.7			-3.6	-2.2	1.5	15.4	16.4	-1.5	-0.6	0.9
n = 3												
MEAN (SE)	8.5 (4.2)	5.3 (6.9)			-4.5 (2.7)	-11.7 (4.9)	-0.8 (2.1)	10.5 (4.8)	25.3 (4.9)	-0.7 (2.0)	-3.4 (-1.9)	-2.6 (-1.5)
n = 6												
MEAN (SE)					****							
7					-6.8 (2.6)	-12.8 (4.6)	-2.9 (2.5)	5.8 (3.0)	12.2 (6.7)	5.1 (3.0)	1.1 (2.4)	-3.8 (2.1)
8								-14.0	-18.4	20.6	12.4	-7.0
n = 8												
MEAN (SE)								3.7	10.2	6.4	4.8	-1.6

9	0.7	22.7	-10.8	-19.6	-22.4	-3.5
10	6.0	31.3	22.3	-11.3	-13.7	-2.8
11	5.9	24.9	19.0	-16.7	-14.0	5.3
12	19.8	7.4	-4.8	-17.7	-12.6	6.0
13	-4.1	-19.8	11.3	-2.3	-7.2	-5.2
14	-10.0	28.7	-29.4	-23.6	-26.1	-3.0
			****	****		
n = 6						
MEAN (SE)	3.1 (4.2)	8.3 (9.9)	1.3 (8.1)	-20.2 (6.1)	-21.0 (5.4)	-0.5 (2.0)

* See Table 1 Details of Doses and treatment.

**** Indicates statistically significant change by Wilcoxon's Summed Rank Sign Test

TABLE 3 CHANGES (AS PERCENT OF INITIAL VALUES) IN MMEF, ERV AND RV FOR SIX VOLUNTEERS AFTER
INHALATION OF 45^{ug} PGE₂

Sub. No.	MMEF	ERV	RV
15	5.0	0	-16.7
16	3.0	-2.4	-26.4
17	-4.1	2.6	-63.1
18	9.0	-3.1	-27.8
19	4.7	-15.3	-
20	20.0	50.0	-83.8
n =	6	6	5
MEAN (SE)	6.3 (3.3)	5.3 (9.3)	-43.6 (12.8) ****

CHAPTER 7

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Table 9

RATING OF INDICES IN ORDER OF ABILITY TO DETECT
CHANGES IN BASELINE VALUES

Table 10a

RATING OF INDICES IN ORDER OF ABILITY TO DETECT
DRUG INDUCED REVERSIBLE CHANGES

Table 10b

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RATING OF INDICES IN ORDER OF ABILITY TO DETECT
DRUG INDUCED REVERSIBLE CHANGES

Table 10b

THE SPECIFICITY AND SENSITIVITY OF SOME INDICES OF SMALL AIRWAY
FUNCTION.

INTRODUCTION

Several workers, (Ingram et.al. 19~~71~~; Woolcock et.al. 1969; Anthonisen et.al. 1963), have shown that obstruction and other functional defects of the airways can exist undetected by routine lung function tests for many years. Until recently by the time these were clinically evident the obstruction of the airways was usually at an irreversible stage. Anthonisen, (1963) showed that some bronchitics whose pulmonary function values from pre-existing tests were within the normal expected range had pathophysiological abnormalities. These abnormalities were attributed to sites in airways having internal diameters less than 2 mm. In the inflated lung the resistance to airflow of such small airways is a relatively small portion of total pulmonary resistance (Macklem et.al. 1967), making it possible to have considerable peripheral obstruction before maximal flowrates fall outside the expected normal range (Woolcock et.al. 1969).

Newer functional tests of small airways have been developed specifically to detect changes in the 'quiet' zone of the lungs (for example Dynamic lung compliance, Cdyn; Closing Volume, CV; Flowrates at low lung volumes measured on maximal expiratory flow-volume curves, MEFV). Maximum benefit from these tests will result if they are shown to have high sensitivity and specificity as well as being simple in execution. Simplicity in execution is of particular importance when such tests are to be used as epidemiological tools since a large number of persons will need to be tested in a relatively short time.

Certain procedures used as small airway tests, for example Radio-

aerosol deposition patterns (Lourenco, 1971), frequency dependence of Cdyn (Woolcock, 1969) and MEFV performed in a body plethysmograph (Bouhuys, 1969), are not easily adaptable for "field use". Others, such as CV (Green, 1972) and MEFV performed without a plethysmograph (Fry, 1960) are more readily amenable to large surveys carried out at sites other than at specialist laboratories.

Various indices of small airway function can be derived from the closing volume and flow volume tests. This study attempts to assess the specificity and sensitivity with which thirteen such indices detect drug induced reversible airway function.

SUBJECTS AND METHODS

Twenty five healthy subjects, classified as smoker and non-smoker according to Medical Research Council criteria (Medical Research Council Questionnaire on Respiratory Symptoms, 1966), were studied. Detailed and mean physical characteristics, tobacco consumption and ventilatory indices are given in tables 1 and 2. No clinical evidence of pulmonary symptoms was revealed by questionnaire. Only one person, (CIE), admitted to having suffered from "Hay fever" in the remote past. Informed consent was obtained from all subjects.

Design of study

Each subject was tested on four consecutive days at the same time of the day. Measurements on three tests:- Closing volume, Air Flow-volume curves and Helium-Oxygen Flow-volume curves, were made before and after the airways were challenged by each of four inhaled broncho-active drugs. The tests were taken in randomised order and the provoking drug chosen by random entry into a Latin Square block of Sessions

and Treatments. The day was divided into morning and afternoon sessions. A pair of volunteers were studied at each session. The members of a pair were of a different classification; male and female or smoker and non-smoker.

The manoeuvres required for each session were explained and where necessary practiced at the start of each session. Measurement of baseline (prechallenge) values was followed by challenge and post challenge measurements, due allowance being made for the expected length of time needed for each drug to produce maximal effects.

Apparatus

The apparatus is fully described elsewhere (Chapter 1). The arrangement was such that rapid sequential performance of the various manoeuvres was possible from two separate breathing ports; one for Closing Volume and the other for both types of Flow-volume curves.

Drugs

Two bronchoconstrictor (Acetylcholine (Ach.) and Prostaglandin F_{2α} (PGF_{2α}) and two bronchodilator (Ipratropium bromide (Sch1000) and Salbutamol (Ventolin) drugs were used as challenging agents. Ach., PGF_{2α} and Sch1000 were nebulised as aqueous solutions in a Wright's nebuliser operating at driving pressures between 5.4 and 6.7 Pa (8 to 10 pounds per square inch). Particles generated under these conditions were of mass median diameter 0.8μm with geometric standard deviation of 2.2μm.

The nebuliser was weighed before and after drug administration and doses were estimated from loss of weight.

Ventolin was administered from a hand held nebuliser (metered dose

inhaler (MDI)) supplied by the makers. The dose delivered per valve depression was given by the manufacturer as 100^µg.

In order to obtain as wide a response range as practicable two distinct doses of each type of drug (constrictors and dilators) were given to each subject. The changes induced by Bronchoconstrictors were assessed by "partial" manoeuvres described below, whilst those following bronchodilators were assessed by "maximal" procedures.

TECHNIQUES

Closing volume

The single breath helium bolus method described by Green, (1972) was used. Bolus volume was 300 ml. Inspiratory and expiratory flow rates were in the range 300 to 500 ml per second. The volunteers tried to maintain steady flow by keeping inspiratory and expiratory indicator lamps glowing at constant brilliance. (See picture of apparatus, Chapter 1). The flow rates were simultaneously recorded on a storage oscilloscope from which polaroid photographs were made. Tracings produced when flow rates fell outside the above range were discarded.

The "partial" Closing volume manoeuvre

Conventional Closing volume technique involves successive expiration to residual volume (RV), inspiration of helium bolus followed by air to total lung capacity (TLC) and subsequent expiration to RV during which phase expired volume is plotted against marker gas (helium) concentration. In a "partial" manoeuvre, inspiration is terminated at a volume below TLC (typically 80% TLC). The slow vital capacity (SVC) needed as a reference volume range is necessarily made in a separate manoeuvre.

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In this study, the SVC was obtained at the end of the set of "partial" replicates; lung volume history was maintained constant throughout by preceding each manoeuvre by a period of quiet breathing. The mean of three replicates was taken.

Air Flow-volume curves

Maximal expiratory flow volume curves (MEFV) or partial expiratory flow volume curves (PEFV) are produced according to whether the vital capacity (VC) or a portion of the VC is expelled with maximal force, flowrate being plotted on the ordinate and volume on the abscissa of an XY recorder. Curves obtained when air is breathed are designated (Air), whilst those traced when a mixture of 80% helium and 20% oxygen is breathed are called (He-O₂) or more briefly (He).

In this study, replicates of PEFV were always obtained before those of MEFV and special attention was paid to the maintenance of constant lung volume history. The reference VC for post-treatment measurements was the mean baseline value.

A modification of the equipment (Chapter 1) allowed the forced expiratory volume in one second (FEV₁) to be obtained from the MEFV (Air). (This is seen as a 1 second "notch" on the graphs Chapter 1). The forced vital capacity (FVC) was also obtained from the MEFV traces.

Helium-Oxygen Flow volume curves

MEFV(He) and PEFV(He) were obtained in a similar manner to Air curves. The He-O₂ mixture was washed into the lung during a period of quiet breathing from a closed system which contained a CO₂ absorber. At the end of 2 to 3 minutes rebreathing, the subject completely emptied the lungs before being connected to another (larger volumed) closed

system having a fresh supply of the He-O₂ mixture. Either two VC's or three "partial" VC's were taken from this supply and nonforcefully expired to the room air before the desired MEFV(He) or PEFV(He) replicates were done. This procedure ensured:-

- a) constancy of lung volume history before each replicate manoeuvre, and
- b) that the composition of the mixture during measurements was similar to that used when calibrating the apparatus
(Chapter 1)

Drug inhalation procedure

The nozzle of the nebuliser was held about 5 cm from the open mouth. A two second "burst" of particles were directed into the mouth as the subject (not wearing noseclips) inhaled slowly but fully starting from RV. A 5 second apnea at TLC was followed by normal breathing. The procedure was repeated. Two "bursts" were given from the Wright's nebuliser and two "puffs" from the MDI.

RESULTS

Some indices were measured directly from the XY recorder charts and others calculated. The directly measured indices were:-

- i) MEF25(Air) and MEF25(P)(Air)...the flowrate at 25%VC
(RV = 0%) measured on the Maximum Expiratory Flow Volume curve (Air) and Partial Expiratory Flow Volume Curve (Air) respectively.
- ii) MEF25(He) and MEF25(P)(He)...the flowrate at 25%VC measured on the Maximum Expiratory Flow Volume Curve (He) and Partial Expiratory Flow Volume Curve (He) respectively.

- iii) $\text{MEF}^{10}\text{O}(\text{Air})$ and $\text{MEF}^{10}\text{O}(\text{P})(\text{Air})$...the flowrate at 10% measured on the Maximum Expiratory Flow Volume Curve (Air) and Partial Expiratory Flow Volume Curve (Air) respectively.
- iv) $\text{MEF}^{10}\text{O}(\text{He})$ and $\text{MEF}^{10}\text{O}(\text{P})(\text{He})$...the flowrate at 10% measured on Maximum Expiratory Flow Volume Curve (He) and Partial Expiratory Flow Volume Curve (He) respectively.
- v) CV...the difference in volume between the "onset of phase iv" and RV. (See closing volume tracing, Chapter 1).
- vi) FEV_1 ...measured on MEFV(Air).
- vii) FVC...measured on MEFV(Air).
- viii) SVC...the slow vital capacity measured on the conventional CV manoeuvre.

The calculated indices were:-

- i) $\text{MEF}25(\text{He}/\text{A})$...the fraction $\text{MEF}25(\text{He})$ minus $\text{MEF}25(\text{Air})$ divided by $\text{MEF}25(\text{Air})$. This fraction was calculated from both MEFV and PEFV.
- ii) $\text{MEF}^{10}\text{O}(\text{He}/\text{A})$...the fraction $(\text{MEF}^{10}\text{O}(\text{He}))$ minus $\text{MEF}^{10}\text{O}(\text{Air})$ divided by $\text{MEF}^{10}\text{O}(\text{Air})$ calculated from MEFV and PEFV.
- iii) IFVP (Isoflow volume point)...the volume, (as percent VC) where flowrates on (He) and (Air) curves are of equal value.
- iv) $\text{FEV}_1/\text{FVC} (\%)$... FEV_1 expressed as percent of FVC.
- v) $\text{CV/SVC} (\%)$...CV expressed as percent of SVC.

Baseline Data

At each session mean values for each index were found as the mean of three manoeuvres; the overall mean of the four such values were also calculated. The overall means for each of the subjects were used in a Regression Analysis. From the baseline data at each session a mean value for the members of the constituent groups was obtained; these are given in tables 2a and 2b for non-smokers and smokers and for males and females respectively. The means for those directly measured indices obtained on both "partial" and "Maximal" manoeuvres are given in table 3 for smokers and non-smokers.

Regression Analysis

Multiple Linear Regression Analysis has the basic assumption that the indices are linearly related to a number (n) independent variables. If Y is the dependent variable and $X_1 \dots X_n$ are the independent variables, then:-

$$Y = a + b_1 X_1 + b_2 X_2 + \dots + b_n X_n$$

where " a " is the intercept and b_i (i taking all values from 1 to n) are the regression coefficients. The coefficients are estimated by the "least square method"; the "goodness of fit" of the regression line is conveniently summarised in the square of the multiple correlation coefficient, R^2 , (Kendall, 1975). R^2 indicates the fraction of the variation accounted for by the regression. The computer program used (Statistical Package for the Social Sciences, (SPSS), subprogram REGRESSION) computes the coefficient R by "forward stepwise inclusion" (SPSS, 1975). The simple coefficient, r , was also calculated and the significance of the regression for each independent variable was estimated from the relationship:- $t = r \sqrt{(N-2)/(1-r^2)}^{1/2}$, where N is the number of observations.

Effect of Height, Weight, Age and Smoking:

With each index as the dependent variable, multiple regression with height, weight, age and smoking (total number of cigarettes or equivalent smoked by a subject) as the independent variables was carried out on the data obtained from males, females, smokers and non-smokers.

For non-smokers data, regressions with the following attained overall statistical significance:- (F is Snedecors ratio). CV ($F = 16.17, P < 0.0001$); SVC ($F = 6.17, P < 0.005$); CV/SVC ($F = 10.15, P < 0.0001$); FEV₁ ($F = 15.26, P < 0.0001$); FVC ($F = 6.94, P < 0.003$) and FEV₁/FVC ($F = 5.54, P < 0.008$). In none of the remaining classifications did the regressions reach overall significance. Tables 4, 5, 6 and 7 show the R^2 and r values for each independent variable and give the significance of the individual regressions for those variables, based on the t distribution. All probabilities less than 5% are shown as *.

Discriminant Analysis

The averaged baseline data were analysed by multivariate analysis methods for each of the four sessions. The Discriminant Function (D.F.) is a multivariate extension of multiple regression. It provides a linear function of the observations which gives maximum discrimination between the groups upon which the observations were made. The D.F.s were obtained using the Statistical Package for the Social Sciences (SPSS) subprogram "DISCRIMINANT". This program calculates standardised D.F. coefficients. In a particular D.F. the magnitude of the modulus of the coefficient reflects the importance of the index in differentiating between the groups. For example, if data from smokers and non-smokers result in a D.F. having the two terms:- 0.9 MEF25 (Air) and +0.8 MEF25 (He)

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then the coefficients are -0.9 and +0.8 with moduli of 0.9 and 0.8. MEF25(Air) therefore plays a more prominent role in distinguishing between the smokers and non-smokers than does MEF25(He).

A typical D.F. equation obtained from the data of a session is as follows:-

Ach Session:-

$$\begin{aligned} \text{D.F.} = & 5.9 \text{ MEF}^{40}(\text{P})(\text{He}) - 4.0 \text{ MEF}^{40}(\text{P})(\text{Air}) - 2.7 \text{ MEF}^{25}(\text{P})(\text{He}) \dots \\ & - 1.2 \text{ MEF}^{40}(\text{P})(\text{He}/\text{A}) + 0.7 \text{ IFVP} + 0.3 \text{ MEF}^{25}(\text{P})(\text{He}/\text{A}) \dots \\ & - 1.3 \text{ CV} - 0.05 \text{ MEF}^{25}(\text{P})(\text{Air}) \end{aligned}$$

When the mean values of the indices are inserted for each of the four groups namely female non-smokers, female smokers, male non-smokers and male smokers, the equation above takes the following four values:-

$$23.87; \quad 31.45; \quad 14.42 \text{ and } 16.7^{\frac{1}{2}}$$

The observed values of the indices for subjects 1 and 2 for that session when substituted in the D.F. gave 65.47 and 15.40. Now since the D.F. for female smokers is 31.47 and that for male non-smokers 14.42, subject No. 1 is classified as a female smoker whilst subject No. 2 is classified as a male non-smoker. These classifications are in fact correct.

Thus for the Ach session a female non-smoker would be expected to have D.F. values between 23.87 and 31.45; and her smoker counterpart a value greater than 31.45. For the male non-smoker a value between 14.42 and 23.87 would be expected and his smoker counterpart should have a value between 14.42 and 16.7 $\frac{1}{2}$.

D.F.s were obtained from the data of each session and the coefficients were ranked in ascending order of magnitude. Indices obtained at sessions

where the treatment was a bronchoconstrictor were ranked separately to the sessions where the treatment was a bronchodilator. The resulting rank order of the indices was tested on the t distribution. Differences in rank order between sessions were not significant at the 5% level.

By summing the ranks of the indices a measure of the specificity with which an index detected differences in the baseline data of smokers and non-smokers was obtained. Two separate specificity orders were compiled, one for the "maximal" and the other for the "partial" indices. The specificity orders were:-

- a) (i) MEF⁴⁰(P)(Air) ii) MEF⁴⁰(P)(He) iii) MEF²⁵(P)(He)
(iv) CV and MEF²⁵(P)(Air) jointly, v) IFVP vi) MEF⁴⁰(P)(He/A)
and vii) MEF²⁵(P)(He/A).
- b) i) CV ii) MEF⁴⁰(He) iii) CV/SVC(%) iv) MEF²⁵(Air) and
MEF⁴⁰(Air) jointly, v) MEF²⁵(He) vi) SVC vii) FEV₁
viii) FEV₁/FVC(%) and ix) MEF⁴⁰(He/A) and IFVP jointly.

At least three D.F.s are required to classify data from four distinct groups; the efficiency of correct classification is summarised in Table 8.

Sensitivity: Baseline data

The coefficient of variation (COV) is the standard deviation expressed as a percentage of the mean value. The COV of each index of pulmonary function was calculated from the pooled baseline data from each treatment session, and ranked in ascending order of magnitude. Indices obtained from "partial" and "maximal" manoeuvres were treated separately, two sessions each contributing to the mean. The between sessions rank order was not significantly different ($r_s = 0.67$, $t = 2.21$)

and $r_s = 0.49$, $t = 1.86$) respectively for "partial" or "maximal" indices.

The magnitude of the summed ranks of COV were then used to arrange the indices in order of sensitivity (low COV being equated to high sensitivity). The following is the resulting order of sensitivities:-

"Partials":- i) MEF⁴⁰(P)(He) ii) MEF²⁵(P)(He) jointly with MEF⁴⁰(P)(Air) iii) MEF⁴⁰(P)(He/A) iv) MEF²⁵(P)(Air) v) CV jointly with MEF²⁵(P)(He/A) and vi) IFVP.

"Maximals":- i) FEV₁/FVC(%), ii) FEV₁ iii) FVC jointly with MEF⁴⁰(Air), iv) SVC v) CV/SVC(%) vi) MEF²⁵(Air), vii) MEF⁴⁰(He/A) jointly with IFVP viii) MEF²⁵(He) ix) CV and MEF⁴⁰(He) jointly and x) MEF²⁵(He/A).

A linear combination of the specificity and sensitivity orders was used to rate the indices (low rating equated with better indices). On the baseline data the indices were rated as follows:-

"Partials":- i) MEF⁴⁰(P)(Air) jointly with MEF⁴⁰(P)(He)
ii) MEF²⁵(P)(He) iii) MEF²⁵(P)(Air) iv) MEF⁴⁰(P)(He/A)
jointly with CV and v) MEF²⁵(P)(He/A) jointly with IFVP.

"Maximals":- i) MEF⁴⁰(Air) ii) CV/SVC(%) iii) FEV₁ jointly with FEV₁/FVC(%) iv) MEF²⁵(Air), CV, and SVC jointly
v) MEF⁴⁰(He) vi) MEF²⁵(He) vii) FVC viii) MEF⁴⁰(He/A)
ix) IFVP and x) MEF²⁵(He/A).

EFFECT OF DRUGS

The effect of the inhaled bronchoactive drugs, (Ach., PGF_{2α} Ventolin and Sal 1000), was measured by the change in each pulmonary

function index from its average baseline value. In the case of some drugs (like Ach) the expected time of duration of bronchoactivity is similar to the time needed to execute all the test manoeuvres. In these circumstances serial tests may not necessarily be replicates.

The change was therefore determined as:-

- i) Maximum...the largest difference between average baseline and post-treatment values.
- ii) Mean...the difference between the average baseline value and the arithmetic mean of the post-treatment values.

Analysis of variance of the mean and maximum changes obtained from the pooled sessional data showed that the two changes were not significantly different at the 5% probability level, (F values were 2.21, 2.85, 1.02 and 0.55 for Ach., PGF_{2α}, Ventolin and Sch1000 sessions respectively). However, t test showed that maximum changes in MEF25(P)(Air) and MEF25(P)(He) were significant at the 1% probability level and mean changes were significant at only the 5% level. Maximum changes may therefore be more efficient in detecting drug response.

Maximum changes were used to assess relative sensitivity and specificity of the indices in detection of the drug induced reversible airway changes. The average maximum changes for each drug are shown in table 9.

Following provocation by Ach., IFVP rose but not significantly ($P>0.05$), all the other indices (MEF25(P)(Air) to MEF₂₅O(P)(He/A) inclusive fell; falls in MEF25(P)(Air) and MEF25(P)(He) were significant by t test, ($P<0.01$ and 0.001 respectively).

After PGF_{2α}, there were rises in MEF25(P)(He/A) and IFVP (both

non significant with $P > 0.05$), and falls in the remaining indices; the falls in MEF_{25(P)}(Air), MEF_{40(P)}(Air) and MEF_{40(P)}(He) were significant ($P < 0.05$, 0.05 and 0.01) respectively.

Inhalation of Ventolin resulted in lowered values of 3 indices and increased values in the remainder; the increases in 7 (including IFVP) were significant, $P < 0.05$. Only the change in MEF_{40(He)} following Sch1000 reached the 5% significance level. The falls in CV, FEV₁ and CV/SVC were small and insignificant.

Drug Doses

All subjects were given 2 puffs of Ventolin (200 μ g). The doses of Ach₄, PGF_{2 α} and Sch1000 given from a Wright's nebuliser were ranked in order of magnitude. Smokers and non-smokers received similar doses (rank order difference not significant at 5% probability level).

Specificity: Drug induced changes

Data of maximum change after each drug were used to calculate Discriminant Functions for the various subgroups of subjects. The D.F. equations for each session were:-

i) Ach.

$$\begin{aligned} \text{D.F.} = & -1.70 \text{ MEF}^{40}(\text{P})(\text{He}/\text{A}) + 1.46 \text{ MEF}^{40}(\text{P})(\text{He}) + 1.05 \text{ IFVP...} \\ & -0.74 \text{ MEF}^{25}(\text{P})(\text{Air}) + 0.61 \text{ MEF}^{40}(\text{P})(\text{Air}) - 0.41 \text{ MEF}^{25}(\text{P})(\text{He})... \\ & + 0.36 \text{ MEF}^{25}(\text{P})(\text{He}/\text{A}) + 0.14 \text{ CV}. \end{aligned}$$

ii) PGF_{2 α}

$$\begin{aligned} \text{D.F.} = & -2.96 \text{ MEF}^{25}(\text{P})(\text{Air}) - 1.76 \text{ MEF}^{25}(\text{P})(\text{He}) - 1.57 \text{ MEF}^{40}(\text{P}) \\ & (\text{Air}) - 0.83 \text{ IFVP} + 0.77 \text{ MEF}^{25}(\text{P})(\text{He}/\text{A}) - 0.68 \text{ CV} - 0.30 \text{ MEF}^{40}(\text{P}) \\ & (\text{He}) - 0.13 \text{ MEF}^{40}(\text{P})(\text{He}/\text{A}). \end{aligned}$$

iii) Sch1000

$$\text{D.F.} = 8.24 \text{ FEV}_1 - 6.06 \text{ FEV}_1/\text{FVC}(\%) + 6.04 \text{ MEF}^{25}(\text{He}/\text{A})...$$

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$$\rightarrow 5.83 \text{ MEF}^{\text{lo}}\text{O(Air)} + 5.78 \text{ MEF}^{\text{lo}}\text{O(He/A)} - 4.02 \text{ MEF}^{\text{lo}}\text{O(He)} - 3.32 \text{ CV} \dots$$

$$- 3.03 \text{ FVC} - 2.60 \text{ MEF}^{\text{lo}}\text{O(He)} - 2.07 \text{ MEF}^{\text{lo}}\text{O(Air)} + 1.61 \text{ IFVP} \dots$$

$$- 0.64 \text{ SVC.}$$

The D.F. soefficients were ranked in ascending order of magnitude.

Rank correlation coefficients (r_s) were calculated for rank order differences between the types of drug (constrictors and dilators and the summed ranks of dilators and constrictors used to arrange the indices in specificity order (high specificity equated with low values of summed ranks). The rank order for the bronchoconstrictors was not significantly different ($r_s = -.33$, $t = 0.87$), but it was different for bronchodilators ($r_s = -0.54$, $t = 2.12$, $P > 0.05$).

Order of specificity were:-

"Partials":- i) MEF^{lo}O(P)Air) ii) IFVP iii) MEF^{lo}O(P)(He) and MEF^{lo}O(P)Air) jointly, iv) MEF^{lo}O(P)(He/A) jointly with MEF^{lo}O(P)(He) v) MEF^{lo}O(P)(He/A) and vi) CV.

"Maximals":- i) MEF^{lo}O(He) ii) MEF^{lo}O(Air) and FEV₁ jointly iii) MEF^{lo}O(He), IFVP and CV conjointly, iv) MEF^{lo}O(He/A) v) MEF^{lo}O(Air) vi) MEF^{lo}O(He/A) jointly with FEV₁/FVC(%) vii) (CV/SVC jointly with FVC) and viii) SVC.

Sensitivity: Drug induced changes

The COV of the maximum post drug changes were calculated from the pooled data (all subjects) and the sensitivity of the indices determined in similar fashion to that used for baseline data. Rank order for bronchoconstrictors ($r_s = 0.52$, $t = 1.49$) and for bronchodilators ($r_s = 0.28$, $t = 0.97$.) were not different for drug types. The order of sensitivity for the indices were:-

"Partials":- i) MEF⁴⁰(P)(Air) ii) MEF⁴⁰(P)(He) iii) MEF²⁵(P)
(Air) jointly with MEF²⁵(P)(He) iv) IFVP and MEF²⁵(P)(He/A)
v) MEF⁴⁰(P)(He/A) and vi) CV ;

"Maximals":- i) MEF⁴⁰(He) ii) MEF²⁵(He) iii) MEF²⁵(Air)
iv) MEF⁴⁰(Air), CV and IFVP conjointly, v) FVC vi) CV/SVC
vii) SVC and MEF²⁵(He/A) viii) FEV₁ and FEV₁/FVC(%) and
ix) MEF⁴⁰(He/A).

Rating: Drug induced changes

A linear combination of the sensitivity and specificity order of each pulmonary function index was used to give the following ratings.

"Partials":- i) MEF²⁵(P)(Air and MEF⁴⁰(P)(Air) ii) MEF²⁵(P)(He),
MEF⁴⁰(P)(He) and IFVP, iii) MEF²⁵(P)(He/A) and MEF⁴⁰(P)(He/A)
and iv) CV.

"Maximals":- i) MEF⁴⁰(He) ii) MEF²⁵(He) iii) MEF⁴⁰(Air)
iv) IFVP and CV v) MEF²⁵(Air) vi) FEV₁ vii) FVC viii)
CV/SVC, MEF⁴⁰(He/A) and MEF²⁵(He/A) ix) FEV₁/FVC(%) and
x) SVC.

Table 10(b) lists the rating of each index, according to ability to detect drug induced reversible changes.

FACTOR ANALYSIS

Factor analysis seeks to establish groups of test variables which measure the same attribute in a group of persons for whom measurements of several test variables are available. Based on the correlation between test variables (pulmonary function indices) new variables ("factors") are created. A "factor" is a linear equation of the test

variables with an additional term which takes account of the random error in the data set. The coefficients of a Factor are chosen such that each Factor has a variance of unity, but account for a proportion of the total variability in the data. Principal Factors (PF), are combinations of Factors which are fewer than the original test variable but account for most of the variability of the data set.

The SPSS computer program Factor Analysis (F.A.) uses Varimax rotation for extraction of PF and was used to analyse data from smokers and non-smokers in this study. In the analysis no account was taken of the differences between indices obtained from "partial" and "maximal" manoeuvres, for example MEF25(P) was not distinguished from MEF25, since both of these measure the same attribute namely flowrate at 25%VC.

The data from non-smokers yielded 5 PFs which accounted for 35.4%, 23.3%, 19.9%, 14.4% and 7% respectively of the data's total variability. Four PFs which accounted for 41.8%, 29.8%, 15.5% and 12.9% respectively were extracted from the smokers' data. The first three PFs from each data set account for more than 75% of the variation in each set (87.1% and 78.6% for smokers and non-smokers respectively). The major components (largest first) of the first three PFs were:-

Non-smokers

- i) (MEF¹O(He), MEF25(He), MEF25(He/A)) ii) (FEV₁/FVC(%), MEF¹O(Air), CV), and iii) (MEF25(Air), CV/SVC, FEV₁).

Smokers

- i) (CV, MEF¹O(He/A), SVC), ii) (MEF¹O(He), MEF25(He), MEF25(Air)) and iii) (FVC, FEV₁, CV/SVC).

In this way the data has been reduced to emphasize the indices which

measure similar types of change in function in both smokers and non-smokers.

DISCUSSION

To evaluate tests of small airway defect subjects are needed who are known to have defective small airways. The defect in small airways can be present alone or in company with defect of the large airways. When both types of defect are present in the same subject, means of measuring and/or allowing for the effect of the large airway defect must be available before the two types of defect can be separated.

Smokers are said to have small airway disease (McFadden, 1972), and cessation of the smoking habit has been shown to lead to improvement in small airway function (McFadden, 1972). Although not ideal smokers can therefore be used as a model of small airway defect.

Provocation tests have previously been used to measure bronchial reactivity in healthy and diseased lungs (Cade, 1971) and in normal controls and patients (Bouhuys, 1969). For this purpose drug doses are generally chosen which produce a minimum response level in the subject. For example, inhalation of a drug of known concentration may be continued until the subject's flow rate changes by "X"%. Another method is to give each subject the same "large" dose of a drug and note the subsequent change in a given index of function. An example of this method is the use of a bronchodilator to assess the reversibility of obstruction in patients.

In this study both the sensitivity and specificity of several indices of small airway function were to be assessed. To assess sensitivity (the ability to detect change) a continuum of sizes of

bronchial response (from zero through to maximum) was desired in order to give both the most sensitive and least sensitive indices a chance to detect a change in airway function. Specificity, (ability to distinguish between types of defect), for its assessment (in the absence of precise knowledge of the site and magnitude of airway defect) required a minimum of two groups of persons distinguishable by variables distinct from those of lung function. These two groups should be expected apriori to have different incidence of small airway disease as for example may be the case when lifelong non-smokers are compared to current and ex-smokers. Superimposing drug-induced reversible functional change upon the pre-existing small airway defect incidence in the two groups, enabled the specificity of the indices to reversible functional changes to be compiled.

The volunteers studied here were healthy, none of them had been diagnosed as suffering from ventilatory impairment and they had no complaints referable to chest disease. Their mean baseline values in the various indices (Table 2) show slightly better small airway function in the non-smokers (higher MEF values) compared to the smokers. The more conventional ventilatory indices (FEV_1 and $FEV/FVC(\%)$) had the same mean value for both groups. Note also that the mean physical variables are almost identical, thus eliminating the need for internal standardisation within the groups.

The challenge to the airways was via inhaled bronchoactive drugs, the particle size and mode of inhalation being such that all sizes of airways were exposed. The two distinct doses of each type of drug (constrictors and dilators) in combination with possible differences in airway reactivity in smokers and non-smokers should have ensured a

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very wide range of responses. The use of two different bronchodilator (and bronchoconstrictor) drugs, should have confounded mode of action of drugs as a variable in the study, as each volunteer acted as his own control. The Ach. dose was approximately $1/_{10}$ of Bouhuys' (1969) midrange dose; that for PGF 2α was the midrange dose of Smith (1975) and the Sch1000 dose was $\frac{1}{4}$ of Francis' (1975) Ventolin was given in the standard dose used in pulmonary function laboratories for assessing reversibility of obstructive airway disease.

The indices compared are those in common use except for MEF 40 which was chosen instead of the more usual MEF 50 since it is less affected by instrumental artefacts and its use allows greater flexibility in the range of submaximal inspiratory volumes which are required for "partial" manoeuvres. Moreover the MEF 50 is highly correlated with MEF 40 ($r = 0.98$, $t = 21.36$, $P < 0.0001$): Regression equations relating the two are given elsewhere in this Thesis, Chapter 2 .

Full inspiration is known to reverse or at least reduce bronchoconstriction (Widdicombe, 1975). Because of this "partial" manoeuvres were used in this study to measure the responses of the constrictors Ach. and PGF 2α . Baseline values of "partial" indices are systematically different from those of "maximal" indices (Bouhuys and this Thesis). This is mainly due to the effect of full inflation in reducing bronchomotor tone (Vincent, 1970). The "partial" CV may additionally be affected by the dependence of CV on inspired volumes (Linn, 1973; Holtz, 1976). In the present study the dependence of CV on inspired volumes was not seen, there being only a small non-significant difference between mean CV for the control constrictor and dilator values (0.66, 0.65, 0.69 and 0.67 LBTPS for the Ach, PGF 2α , Ventolin and Sch1000

sessions respectively). A systematic difference is seen in that the "partial" CV means are consistently lower in value than the "full" CV means (0.66 ± 0.29 and 0.65 ± 0.25 compared to 0.69 ± 0.26 and 0.67 ± 0.29). However, these means are again indistinguishable statistically, indicating that in this study the systematic difference could not be separated from "circadian" or random variations. This finding agrees with that of Liang, (1973), who found no significant decrease in CV with inspired volume (V_I) but contrasts with that of Holtz, (1976) who found significant differences between CV measured at V_I between 75% and 100% VC.

All the indices compared in this study are determined by flow through the airways because measurements were made at the mouth. Among the many variables which affect flow through airways and hence the derived indices are calibre, length, muscle tone and thickness of mucus layer (Clarke, 1973). Since length of airways and the consistency of mucus lining are determined mainly by the physical size and the healthiness of the lung respectively, these variables should have remained fairly constant throughout this study. Airway calibre is affected by both the degree of lung inflation and lung volume history (Bouhuys, 1967). Since special care was taken in this study to ensure constant lung volume history and degree of lung inflation, change in airway calibre by drugs should provide a valid basis for comparison of the indices.

Baseline data were collected from the subjects in this study under conditions which:-

- (i) Allowed differences of airway calibre to be measured in the presence of normal airway smooth muscle tone ("partial" indices).

(iii) Allowed differences of calibre to be measured in the presence of reduced tone (maximal indices).

Condition (ii) follows because full inflation reduces bronchomotor tone (Vincent, 1970).

In healthy persons, the role of airway smooth muscle tone may be of marginal significance for example in regulating gas distribution, its relaxation causing subtle changes but no gross impairment of lung function (Bouhuys, 1974a). However, since markedly nonuniform distribution of inspired gas, as defined by the single-breath nitrogen test, (a test very similar to the CV test), is reported to be found only in older persons and in patients with lung disease (Bouhuys, 1974b), change in tone in smokers may be important in revealing early lung disease. If the CV test is capable of detecting differences in calibre in the presence of normal and reduced tone, then indices derived from it should be well rated in the baseline ratings of the indices derived in this study since difference in calibre is the common factor in the data. On the other hand, if the test is better at detecting other aspects of tonal change apart from calibre change (e.g. gas distribution) CV indices should rate better under condition (ii) than under (i) above. In the present study the baseline ratings suggest that MEFV curves may be better at detecting airway calibre changes than the CV test and there is confirmatory evidence from the ratings of the indices in relation to the drug induced changes. Reports from other workers (Macklem, 1972) indicate that both the CV test and Flow-volume curves can detect early signs of small airway disease. Some of the variation in the results obtained when smokers have been used as a model of small airway disease (McFadden, 1972; Ingram, 1971; McCarthy, 1972; Bode, 1975;

Martin, 1975) may however be better explained by difference in specificity of the indices, (as suggested in the present study), rather than by difference in sensitivity postulated by Macklem (1972). Some recognition of the different roles which Specificity and Sensitivity play in the rating of tests of small airway function was demonstrated by the participants in "WORKSHOP ON SCREENING PROGRAMS FOR EARLY DIAGNOSIS OF AIRWAY OBSTRUCTION" held at Virginia, U.S.A., 1975. They recommended Closing Volume as the "test of choice" since although it was not necessarily better than other tests, "more is known about its source of variance and the physiologic factors that influence it than most other tests".

In arriving at ratings for the pulmonary function indices compared in this study, equal weighting has been given to the two components (linear combination of measures of sensitivity and specificity). Sensitivity has been assessed in terms of the indices' COV and this is justified by the consideration that for many lung function indices the variability is proportional to the magnitude of the index (Cotes, 1975). Specificity has been determined on the basis of the indices' discrimination between distinct populations, for example smokers and non-smokers. The underlying assumption of the statistical technique of Discriminant Analysis is that the data should consist of random samples from normally distributed parent populations (Kendall, 1975). Since the smokers and non-smokers in this study were not selected by random sampling methods, the analysis of the resulting data may not have been strictly valid making statistical tests of the "goodness of fit" of the Discriminant Functions (DF) difficult to interpret. Indeed the DFs reported here may not be the best statistically, and the weightings of the indices may be in error. The

technique of ranking, being nonparametric, is largely independent of the form of the sample distribution (Hayslett, 197^h). If the Specificity calculated from DFs for smokers and non-smokers under similar conditions could be compared nonparametrically, it might be possible to detect large errors in the relative weightings of the indices. An opportunity to make this comparison was available in this study since the type of change which occurs in the baseline data of the maximal indices would be expected to differ in magnitude only from that which occurred after treatment with the bronchodilators. Rank order correlation of specificity was not significantly different when baseline data was compared to post-dilator data. For these comparisons, spearman correlation coefficient, r_s , was 0.53 and 0.37 for the post-ventolin and post-dilator (ventolin and Sch1000 summed) respectively; corresponding t values of 1.87 and 1.33 indicates that the r_s was not statistically different at 5% probability level. This finding implies that the weighting of the indices in the DFs were not grossly in error.

During deep and forced breathing, the calibre of the airways is a function of the compliance of the airway wall. In small airways, a change in smooth muscle tone may produce a different functional response to that which results from similar tonal changes in large airways, (Bouhuys, 197⁴). The overall response to bronchoconstrictors and bronchodilators found in this study suggests that the main region of the lungs affected by the drugs was in the small airways. Since factor analysis groups indices which measure similar attributes, the result of the analysis should highlight areas of the lungs where the drugs have been effective. Considering the non-smokers' data, the PF

which accounts for 35% of the total variability of the data, is heavily weighted in indices which reflect changes occurring at lung volumes at or below 60% VC. The equivalent PF for the smokers' data accounts for about 30% of the data's variability. Although the smokers' first PF accounts for more of the variation than the second, (42% compared to 35%), it is also heavily weighted in small airway indices (CV and MEF⁵⁰(He/A)). Factor analysis is therefore in agreement with the response suggestion; namely that the small airways were challenged by the inhaled drugs. Independent evidence from workers using gamma emitting radionuclide tagged particles show that inhaled particles can be deposited in peripheral lung airways. Using external radiation detectors positioned over the chest, several workers have shown that monodisperse solid particles have been deposited in the alveolar region (Thomson, 1974; Pavia, 1976; Lippman, 1971). By imaging the lung field on Gamma camera, 5^{um} monodisperse and heterodisperse (size range 0.2 to 2^{um}) particles have been shown to be present in lung periphery, (Lourenco 1971; Short 1978; Fazio 1977). Remembering that the drugs in this study were administered as heterodisperse droplets (0.8^{um} mmd), that less than 10% of inhaled particles are deposited in the entire respiratory tract (Connolly, 1971) and that the drug doses (with the exception of ventolin) were estimated by loss of weight from the nebulisers, only very small quantities of drug could have reached the small airways. The overall impression from this study is therefore that small airway tests are very sensitive indeed. This high sensitivity may in part be responsible for the large variability found when normal values for the indices are calculated. Large intersubject variability between times and intrasubject variations have been cited as evidence of limitations in epidemiology (Clark, 1977). Whilst it is true that the variability associated with flow volume curves is generally greater than those

found in the CV and the FEV₁ tests, this study indicates that indices derived from flow volume data may be highly specific to reversible changes.

Reversible changes are the only ones likely to benefit from medical intervention in the course of chronic obstructive disease. It may, therefore become worthwhile to consider ways of combining data from "partial" and "maximal" manoeuvres, to reveal the range of tonal variation present in health and disease. This range of variation may even be useful in narrowing the range of predicted "normal" values.

TABLE I

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PHYSICAL CHARACTERISTICS AND TOBACCO CONSUMPTION FOR TWENTY FIVE SUBJECTS

Subject	Sex	Age Yr	Height (m)	Weight (Kg)	Equivalent cigarette consumption Rate (Pk-Yr)*	Total No. x 10 ³
1 MAC	M	36	1.83	70.3		
2 GWC	M	42	1.74	72.0		
3 GVC	M	51	1.82	72.2		
4 DJD	M	39	1.84	75.0		
5 JDF	M	55	1.82	76.1		
6 MK	M	41	1.90	74.2		
7 HP	M	62	1.79	83.7		
8 JW	M	55	1.79	71.6		
9 ED	F	25	1.69	63.9		
10 LD	F	22	1.59	54.1		
11 MC	F	31	1.68	56.1		
12 JS	F	23	1.62	52.1		
13 SW	F	24	1.72	55.0		
14 DW	F	32	1.67	58.1		
15 CIE	M	41	1.88	73.1	0.36	45.6
16 KLK	M	39	1.73	68.1	0.36	49.5
17 GM	M	41	1.75	84.4	2.0	306.6
18 AP	M	56	1.80	74.6	0.90	249.7
19 MS	M	34	1.81	74.0	0.33	38.6
20 PW	M	64	1.71	77.4	0.21	70.6
21 AZ	M	31	1.80	70.1	0.51	47.5
22 HG	F	22	1.59	66.1	0.93	27.4
23 JK	F	31	1.70	66.7	0.70	25.6
24 ET	F	26	1.69	49.0	0.37	21.5
25 PB	F	22	1.65	60.9	1.0	43.8

* 1 pk-yr = 7.3×10^3 cigarettes in 1 year

TABLE 2a

GROUP CHARACTERISTICS: 14 NON-SMOKERS AND 11 SMOKERS

Physical:- Mean (S.D.)

Non-smokers: Ht.(m) 1.75(0.09); Wt.(Kg) 66.6(9.8); Age(Yr) 39.1(12.3)

Smokers: Ht.(m) 1.74(0.08); Wt.(Kg) 70.0(8.5); Age(Yr) 37.4(13.0)

Pulmonary Function Indices: Mean (SE_m)

	NON-SMOKERS	SMOKERS	
MEF25(Air)	144 (9.3)	122 (9.0)	L/min.
MEF25(He)	169 (11)	142 (12)	L/min.
MEF ¹⁴ O(Air)	240 (13)	223 (14)	L/min.
MEF ¹⁴ O(He)	306 (16)	287 (19)	L/min.
CV	0.68 (.04)	0.64 (.04)	L BTPS
SVC	4.23 (.23)	3.89 (.36)	L BTPS
CV/SVC	15.9 (.97)	16.9 (1.4)	%
FEV ₁	3.94 (.18)	3.75 (.21)	L BTPS
FVC	5.16 (.28)	4.65 (.23)	L BTPS
FEV ₁ /FVC	77.4 (2.2)	79.6 (1.8)	%
MEF25(He/A)	0.28 (.02)	0.32 (.03)	
MEF ¹⁴ O(He/A)	0.36 (.02)	0.38 (.04)	
IFVP	19.3 (1.7)	25.9 (3.1)	% VC

TABLE 2b

GROUP CHARACTERISTICS: 15 MALES AND 10 FEMALES

Physical:- Mean (S.D.)

Males:- Ht.(m) 1.80(0.05); Wt.(Kg) 74.5(4.5); Age(Yr) 45.9(10.2)
 Smoking (Cigarettes smoked $\times 10^3$) 115.4(106.3)

Females:- Ht.(m) 1.66(0.04); Wt.(Kg) 58.2(5.6); Age(Yr) 26.7(4.1)
 Smoking (Cigarettes smoked $\times 10^3$) 30.4(9.0)

Pulmonary Function Indices: Mean (SE_m)

	MALES	FEMALES	
MEF25(Air)	137 (9.7)	131 (7.5)	L/min.
MEF25(He)	164 (11.1)	163 (10.1)	L/min.
MEF ¹⁰ O(Air)	249 (14.1)	214 (10.1)	L/min.
MEF ¹⁰ O (He)	317 (18.1)	269 (11.1)	L/min.
CV	0.80 (0.03)	0.64 (0.02)	L BTPS
SVC	4.58 (0.19)	3.39 (0.20)	L BTPS
CV/SVC	18.3 (0.95)	13.1 (1.1)	%
FEV ₁	4.33 (0.15)	3.31 (0.16)	L BTPS
FVC	5.60 (0.23)	4.06 (0.18)	L BTPS
FEV ₁ /FVC	74.8 (2.1)	82.6 (1.6)	%
MEF25(He/A)	0.28 (0.05)	0.32 (0.02)	
MEF ¹⁰ O(He/A)	0.40 (0.03)	0.32 (0.02)	
IFVP	20.0 (1.1)	25.7 (4.1)	SVC

TABLE 7

BASELINE DATA: "PARTIAL" AND "MAXIMAL" MANOEUVRES.

MEAN SESSIONAL VALUES FOR NON-SMOKERS AND SMOKERS.

	NON-SMOKERS				SMOKERS				L/min.
	Mean	(SE _m)	Mean	(SE _m)					
MEF ₂₅ (P)(Air)	1 ^{b7}	(19)	135	(17)	133	(17)	116	(14)	"
MEF ₂₅ (Air)	127	(15)	121	(12)	122	(13)	102	(8)	"
MEF ₂₅ (P)(He)	173	(16)	16 ^{b4}	(18)	16 ^{b4}	(25)	14 ^{b7}	(18)	"
MEF ₂₅ (He)	14 ^{b5}	(16)	14 ^{b6}	(15)	14 ^{b1}	(22)	123	(14)	"
MEF ^b O(P)(Air)	259	(31)	218	(23)	238	(26)	223	(28)	"
MEF ^b O(Air)	218	(21)	218	(20)	216	(17)	214	(18)	"
MEF ^b O(P)(He)	317	(30)	305	(29)	30 ^{b4}	(33)	306	(32)	"
MEF ^b O(He)	291	(29)	290	(29)	280	(26)	277	(31)	"
CV (P)	0.71	(.08)	0.57	(.09)	0.56	(.08)	0.58	(.10)	L BTP
CV	0.74	(.08)	0.63	(.12)	0.63	(.07)	0.57	(.10)	"

* "Partial" manoeuvres were performed at two sessions and "maximal" at two sessions; each column relates to a session.

TABLE 1

BASELINE DATA: MULTIPLE CORRELATION COEFFICIENTS (R^2), SIMPLE CORRELATION COEFFICIENTS (r) AND SIGNIFICANCE OF REGRESSION.

Females

	HEIGHT		WEIGHT		AGE		SMOKING	
	R^2	r	R^2	r	R^2	r	R^2	r
MEF25(Air)	0.01	0.10	0.01	-0.06	0.07	-0.22	0.72	0.75
MEF25(He)	0.01	0.09	0.02	-0.12	0.16	-0.34	0.76	0.77
MEF40(Air)	0.01	0.12	0.02	0.03	0.05	-0.18	0.37	0.54
MEF40(He)	0.01	0.12	0.03	-0.12	0.11	-0.26	0.61	0.67
MEF25(He/A)	0.03	0.18	0.06	0.14	0.12	-0.24	0.22	0.39
MEF40(He/A)	0.01	0.11	0.03	-0.15	0.07	-0.17	0.42	0.53
CV	0.05	0.23	0.32	0.51	0.38	0.20	0.57	-0.42
SVC	0.22	0.47	0.36	0.36	0.38	-0.15	0.39	-0.02
FEV ₁	0.35	0.56	0.37	0.14	0.43	-0.13	0.72	-0.41
FVC	0.30	0.55	0.42	0.33	0.44	-0.12	1.0	-0.92
CV/SVC(%)	0.01	0.09	0.15	0.37	0.47	0.52	0.47	-0.25
FEV ₁ /FVC(%)	0.02	0.13	0.29	-0.53	0.30	-0.04	0.82	0.51
IFVP	0.09	-0.30	0.17	0.29	0.19	-0.22	0.32	0.74

TABLE 5

BASELINE DATA: MULTIPLE CORRELATION COEFFICIENTS (R), SIMPLE CORRELATION COEFFICIENTS (r) AND SIGNIFICANCE OF THE REGRESSION.

<u>Males</u>	HEIGHT		WEIGHT		AGE		SMOKING	
	R ²	r						
MEF25(Air)	0.03	0.16	0.10	-0.31	0.12	-0.23	0.13	-0.27
MEF25(He)	0.03	0.18	0.12	-0.33	0.14	-0.30	0.15	-0.27
MEF ¹⁴ O(Air)	0.10	0.32	0.14	-0.28	0.20	-0.39	0.22	-0.32
MEF ¹⁴ O(He)	0.12	0.35	0.20	-0.36	0.27	-0.46	0.30	-0.39
MEF25(He/A)	0.00	-0.01	0.00	-0.05	0.01	-0.07	0.01	-0.01
MEF ¹⁴ O(He/A)	0.01	-0.01	0.03	-0.22	0.08	-0.05	0.13	0.07
IFVP	0.01	-0.21	0.03	0.23	0.22	0.47	0.27	0.30
CV	0.03	0.17	0.06	0.13	0.11	0.20	0.42	0.42
SVC	0.06	0.25	0.20	-0.42	0.28	-0.46	0.46	-0.59
CV/SVC	0.00	0.07	0.19	0.40	0.35	0.48	0.29	0.60
FEV ₁	0.02	0.13	0.18	-0.43	0.42	-0.61	0.69	-0.63
FVC	0.01	0.07	0.03	-0.39	0.10	-0.26	0.80	-0.75
FEV ₁ /FVC	0.10	0.31	0.10	-0.10	0.37	-0.57	0.43	-0.23

TABLE 6

BASELINE DATA: MULTIPLE CORRELATION COEFFICIENTS (R), SIMPLE CORRELATION COEFFICIENTS (r) AND SIGNIFICANCE OF THE REGRESSION.

<u>NON-SMOKERS</u>	HEIGHT		WEIGHT		AGE	
	R ²	r	R ²	r	R ²	r
MEF25(Air)	0.04	0.20	0.30	-0.37	0.32	-0.05
MEF25(He)	0.01	0.09	0.30	-0.15	0.30	-0.20
MEF ¹ O(Air)	0.13	0.36	0.37	-0.29	0.38	0.02
MEF ¹ O(He)	0.13	0.37	0.35	-0.27	0.36	-0.05
MEF25(He/A)	0.00	0.03	0.05	-0.19	0.08	-0.22
MEF ¹ O(He/A)	0.04	0.21	0.05	0.18	0.06	0.09
IFVP	0.18	-0.43	0.19	-0.14	0.19	-0.14
CV	0.18	0.43	0.42	0.62	0.43	0.51
SVC	0.29	0.54	0.32	0.09	0.34	0.28
CV/SVC	0.10	0.31	0.40	0.63	0.41	0.33
FEV ₁	0.37	0.61	0.38	0.16	0.38	0.28
FEV ₁ /FVC	0.00	0.05	0.12	-0.29	0.18	-0.33

TABLE 6

BASELINE DATA: MULTIPLE CORRELATION COEFFICIENTS (R), SIMPLE CORRELATION COEFFICIENTS (r) AND SIGNIFICANCE OF THE REGRESSION.

<u>NON-SMOKERS</u>	HEIGHT		WEIGHT		AGE	
	R ²	r	R ²	r	R ²	r
MEF25(Air)	0.04	0.20	0.30	-0.37	0.32	-0.05
MEF25(He)	0.01	0.09	0.30	-0.15	0.30	-0.20
MEF40(Air)	0.13	0.36	0.37	-0.29	0.38	0.02
MEF40(He)	0.13	0.37	0.35	-0.27	0.36	-0.03
MEF25(He/A)	0.00	0.03	0.05	-0.19	0.08	-0.22
MEF40(He/A)	0.04	0.21	0.05	0.18	0.06	0.09
IFVP	0.18	-0.43	0.19	-0.14	0.19	-0.14
CV	0.18	0.43	0.42	0.62	0.43	0.51
SVC	0.29	0.54	0.32	0.09	0.34	0.28
CV/SVC	0.10	0.31	0.40	0.63	0.41	0.33
FEV ₁	0.37	0.61	0.38	0.16	0.38	0.28
FEV ₁ /FVC	0.00	0.05	0.12	-0.29	0.18	-0.33

TABLE 7

BASELINE DATA: MULTIPLE CORRELATION COEFFICIENTS (R), SIMPLE CORRELATION COEFFICIENTS (r) AND SIGNIFICANCE OF THE REGRESSION.

<u>SMOKERS</u>	HEIGHT		WEIGHT		AGE	
	R ²	r	R ²	r	R ²	r
MEF25(Air)	0.00	0.06	0.01	-0.02	0.04	-0.04
MEF25(He)	0.00	-0.01	0.02	-0.01	0.04	-0.12
MEF ¹⁴ O(Air)	0.07	0.26	0.09	0.05	0.09	0.14
MEF ¹⁴ O(He)	0.09	0.30	0.15	0.01	0.15	0.13
MEF25(He/A)	0.02	-0.13	0.02	-0.11	0.12	-0.01
MEF ¹⁴ O(He/A)	0.02	0.13	0.02	0.07	0.02	0.13
IFVP	0.01	-0.09	0.04	0.09	0.04	0.02
CV	0.63	0.79	0.65	0.62	0.74	0.83
SVC	0.63	0.65	0.45	0.29	0.52	0.57
CV/SVC	0.31	0.55	0.47	0.66	0.64	0.78
FEV ₁	0.49	0.70	0.56	0.23	0.73	0.61
FVC	0.42	0.65	0.42	0.40	0.55	0.66
FEV ₁ /FVC	0.18	-0.42	0.49	-0.70	0.49	-0.60

TABLE 7

BASELINE DATA: MULTIPLE CORRELATION COEFFICIENTS (R^2), SIMPLE CORRELATION COEFFICIENTS (r) AND SIGNIFICANCE OF THE REGRESSION.

<u>SMOKERS</u>	HEIGHT		WEIGHT		AGE	
	R^2	r	R^2	r	R^2	r
MEF25(Air)	0.00	0.06	0.01	-0.02	0.04	-0.04
MEF25(He)	0.00	-0.01	0.02	-0.01	0.04	-0.12
MEF Δ O(Air)	0.07	0.26	0.09	0.05	0.09	0.14
MEF Δ O(He)	0.09	0.30	0.15	0.01	0.15	0.13
MEF25(He/A)	0.02	-0.13	0.02	-0.11	0.12	-0.01
MEF Δ O(He/A)	0.02	0.13	0.02	0.07	0.02	0.13
IFVP	0.01	-0.09	0.04	0.09	0.04	0.02
CV	0.63	0.79	0.65	0.62	0.74	0.83
SVC	0.43	0.65	0.45	0.29	0.52	0.57
CV/SVC	0.31	0.55	0.47	0.66	0.64	0.78
FEV ₁	0.49	0.70	0.56	0.23	0.73	0.61
FVC	0.42	0.65	0.42	0.40	0.55	0.66
FEV ₁ /FVC	0.18	-0.42	0.49	-0.70	0.49	-0.60

TABLE 3

THE PERCENTAGES OF VOLUNTEERS IN EACH OF FOUR GROUPS WHICH WERE
CORRECTLY CLASSIFIED USING DISCRIMINANT FUNCTIONS DERIVED FROM
DATA BEFORE (BASELINE, B) AND AFTER TREATMENT (DRUG, D).

SESSION/GROUP	FEMALE NON-SMOKER	FEMALE SMOKER	MALE NON- SMOKER	MALE SMOKER	BASELINE = B, AFTER DRUG = D.
Ach	83.3%	50%	87.5%	57.1%	B
	50%	75%	87.5%	57.1%	D
PGF _{2α}	66.7%	100%	85.7%	42.9%	B
	66.7%	100%	71.4%	71.4%	D
Sch1000	66.7%	75%	100%	71.4%	B
	83.3%	100%	100%	57.1%	D
VENTOLIN	100%	75%	50%	57.1%	B
	83.3%	75%	50%	42.9%	D

TABLE 9

THE AVERAGE OF THE MAXIMUM CHANGE FROM BASELINE VALUE OF PULMONARY
FUNCTION INDICES AFTER PROVOCATION BY EACH OF FOUR DRUGS.

	Mean \pm (SEM)	Ach.	PGF _{2α}	Ventolin	Schl000
MEF25(P)(Air)	-26.1 (13.3)*	-20.6 (9.6)*	-	-	-
MEF25 (Air)	-	-	20.0 (8.1)*	9.0 (6.2)	
MEF ¹⁴ O(P)(Air)	-39.8 (21.2)	-35.8 (13.2)*	-	-	-
MEF ¹⁴ O(Air)	-	-	41.8 (9.2)*	4.9 (9.5)	
MEF ¹⁴ O(P)(He)	-21.4 (22.1)	-59.4 (15.4)*	-	-	-
MEF ¹⁴ O(He)	-	-	51.4 (15.1)*	27.1 (13.7)*	
MEF25(He/A)(P)	-0.26 (0.24)	0.22 (0.24)	-	-	-
MEF25(He/A)	-	-	0.160 (0.13)	0.03 (0.05)	
MEF ¹⁴ O(He/A)(P)	-0.14 (0.25)	-0.03 (0.10)	-	-	-
MEF ¹⁴ O(He/A)	-	-	-0.004 (0.07)	0.05 (0.08)	
IFVP(P)	1.76 (6.1)	13.6 (7.4)	-	-	-
IFVP	-	-	13.7 (6.8)*	2.9 (4.6)	
CV (P)	-0.011 (0.05)	-0.032 (0.05)	-	-	-
CV	-	-	-0.061 (0.09)	-0.039 (0.85)	
MEF25(P)(He)	-40.2 (13.4)*	-23.6 (15.0)	-	-	-
	-	-	28.6 (8.2)*	10.1 (9.7)	
FEV ₁	-	-	0.046 (0.16)	-0.044 (1.04)	
FVC	-	-	-0.236 (0.45)	0.059 (0.89)	
FEV ₁ /FVC	-	-	1.2 (0.45)*	0.2 (1.55)	
CV/SVC	-	-	4.6 (1.02)*	-1.33 (1.31)	

* Significant at least the 5% level.

TABLE 10a

RATING OF INDICES IN ORDER OF ABILITY TO DETECT CHANGES IN
BASELINE VALUES.

Partial Indices

1. MEF40(P)(Air) and MEF40(P)(He)
2. MEF25(P)(He)
3. MEF25(P)(Air)
4. MEF40(P)(He/A) and CV
5. MEF25(P)(He/A) and IFVP

Maximal Indices

1. MEF40(Air)
2. CV/SVC%
3. FEV₁ and FEV₁/FVC%
4. MEF25(Air), CV and SVC
5. MEF40(He)
6. MEF25(He)
7. FVC
8. MEF40(He/A)
9. IFVP
10. MEF25(He/A)

TABLE 10b

RATING OF INDICES IN ORDER OF ABILITY TO DETECT DRUG INDUCED
REVERSIBLE CHANGES.

Partial Indices

1. MEF₂₅(P)(Air) and MEF¹⁰O(P)(Air)
2. MEF₂₅(P)(He), MEF¹⁰O(P)(He) and IFVP
3. MEF₂₅(P)(He/A) and MEF¹⁰O(P)(He/A)
4. CV

Maximal Indices

1. MEF¹⁰O(He)
2. MEF₂₅(He)
3. MEF¹⁰O(Air)
4. IFVP and CV
5. MEF₂₅(Air)
6. FEV₁
7. FVC
8. CV/SVC%; MEF¹⁰O(He/A) and MEF₂₅(He/A)
9. FEV₁/FVC(%)
10. SVC

C H A P T E R 3

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THE USE OF THE DEPOSITION OF RADIONUCLIDE TAGGED AEROSOLS TO MEASURE
THE FUNCTION OF HUMAN AIRWAYS.

INTRODUCTION

The deposition of particles in the human respiratory tract is known to be governed by three mechanisms. Altshuler (1960) stated that particles entrained in respired air are removed by:-

- a) Impaction the inertia of the particles carry them to airway wall when airflow direction changes. Deposition by impaction is therefore predominantly in large airways, for example, at the major bronchial bifurcations.
- b) Sedimentation under the effect of gravity particles settle out of the airstream and are deposited on airway walls. This mode of deposition would be expected to be prominent in airways of small internal diameter, where linear velocity is low.
- c) Brownian motion entrained particles experience random collision with gas molecules and drift towards airway walls. Removal of particles by this mode is generally small, taking place only where the mean free path is of similar order to molecular separation, as for example in very small airways in which airflow is almost stagnant.

In this study, the deposition of a radionuclide (Technetium 99m) tagged uniform sized polystyrene particles inhaled under controlled conditions was expressed in two ways. Total deposition in the lung was used to derive an index which reflects airway function. This index, the initial lateral penetration index (ILPI) is large for healthy airways reflecting relatively high deposition in the peripheral lung air spaces. Another index, the six hour retention value (SHRV), is based on particles

deposited on non-ciliated airways. Obstructed airways would be expected to yield low SHRV.

The deposition derived indices in eleven asymptomatic adults were compared with results of other pulmonary function tests including ; Closing Volume (a small airway test) and FEV_1 which is generally regarded as a large airway test.

SUBJECTS AND METHODS

Subjects

The eleven volunteers (7 males and 4 females) who took part in this study had volunteered for a trial of a new anticholinergic bronchodilator. They were all asymptomatic at the time of the study and gave no history of asthma or chronic bronchitis (Medical Research Council Questionnaire on Respiratory Symptoms, 1970). Three were lifelong non-smokers, 2 ex-smokers and the remaining 6 currently smoked. Table 1 gives the physical characteristics, tobacco consumption and the ventilatory capacities of the subjects. The predicted values were calculated on tests carried out in the month preceding the start of the study.

Methods

a) Pulmonary function

The methods and apparatus were previously described (Chapter 1). Maximal and Partial expiratory flow volume curves (MEFV and PEFV) were measured on the Mark 1 flow volume apparatus (Chapter 1); the peak expiratory flow rate (PEFR) and the flowrate at 50% of the vital capacity (MEF50) were obtained from the MEFV curves. The flowrate after 75% of the vital capacity had been expelled forcefully following a submaximal inspiration (MEF 25(P)) was measured on the PEFV curves. Closing volume was expressed as a percent of vital capacity (CV/VC%). FEV_1 and FVC were measured on a dry bellows spirometer (Drew and Hughes, 1969).

b) Radionuclide tagged aerosol tracer technique

The tracer technique used has been described by Thomson et.al. 1974. A monodisperse aerosol of polystyrene unleachably tagged with Technetium 99m(Few et.al. 1970) was generated in an airtight tank by a spinning disk (May, 1949). A wedge spirometer connected in series with the tank allowed the volume of aerosol withdrawn from the tank to be preset and automatically controlled by solenoid valves. A series connected pneumotachograph measured the flowrate with which air left the tank. A continuous recording of this flowrate (inspiratory flowrate (V_1)) was made on a chart recorder. A tank by-pass circuit enabled subjects to take alternate breaths from the tank (active breath) and from room air. The particles were of 5 μm ($\pm 0.8 \text{ SD}$) diameter.

The ^{140}O Kev gamma ray emitted by the isotope (TC-99m, half life six hours) was externally detected by scintillation detectors using sodium iodide (thallium activated) crystals 3.8 cm diameter and 2.5 cm thick. Lung burden assessment was obtained from two diametrically opposed probes centered on the chest anteriorly and posteriorly. The collimation of the probes was such that their field of view included most of the chest but very little of the stomach (Pavia et.al. 1971). Topographical distribution of the inhaled aerosol was determined from a posterior rectilinear scan. The locally built scanner (Dawson et.al. 1971) traversed vertical slices of the thorax in widths 1 inch (2.5 cm). The whole field was scanned in approximately half an hour, each traverse taking about three minutes. A continuous recording of the counts in each traverse was made on a potentiometric recorder fed from a ratemeter.

PROCEDURE

On three days, one week apart, each subject undertook an experimental run. A run consisted of:-

- a) pulmonary function tests and determination of background radiation counts over the chest in the hour immediately preceding
- b) aerosol inhalation
- c) initial lung burden was determined within three minutes of the start of inhalation. Assessment of the topographical distribution was begun about two minutes later.
- d) determination of the lung burden retained at six hours after aerosol inhalation

Pulmonary function tests were performed in the same order in each run. The first test was the partial expiratory flow volume curve from which MEF25(P) was obtained as the mean of the last three of five 'blows'. The reference vital capacity was taken as the mean of the last three (of five) blows on the maximum expiratory flow volume curves which followed the partial curves. A set of five maximum curves were completed within five minutes of completing a set of partial curves. The PEFR and the MEF50% was the mean of the values on the last three MEFV curves. The closing volume manoeuvre was next performed. CV/VC% was the mean of three technically competent manoeuvres. Finally FEV₁ was obtained as the mean of the last three of five attempts.

Aerosol inhalation consisted of the taking of eight breaths each of 500 ml volume from the tank. Nose clips were worn, there was no breath holding pause and each active breath was followed by one from the room air, each inspiration beginning at the FRC level. At the end of inhalation particles were cleared from the mouth, pharynx and oesophagus by a water mouthwash some of which was swallowed.

Initial lung burden was measured with the anterior probe centered on

the midsternum in the medial line and in close proximity to the body surface. Two counts each of 100 seconds duration were made. For each count the arithmetic mean of the two detectors was taken due allowance being made for differences in detector efficiencies determined beforehand. The initial lung burden was the mean of two counts.

An initial lateral penetration index (ILPI) was calculated from the topographical distribution of particles in the right lung. The right lung was selected since its field is less likely to be influenced by swallowed particles in the stomach. The total counts in each traverse was corrected for background and radioactivity decay. ILPI is the ratio of the summed counts in traverses 4 and 5 to the summed counts in traverses 1 and 2. Because each traverse width is 2.5 cm, two vertical 5 cm slices are used in calculating ILPI.

This method of calculating ILPI ensures a large value of the index when the particles are distributed in the lung periphery (patent smaller airways) and a small value when the particles are predominantly centrally deposited (obstructed small airways). The effect of taking narrower vertical slices is discussed later (see DISCUSSION).

At six hours after aerosol inhalation, the subject was repositioned as for the initial lung burden measurement and a single 100 second count taken. This count was decay corrected and expressed as a percentage of the initial lung burden. The value resulting is the six hour retention value (SHRV).

All subjects were closely observed during the entire experimental run. Smoking was not permitted for an hour before or during a run.

RESULTS

The mean value (the average of three runs), the standard error of

the mean (SEM) and the coefficient of variation (COV - the standard deviation expressed as a percentage of the mean) were calculated for all the parameters measured in an individual.

Flowrate during inhalation was analysed as maximum ($V_{I\max}$) and average (V_{Iavg}). The overall mean \pm SEM was $76 \pm 3 \text{ L min}^{-1}$ and $34 \pm 2 \text{ L min}^{-1}$ for $V_{I\max}$ and V_{Iavg} respectively. The corresponding COV were 15% and 12% indicating that inspiratory flowrates were highly reproducible overall. Analysis of variance confirmed that there were no significant difference in flowrates between the three runs. Inspiratory flowrate values are listed in Table 3.

Pulmonary function test results are shown in Table 2. The grand mean COV indicates that FEV₁ was the least variable test between runs, (COV 3.6%). Of the tests regarded as measuring small airway function. Closing volume (COV 11%) was less variable than (MEF25(P)) (COV 24%). The PEFR (COV 14%) and the MEF50 (COV 14.7%) were about equally variable in this study.

In Fig. 1 the mean observed CV/VC% (\pm 1SE) is shown for the 11 subjects. From a comparison with the predicted values of McCarthy et.al. (1972) whose predicted values are shown as the solid line \pm 2SE (dotted lines), none of the subjects gave evidence of small airway disease.

Fig. 2 shows the topographical distribution of the inhaled particles in the right lung of the volunteers for each of the three runs. The mean distribution was not significantly different on the 3 runs indicating no detectable change in the lung behaviour over the three weeks of study, since the physical variables (flowrate and volume) were unchanged.

Scatter diagrams showing the relationship of the initial lateral penetration index with lung function indices (CV/VC%, MEF25(P), MEF50,

FEV_1 and PEFR), and with age are given (Figs. 3 to 7). The expected positive correlations of ILPI with PEFR, FEV_1 and MEF50 were found as was the negative correlation with CV/VC%. The correlation coefficient (r) for these indices were 0.52, 0.57, 0.44 and -0.49 respectively. For MEF25(P) the theoretically expected positive correlation was demonstrated but the r value (0.23) was not as high as for the other function indices. ILPI was found to be independent of age. In view of the healthiness of the subjects and the small number in the group it is not surprising that none of the r values was significant at the 5% by t test.

ILPI was also found to be positively correlated with SHRV ($r = 0.11$), V_{Imax} ($r = 0.50$) and V_{Avg} ($r = 0.20$). The mean \pm SEM of ILPI in the lifelong non-smokers was 0.58 ± 0.10 with a mean COV of 29.5%. The smokers and ex-smokers gave the following corresponding results:- ILPI 0.69 ± 0.13 and COV of 52.7%. For this latter group the number of cigarettes smoked was negatively correlated with ILPI ($r = -0.355$ $n = 8$, $t = -0.93$). Again, the correlation coefficients were not statistically significant but indicate a trend.

The mean six hour retention value of $61\% \pm 3\%$ of initial lung burden is in accord with other authors (Pavia et.al. (1976) Short et.al. (1978)) and indicate that only about 40% of the particles were initially deposited on ciliated airways. The mean COV of the SHRV (11%) was much less than that for ILPI and might be explained in part by differences in individuals' clearance rate patterns. The SHRV was independent of inspiratory flow rates ($r = -0.005$ for V_{Imax}) and negatively correlated with age ($r = -0.28$). The non-smokers gave a mean SHRV of $59.7\% \pm 9.4\%$ and the smokers and ex-smokers $61.9\% \pm 3.0\%$.

DISCUSSION

Several workers (Thomson & Short, 1969; Lippmann et.al. 1970 Goldberg and Lourenco, 1973; Thomson and Pavia; 1974; Dolovich et.al. 1976; Short et.al. 1978) have shown that airway disease of both the restrictive and obstructive type cause premature particle deposition in the lung inhaled. Both monodisperse (Pavia et.al. 1977) and heterodisperse (Fazio, 1977) aerosols have been used to study lung deposition patterns. Pavia et.al. 1977) have pointed out that despite sophistication of aerosol inhalation technique particles will be deposited on either side of a target zone in the airways. Thus an aerosol chosen to give maximal deposition in large airways will also give some deposition in finer airways; some 10 μm particles have been shown to reach and be deposited in small airways (Thomson and Pavia, 1974).

From the topographical distribution of particles inhaled by the normal subjects in the present study, a 5 μm diameter particle appear to have an almost gaussian distribution along the lung airways. It is therefore assumed that differences in the individual's distribution is due largely to differences in the state of airway health and thus in airway patency. As an example consider two pairs of the subjects in this study. The first pair (females) consists of a lifelong non-smoker (No. 9) and a current smoker (No. 11). The smoker was older by 7 years (TLPI independent of age) taller by 2 cm. and had a predicted FEV₁, which although within the normal range, was 25% less than that of the non-smoker. Closing volume was marginally higher (by 3%) and flowrates at all lung volumes lower (MEF 25(P) by 12, MEF50 by 100 and PEFR by 27 L min^{-1}) in the smoker. Inhalation flowrates were higher by only 5 L min^{-1} in the smoker, but both TLPI and SHRV were substantially different. It is not thought that the 24% difference in SHRV is fully accounted for by impaction in central airways due to the increase of 5 L min^{-1} in

inspiratory flow rates. Notice also that the smoker's lower ILPI was less variable (COV less by 31%) than the non-smokers higher ILPI and that the variability in SHRV is reversed (smokers variability greater by 23%).

The second pair (males) consists of a current smoker (No. 1) and an ex-smoker (No. 10). The current smoker had consumed approximately 4 times the number of cigarettes as the ex-smoker, but the physical characteristics were almost identical (Table 1). The ex-smoker had a slightly higher predicted FEV₁ (by 6%), a lower closing volume (FEVC) and higher MEF25(P) (20 L min^{-1}), MEF50 (65 L min^{-1}) and PEFR (223 L min^{-1}). Both maximum and average inspiratory flow rate during aerosol inhalation were lower in the ex-smoker by 15 and 8 L mm^{-1} compared to the current smoker. However, ILPI was higher in the ex-smoker (0.94 compared to 0.62) and SHRV less 62% compared to 70% initial lung burden. Comparing the variabilities of SHRV and ILPI in the two pairs of subjects, it is seen that the healthier lungs (9 and 10) are less variable in SHRV than the less healthy lungs (1 and 11). No such clear cut trend is exhibited by ILPI. One possible explanation of the greater COV of ILPI compared to SHRV may be that tonal variations of the airways affect ILPI more than SHRV. Although the mucociliary clearance depends on site of deposition of the aerosol in that particles deposited nearer to the mouth are cleared more quickly (shorter distance to travel) small changes in deposition site as might be the result of tonal diameter changes in airways would be smoothed in the overall clearance curve up to 6 hours after inhalation. The effect of tonal changes might explain some of the observed differences in the initial part (up to about 2 hrs) of the clearance curve. In order to demonstrate this conclusively it would be required that inhalation flowrates are more closely controlled than was the case in the present study. The range of maximum inspiratory flowrates in this study was

53 to 99 L min⁻¹ and it is possible that this range of flowrates could be accommodated by changes in airway smooth muscle.

If the deposition of radionuclide tagged particles is to be used as a screening procedure for detecting early small airway disease, then the technique must be such that a high throughput can be obtained. The slowest data acquisition process in the present study was that involved in establishing the initial lateral penetration of the aerosol. With a view to speeding up this part of the process, simple vertical slices of the lung field was selected to form the basis of the initial lateral penetration index, as previously mentioned. It will be recalled that the ILPI here reported was based on slices of width 5 cm. (2 traverses each of an inch in width adjacent to the midline and periphery of the chest). An index calculated on slice width of 2.5 cm. (sum of traverse 5 counts divided by sum of traverse 1 counts) gave results which were similar to the ILPI which is tabulated (Table 3). As would be expected the statistical accuracy of the counting over the narrower slice was inferior to that and the wider slice, and this resulted in a larger COV for the 2.5 cm. slice. The mean difference between the two COV was 10% and this difference was significant at the 5% level by t test. This observation is encouraging since the possibility of using a simpler monitoring jig than the homemade precision scanner is not precluded. Feasibility studies using a minimonitor as the detector is planned.

The six hour retention value, because of its dependence on mucociliary clearance as well as on initial site of deposition of the aerosol, would probably be less viable as a screening procedure even though it gave a lower COV than ILPI in the volunteers studied.

It is concluded that the deposition of radionuclide tagged particles in the human lung can be used to study the function of large as well as

small airways. The technique used in this report would need to be simplified if mass screening was to be contemplated. Problems associated with the level of radioactivity being inhaled would be an important consideration.

TABLE 1 PHYSICAL CHARACTERISTICS, SMOKING HISTORY AND PRESTUDY OBSERVED AND PREDICTED VENTILATORY

SUBJECT NUMBER	SEX	AGE (Yr)	FUNCTION OF ELEVEN ADULT VOLUNTEERS			FEV ₁ (L)	**Obs. % Pred.	FEV ₁ FVC(%)
			HEIGHT (M)	WEIGHT (Kg)	SMOKING (10 ³ Pk - Yr)			
01	M	71	1.76	78.2	9.1	3.3 ^b	11 ^b	66 101
*02	M	43	1.75	70.8	None	4.45	103	80 106
03	F	61	1.51	47.8	8.8	2.1 ^b	118	80 105
04	M	64	1.72	65.7	9.9	2.45	81	67 93
05	M	67	1.72	84.1	1.4	3.5	121	76 11 ^b
*06	F	71	1.57	56.2	None	1.59	90	73 99
07	M	70	1.71	62.0	7.7	2.66	95	93 141
08	M	74	1.74	76.5	15.0	2.8	101	76 118
*09	F	62	1.64	62.0	None	2.4	11 ^b	78 102
10	M	72	1.78	77.4	2.5	3.60	120	77 119
11	F	69	1.66	75.7	4.4	1.79	89	64 86
MEAN			65.8	1.69	68.8	7.4	2.79	105 75.5 101
S.D.			8.7	0.08	11.0	4.4	0.86	14.1 8.1 34.4
SE(M)						0.26	4.3	2.4 10.4

Key

- * Non-Smokers
- * Obs = observed value
- % Pred. = predicted value (Cotes,)

TABLE 2 PULMONARY FUNCTION TEST RESULTS: POOLING OF OBSERVATIONS TAKEN AT WEEKLY INTERVALS OVER 3 WEEKS

SUB. No.	CLOSING VOLUME			MEF25(P)			FEV ₁			MEF50			PEFR		
	MEAN	SEM	COV	MEAN	SEM	COV	MEAN	SEM	COV	MEAN	SEM	COV	MEAN	SEM	COV
C1	35.5	2.8	19	19.9	1.7	11.3	3.4	0.02	0.89	132	4.2	5.5	325	21	11
*02	11.8	0.8	13.2	69.6	3.2	7.9	4.3	0.10	1.4	236	4.3	3.2	401	6	3
03	18.3	0.5	4.9	19.4	1.9	17	2.10	0.05	3.9	84	7.4	15.1	244	24	17
04	33.6	1.2	10.0	17.6	1.5	14.8	2.50	0.02	1.2	85	4.7	9.6	489	15	5
05	21.7	0.6	9.3	40.8	4.6	19.5	3.40	0.01	3.9	169	6.3	6.5	406	64	27
*06	28.0	1.5	18.1	21.0	4.2	34.8	1.70	0.05	4.7	88	10.0	19.8	154	23	26
*07	32.5	1.2	12.2	29.3	3.9	23.1	2.90	0.10	7.2	155	22.6	25.3	282	14	8
*08	30.2	1.1	8.8	28.5	1.5	7.4	2.60	0.04	2.3	131	21.1	23.2	353	28	11
*09	30.0	0.8	6.6	41.2	10.3	43.3	2.50	0.06	4.4	170	41.1	24.2	231	12	9
10	24.6	0.5	7.3	39.3	7.9	34.8	3.60	0.02	0.8	197	25.3	22.3	548	78	25
11	33.1	1.2	13	28.7	2.6	15.7	1.90	0.07	5.8	70	2.7	6.8	204	9	7
GRAND MEAN	26.5	1.3	11	32.3	4.6	24	2.80	0.24	3.6	138	16.0	14.7	331	37	14

Closing Volume (%VC); FEV₁ in Litres BTPS; MEF25(P); MEF50 and PEFR in Litres/min

* Non-smokers

TABLE 2 PULMONARY FUNCTION TEST RESULTS: POOLING OF OBSERVATIONS TAKEN AT WEEKLY INTERVALS OVER 3 WEEKS

SUB. No.	CLOSING VOLUME			MEF25(P)			FEV ₁			MEF50			PEFR		
	MEAN	SEM	COV	MEAN	SEM	COV	MEAN	SEM	COV	MEAN	SEM	COV	MEAN	SEM	COV
C1	35.5	2.8	19	19.9	4.7	21.3	3.4	0.02	0.89	132	4.2	5.5	325	21	11
*02	11.3	0.8	13.2	69.6	3.2	7.9	4.3	0.10	1.4	236	4.3	3.2	401	6	3
03	18.3	0.5	4.9	19.4	1.9	17	2.10	0.05	3.9	84	7.4	15.1	244	24	17
04	33.6	1.2	10.0	17.6	1.5	14.8	2.50	0.02	1.2	85	4.7	9.6	439	15	5
05	21.7	0.6	9.3	40.8	4.6	19.5	3.40	0.01	3.9	169	6.3	6.5	406	64	27
*06	28.0	1.5	18.1	21.0	4.2	34.8	1.70	0.05	4.7	88	10.0	19.8	154	23	26
*07	32.5	1.2	12.2	29.3	3.9	23.1	2.90	0.10	7.2	155	22.6	25.3	282	14	8
*08	30.2	1.1	8.8	28.5	1.5	7.4	2.60	0.04	2.3	131	21.4	23.2	353	28	11
*09	30.0	0.8	6.6	41.2	10.3	43.3	2.50	0.06	4.4	170	41.1	24.2	231	12	9
10	24.6	0.5	7.3	39.3	7.9	34.8	3.60	0.02	0.8	197	25.3	22.3	548	78	25
11	33.1	1.2	13	28.7	2.6	15.7	1.90	0.07	5.8	70	2.7	6.8	204	9	7
GRAND MEAN	26.5	1.3	11	32.3	4.6	24	2.80	0.24	3.6	138	16.0	14.7	331	37	14

Closing Volume (%VC); FEV₁ in Litres BTPS; MEF25(P); MEF50 and PEFR in Litres/min

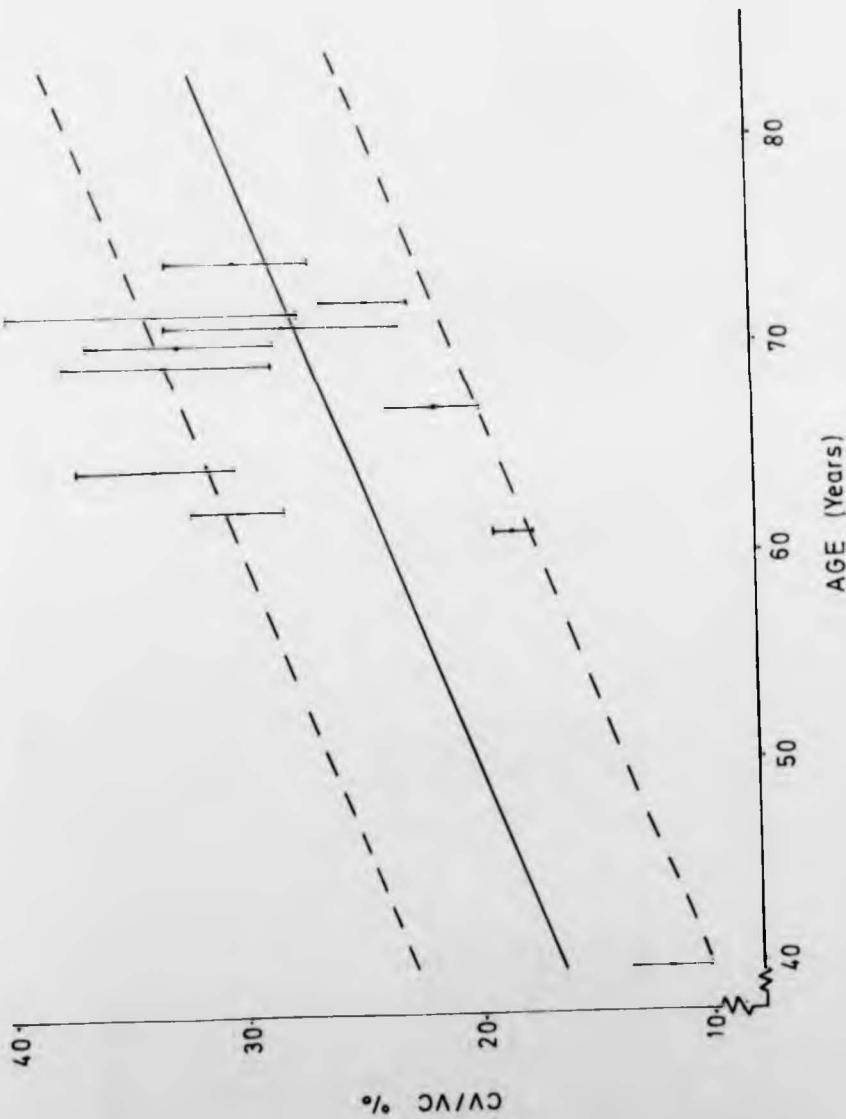
* Non-smokers

TABLE 3 MAXIMUM AND AVERAGE INHALATION FLOWRATES, INITIAL LATERAL PENETRATION INDEX AND SIX HOUR RETENTION
VALUE: POOLING OF OBSERVATIONS TAKEN AT WEEKLY INTERVALS OVER THREE WEEKS

Sub. No.	Inhalation flowrates (L/min)						ILPI	SHRV (%)	Lung				
	Maximum			Average					Mean	SEM	COV		
	Mean	SEM	COV	Mean	SEM	COV							
01	92	13	24	41	4	16	0.82	0.30	63	10	5	12	
02	88	10	19	38	4	19	0.74	0.07	16	67	5	14	
03	53	4	15	23	1	9	0.71	0.14	35	59	3	9	
04	72	4	9	37	1	4	0.52	0.09	29	74	2	5	
05	80	10	21	36	4	18	1.35	0.30	38	57	5	15	
06	85	6	13	39	2	7	0.61	0.22	62	41	6	23	
07	79	2	5	35	1	7	0.40	0.16	69	59	3	10	
08	73	12	28	33	4	20	0.66	0.16	42	67	2	6	
09	65	3	9	27	1	9	0.40	0.23	99	71	1	2	
10	77	4	9	33	2	12	0.94	0.20	37	62	2	5	
11	70	5	13	33	2	12	0.14	0.06	65	47	7	25	
MEAN	76	3	15	34	2	12	0.66	0.10	50	61	3	11	

ILPI is initial lateral penetration index; SHRV is six hour retention value as percentage of initial lung burden;
SEM is standard error of the mean and COV is the coefficient of variation - the standard deviation as percentage of the
mean.

Mean observed CV/VC % ± 1 SE compared with values predicted by Mc Carthy et al (± 2 SE)



Topographical Distribution of Inhaled Particles on 3 days, one week apart

SCH 1000 MDI STUDY 11 subjects

Right Lung

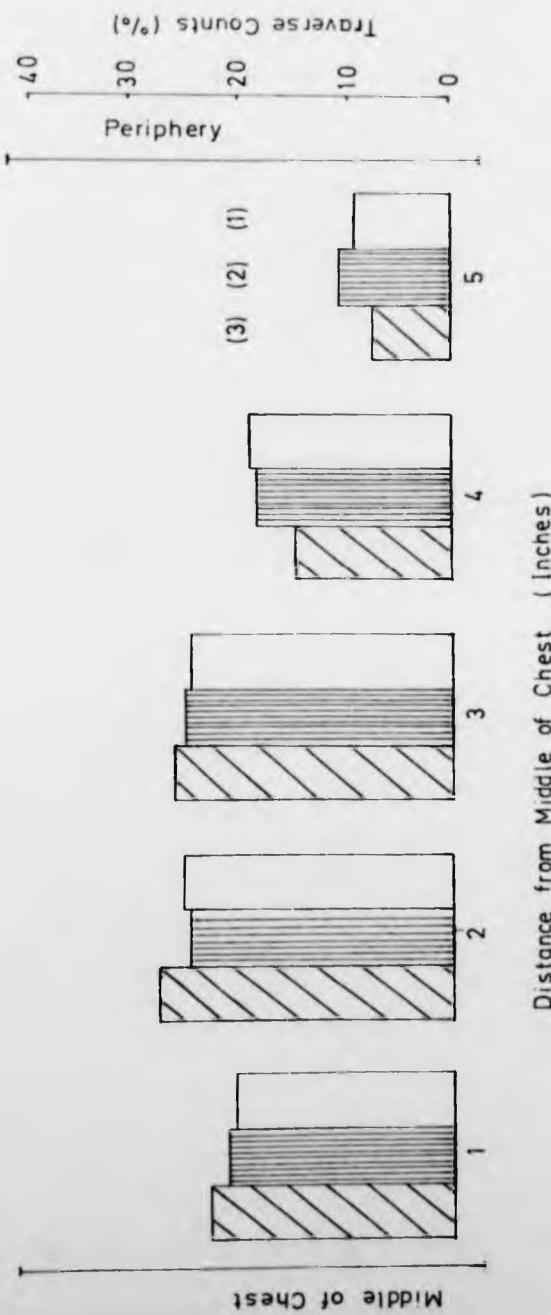


FIG.3 Scatter Diagram of Initial Penetration Index and CV/VC % in 11 Asymptomatic Subjects

$$\text{ILPI} = 1.26 + 0.02 \text{ CV/VC \%}$$
$$n = 11 \quad r = 0.49$$

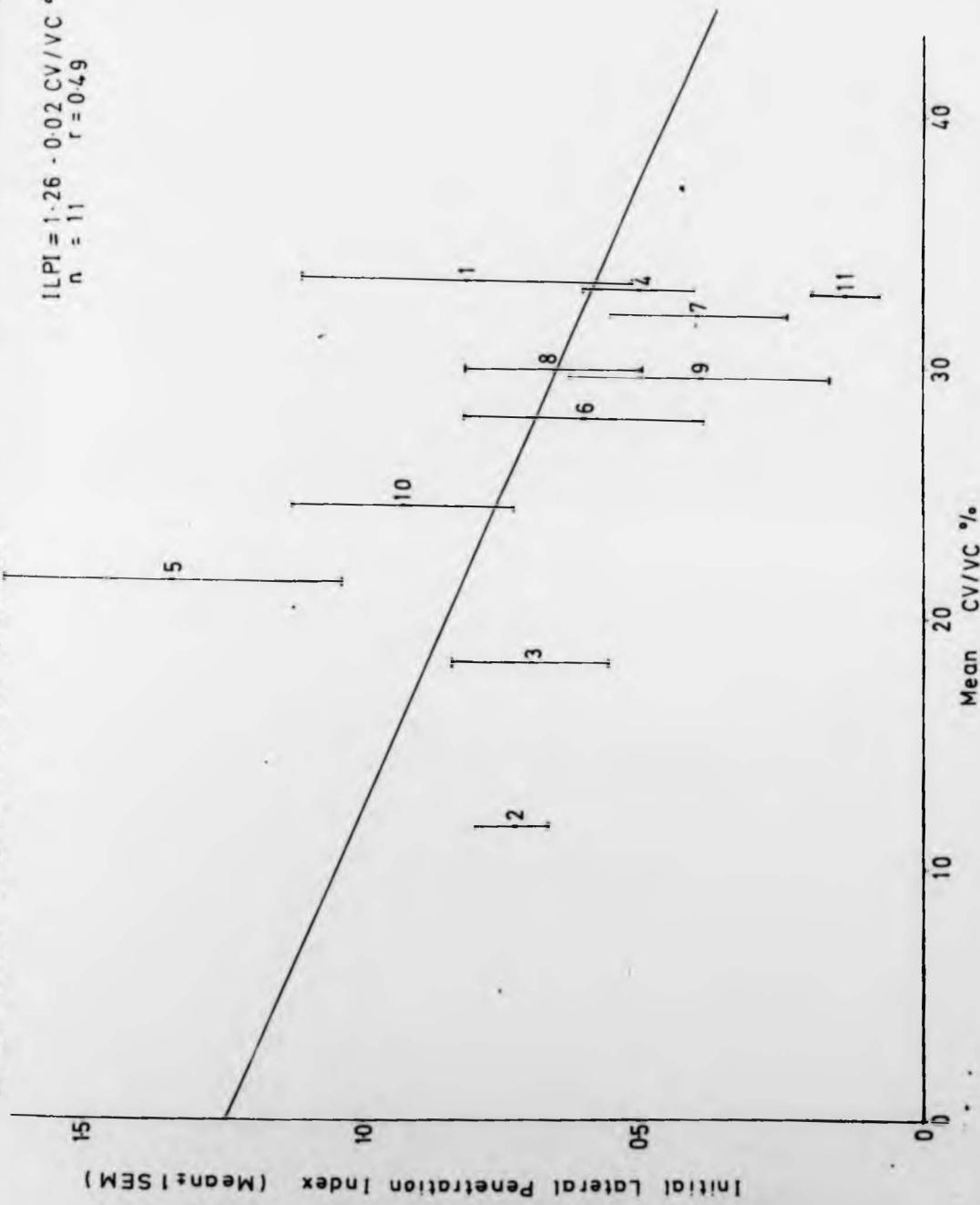


FIG 4 Scatter Diagram of Initial Lateral Penetration Index and MEF 25(P) in 11 Asymptomatic Subjects

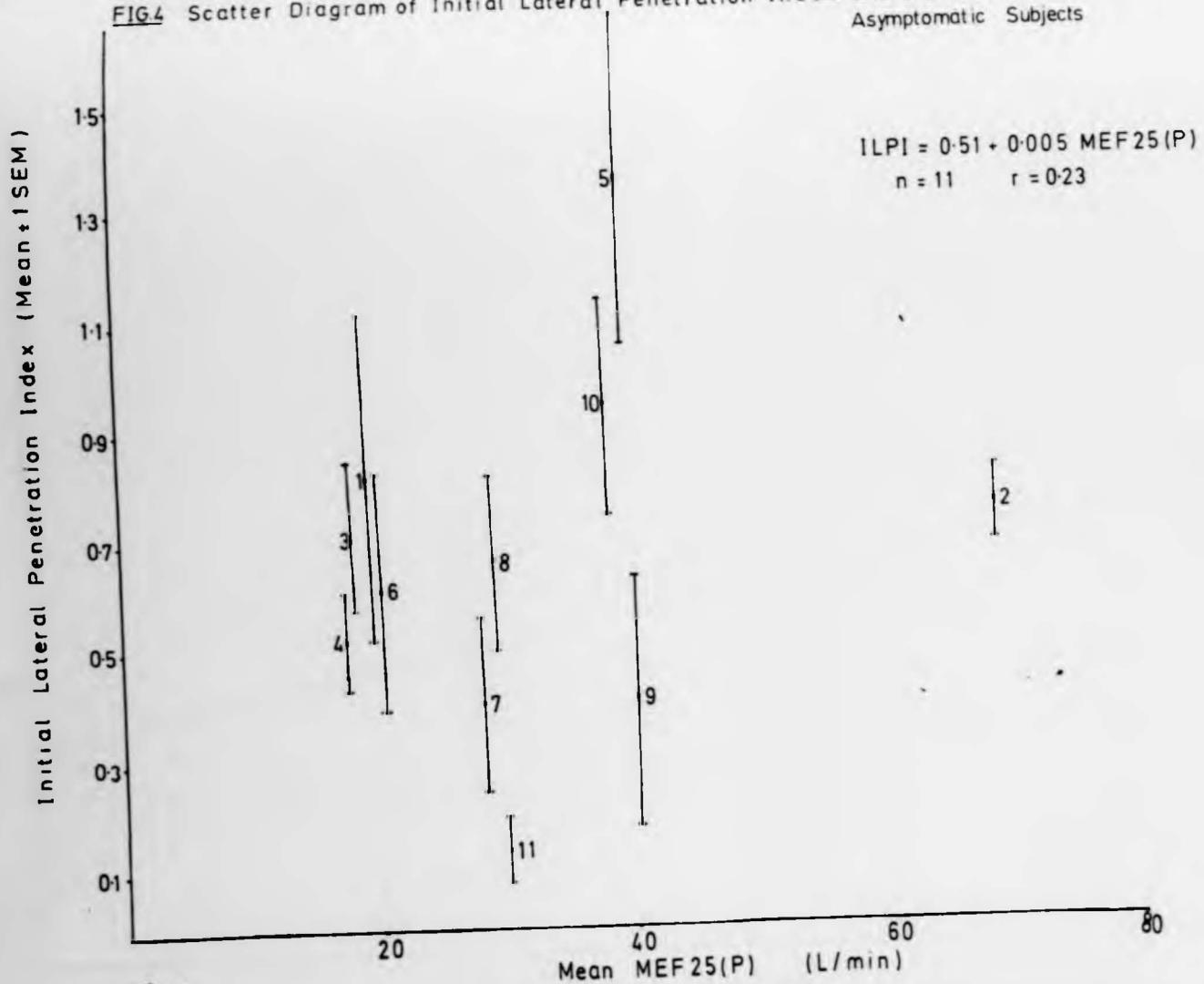


FIG.5 Scatter Diagram of Initial Lateral Penetration Index and MEF 50 in 11 Asymptomatic Subjects.

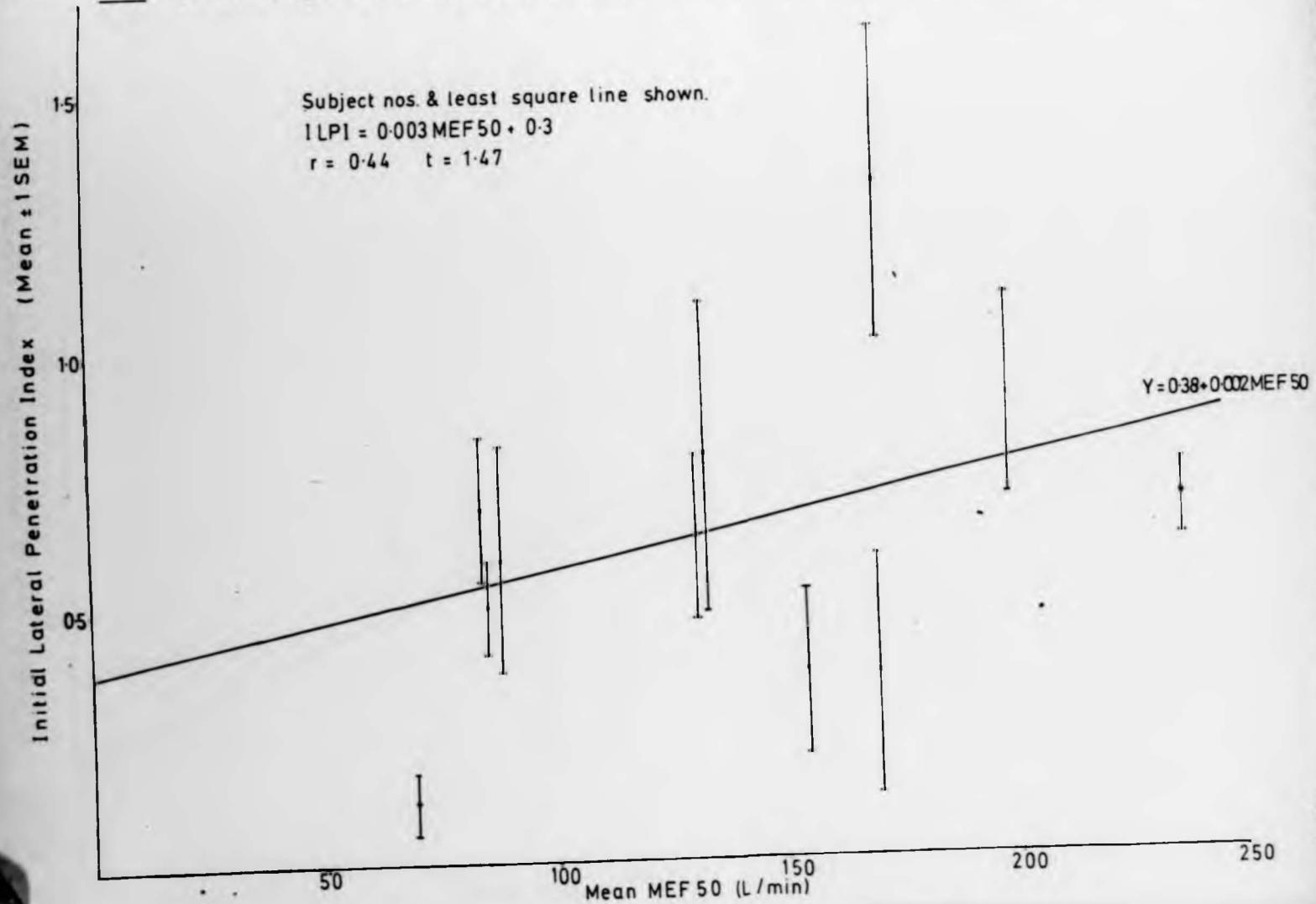


FIG.6 Scatter Diagram of Initial Lateral Penetration Index and FEV, in 11 Asymptomatic Subjects.

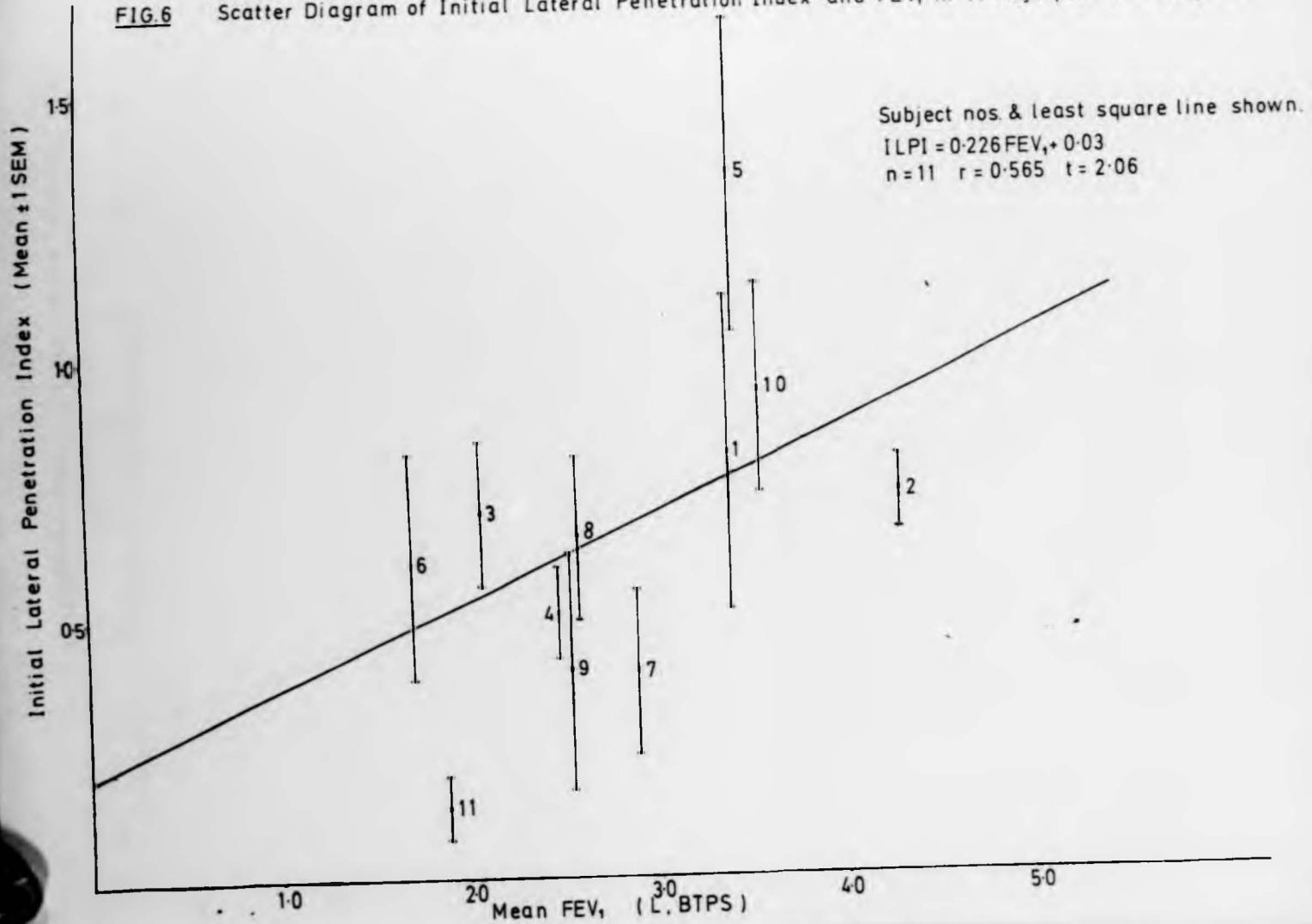
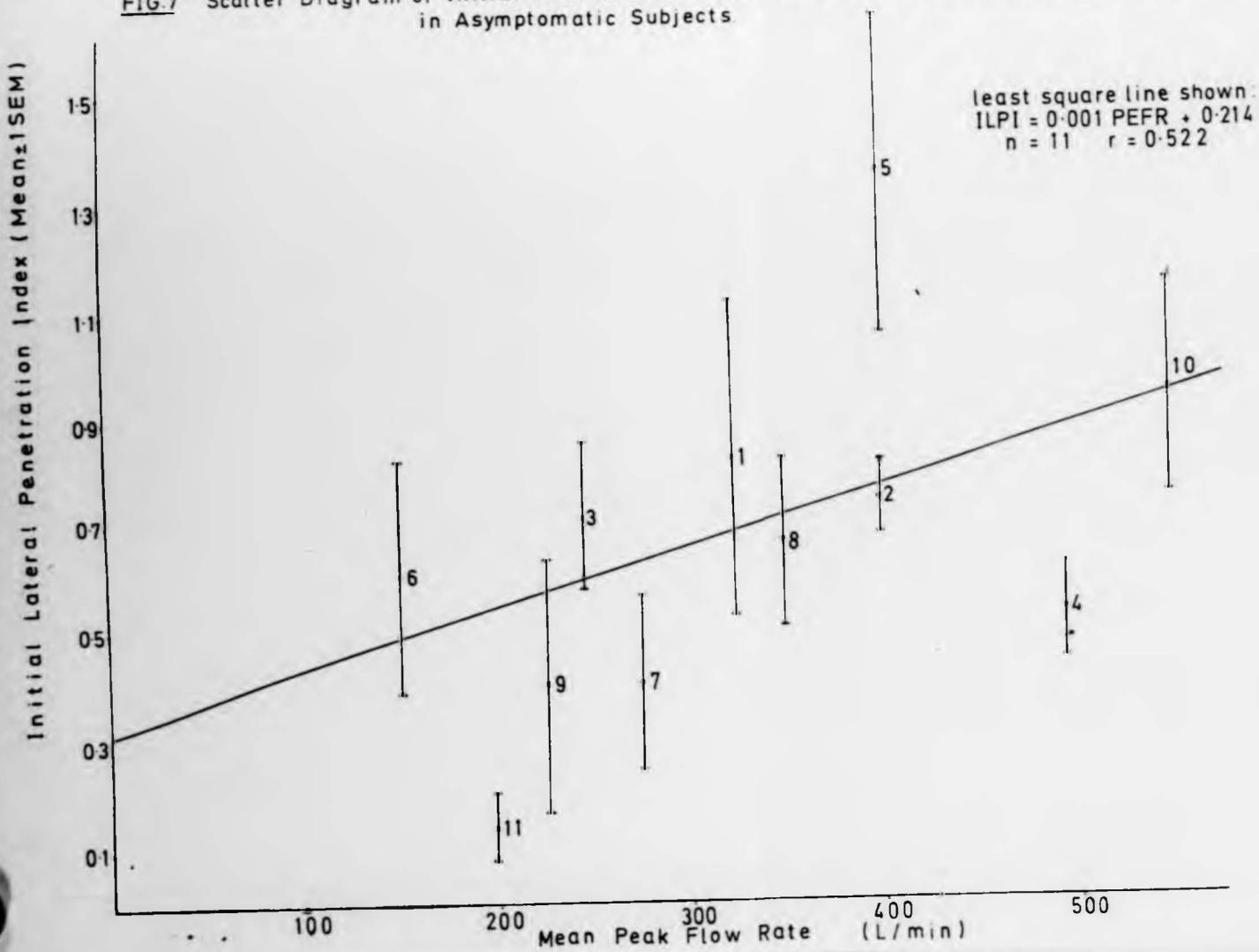


FIG.7 Scatter Diagram of Initial Lateral Penetration Index and Peak Expiratory Flow Rate
in Asymptomatic Subjects

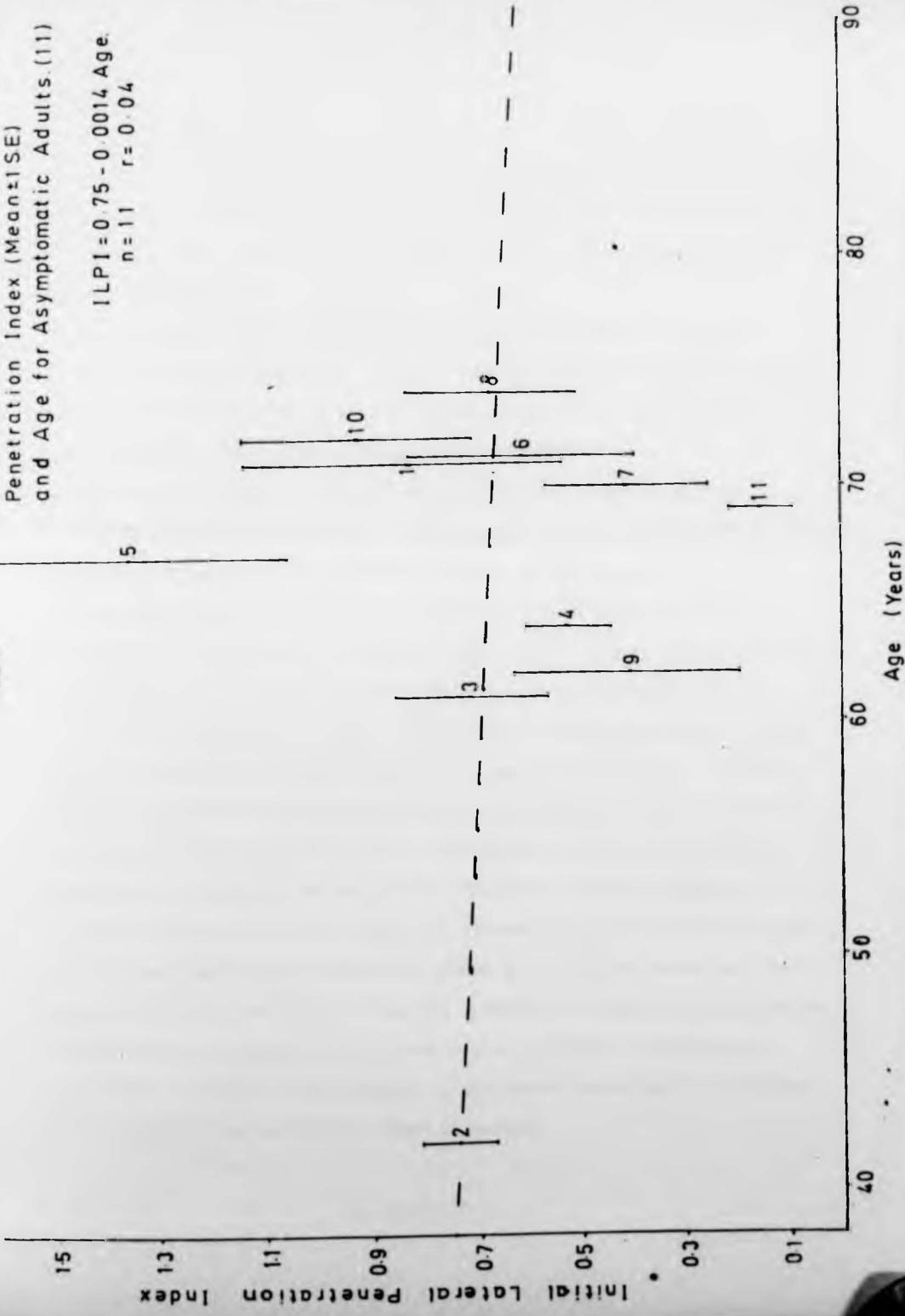


Scatter Diagram of Initial Lateral Penetration Index (Mean \pm SE) and Age for Asymptomatic Adults (11)

$$\text{ILPI} = 0.75 - 0.0014 \text{ Age.}$$

n = 11 r = 0.04

FIG 8



EQUIPMENT

The apparatus used for the major part of the work on which this thesis is based was assembled from easily available components and the total capital cost was less than £100. Analogue electronic circuitry was used because of the relative expense of digital components at the time of assembly (1973 - 1974).

The design of the equipment was such that it could be easily disassembled for sterilisation, could be transported with ease (compact modular design to fit into boot of average family saloon), and was robust enough to maintain stability and reliability.

Over the two years in which the apparatus was used it proved remarkably stable and reliable. The calibration was assessed at frequent and regular intervals but adjustments were rarely needed.

To reduce the hazard of cross infection from equipment used in epidemiology the ability to disinfect the apparatus between subjects is of considerable importance. This was achieved in my apparatus by the inclusion of disposable sterile gauze pads at strategic points. These pads were changed after each subject had used the machine. The whole apparatus was disassembled and sterilised thoroughly at weekly intervals. The advice of the Microbiological Hazards Safety Officer was sought and followed regarding the method and frequency of sterilisation.

Both the short term and long term comparison of the equipment with standard manufactured apparatus (chapters 2 and 3) has underlined the inherent quality. Meticulous care in approach and methodology, (choice of electronic components and circuit layout) however contributed to the results obtained. No evidence is presented regarding the behaviour of the apparatus in a multiple user situation.

The rapid expansion in the variety and availability of digital circuitry including microprocessors in recent years would probably make a repeat design of similar analogue equipment less than cost effective.

Collection of Data

The volume of data which can be handled by equipment having an analogue output format is restricted because transcription is by clerical rather than automatic mechanical or electrical means. The quality of the data is not impaired however compared to that which is acquired in digital form. This is confirmed by the evidence in chapter 4 since the FM tape recorder acquires information in a mode which is closely analogous to digital acquisition.

Design of Experiments

The design of the experiments (chapters 5, 6 and 7) took account of the above limitations. The statistical methods by which the resulting data was to be analysed determined the method of collection.

Choice of Indices of Pulmonary Function

The thirteen indices (subject of chapter 7) were chosen having regard to the current (1975) popularity and the ability of the apparatus to produce them reliably. Additional indices, in particular the slope of the alveolar plateau on closing volume tracings, could have been chosen, but these were not of current interest at the time of planning the experiments.

Chapter 7 reports an assessment of the relative merits of pulmonary function indices. The method was designed to 'test' the Tests rather than testing the volunteers. This approach is different from the more usual epidemiological approach in which large unselected populations are involved, and the quality of the tests inferred from the results.

The merit of the approach used is that information on the 'tests' is gained directly.

In a thesis dealing with the evaluation of small airway disease, the absence of experimental work on clinically defined disease states such as Asthma and Bronchitis is perhaps unusual. The lack of Ethical Committee permission to pursue such studies however did not severely limit the scope of the work, but focussed attention on the 'Tests'

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 - to my son, Julian, who has not always understood the reason why I could not come out to play
- and
- to the typists who have so carefully typed the script.

Bronchoactive drugs and other assistance was obtained from:-

Messrs. Boehringer Ingelheim Ltd., (Sch 1000)

and

Messrs. Upjohn Ltd.,

(Prostaglandins)

CRITICAL REVIEWIntroduction

The apparatus which was designed and constructed by the Author for use in the field has been described and evaluated in Chapters 1 - 3 above. The sources of error which are now described are mainly inherent in the methods especially where a body plethysmograph is not available or its use impracticable.

Sources of Experimental Error in the Tests1) Closing Volume

A major source of error, up to 100 ml, in this test is associated with the determination of the point of intersection of Phases III and IV. A contributing factor is the error in defining the 'best fit' line through the alveolar plateau. The error is greater in subjects with smaller vital capacities because of the shortness of the plateau. Aid in locating the point is given by the diminution in the amplitude of cardiogenic oscillations as the intersection is approached. However these oscillations are not always present even in the same person performing replicate manoeuvres.

Expressing the closing volume as a percentage of the vital capacity and rejecting tracings in replicate tests where the vital capacities exceeds $\pm 5\%$ of the mean has helped to reduce the variation in the test. Subjects tested at the same time of day on three occasions over a period of three weeks gave for the CV/SVC% coefficients of variation (COV) in the range 5 - 19% with a mean of 11% (Ch.8). This compares with an average COV of 18% derived from the published standard values of McCarthy et al.

Perhaps the greatest drawback of this test is that some subjects do not appear to have a closing volume especially between the ages 18 and 25.

ii) Flow Volume

Unlike the closing volume test most persons produce an apparently technically competent maximal or partial expiratory flow volume tracing whether air alone or a mixture of helium-oxygen is being breathed. However variation in the location of the tracing within the vital capacity range may introduce appreciable error. This variation is greater where the flow volume manoeuvre is done to assess the effect of agents which may alter the vital capacity (e.g. bronchoconstrictors/bronchodilators). Such variation can be reduced by performing the tests within a body plethysmograph.

Because the Residual Volume (RV) is more influenced by small airway calibre and compressibility a larger systematic error will be involved if 'before' and 'after' curves are aligned at RV than if they were aligned at total lung capacity (TLC). Notional alignment at TLC, (as done in this study), using a separate vital capacity manoeuvre, is also preferable when 'partial' curves are required.

It is doubtful whether the Function Residual Capacity (FRC) obtained by subsequent tidal volume trace is more accurate for this purpose although return to FRC in replicate spirograms is normally less variable than RV or TLC.

Another source of error is involved in the precision with which the instantaneous flow rates can be determined from the flow-volume tracings. In general the closer to the RV the flow rate ordinate is measured the larger the reading error because of the finite thickness of the recorded line. For example a mean COV for MEF25 (repeat studies on the same subjects on 3 occasions) of 24% (range 8 - 43%) was found and the corresponding figures for MEF50 were mean 15% (range 3-41%). This type of error must be taken into account when indices reflecting small airway behaviour are chosen from the expiratory flow volume curve. Doubt has been cast on the MEF50 as

representing small airway behaviour by Macklem (personal communication) although this index has increased in smokers after cessation of the smoking habit. In this respect the MEF⁴⁰ may be better at representing small airway behaviour than MEF⁵⁰ despite a larger reading error in the former. The MEF⁴⁰ is also known to have a fairly constant relationship to the absolute lung volume scale determined plethysmographically, being TLC-60% of initial vital capacity.

The effect of using a helium-oxygen mixture to trace a flow volume curve is to increase the flowrates at all volumes where turbulent flow predominates. The flow volume trace would therefore be amplified (compared to air) in about the upper two thirds of the vital capacity range resulting in a proportionate reduction in the flow rate ordinate reading error. In the lower third of the vital capacity this error is either unaffected or become slightly greater because flow rates are less than or equal to those on air.

By incorporating both air and helium-oxygen flow volume curves in the same test, and index, the isoflow volume point (IFVP) can be obtained. The average value of this index in the smokers studied was $26 \pm 3\%$ compared to $19 \pm 2\%$ for the non-smokers (Ch.7) supporting the view that a flowrate ordinate in the last third of the vital capacity is more likely to reflect small airway changes than an ordinate in the upper or middle third.

iii) Radioactive Tracer Aerosol Inhalation

The initial lateral penetration index (ILPI) and the six hour retention value (SHRV) (which to some extent depends on ILPI) are recent innovations which are as yet not validated. They are potentially valuable as being independent of measurements on expired air. The largest errors on this test will be associated with the reproducibility of flow rate during inhalation and volume of aerosol inhaled. Both the lung volume at which inhalation begins

and the volume of aerosol inhaled can be closely controlled, but even under these constraints inspiratory flow rates can vary widely. The COV of both ILPI (mean 50%, range 16-99%) and SHRV (mean 11%, range 2-25%) on repeat studies of the same individuals are largely due to the differences in inspiratory flowrates attained by an individual in replicate manoeuvres.

The simple ratio used to calculate ILPI probably gives rise to non-linearity and a better index will almost certainly be found with increased experience in this technique. The SHRV may be less promising than ILPI as a measure of small airway function as it is greatly influenced by cough and mucociliary function.

Consistency of Results

Smokers are often used as a model of small airway disease on the assumption that adverse effects will be first manifest in the small lung airways. It is more likely however that the influence will be felt over the entire airway generation since indices such as MEF25, peak expiratory flow rate and FEV₁ are reported to have increased following cessation of the smoking habit. When smokers and non-smokers have had their pulmonary function assessed, consistency in the results is often judged by observing whether the smokers' results are overall inferior to those of the non-smokers. Such has been the case in this Thesis.

Another way of checking consistency is by noting correlation between indices and anthropometric data. An apparent inconsistency exists in Table 7 of Ch.7 where the FEV₁, instead of declining with increasing age, gave a positive correlation $r = 0.61$ $P < 0.05$ for smokers. This may be due to the differences in tobacco consumption which was greater in the younger smokers.

A different unexplained feature is observed, (Table 3, Ch.7) where repeat determinations on trained subjects have resulted in a 'significant' trend in obtaining lower values on the second occasion. If this

observation is not an artifact of the methodology, it may imply that even in trained subjects apprehension, anxiety or other emotional state, possibly associated with the first test, may affect the results.

General Assessment

Probably the most significant observation in this Thesis is the importance of the role of airway smooth muscle tone in contributing to variability in flow volume curves and other pulmonary function tests. Even where forced expiration is not involved, as in radioactive tracer aerosol inhalation, variation in airway muscle tone may have affected function findings by varying the inhalation velocity.

The significance of tone in the production of the characteristic flow volume curve, 'notch' has not been fully evaluated and needs further research.

A systematic use of both 'partial' and 'maximal' flow volume curves to chart the effects of bronchoactive agents on airway smooth muscle may lead to the establishment of the range of calibre change which can be expected in health and disease. The use of histamine, which may mainly constrict medium sized airways, could be a useful starting point.

Recommended Test

Tests of lung function should be appropriate to the circumstances for which they are required. They should therefore discriminate between relevant and irrelevant aspects of function in a particular situation. In assessing small airway disease early detection is more relevant only where reversibility is a possibility since only palliative measures can be instituted against chronic disease.

On the basis of the analysis (Ch.7) certain indices seem more appropriate than others for detecting reversible effects. The discriminant functions which are central to the ranking of these indices have yet to be verified in an independent population of persons (two distinct populations of changes existed in the study). The isoflow volume point (IFVP) appeared to be as good as closing volume in detecting reversible changes, so that if only a single test is to be recommended, it would be the combination of air and He-oxygen flow volume curves. The additional information which can be gained by the use of helium more than compensates for the slightly increased effort and cost.

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IPRATROPIUM BROMIDE: MUCOCILIARY CLEARANCE RATE AND AIRWAY RESISTANCE IN NORMAL SUBJECTS

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Summary

The effect of ipratropium bromide, a new anticholinergic drug, on the rate of clearance of secretions from the lung and on airways resistance was investigated in 12 healthy subjects in a double-blind cross-over trial with placebo and a control run without aerosol. Before taking the drug the subjects inhaled uniform $5\text{ }\mu\text{m}$ tracer particles of polystyrene in which ^{99m}Tc had been unleachably incorporated. The initial depth of deposition and the rate of clearance of the particles were obtained from serial gamma counts made externally to the chest over six hours.

The difference between drug, placebo and control runs in the deposition patterns of the tracer particles and their subsequent rates of clearance were not significant. The drug treatment resulted in statistically significant falls in specific airway resistance at 1, 2, 3 and 6 hours ($P < 0.02$). There was no objective or subjective evidence of side effects from the drug.

INTRODUCTION

Ipratropium bromide is a synthetic anticholinergic agent, chemically a quaternary ammonium compound. The study reported here was designed to examine the effect of ipratropium bromide and of the propellants in which it was administered on lung mucociliary clearance and the effect of the drug as a bronchodilator in healthy subjects.

Subjects and Methods

Twelve healthy volunteers were studied. Table I summarizes the physical characteristics, tobacco consumption and ventilatory capacities of the 12 volunteers. All subjects gave informed consent.

In a random double-blind manner each of the 12 subjects took single doses of ipratropium bromide and a placebo consisting of the propellants only (freon 11, 12 and 114 in proportions 1:2:1 plus soya lecithin surfactant). The manufacturer's instructions were followed. For each puff of the dispenser the subjects expired fully, inhaled the aerosol slowly and maximally, then held their

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breath for 5 seconds (mass median diameter: $0.8 \mu\text{m}$; geometric standard deviation: $2.2 \mu\text{m}$). In a third control run no medication was given. (The control as well as the drug and the placebo are here called treatments.) Three doses of the drug were compared: 0.04, 0.08 and 0.16 mg in 2, 4 and 8 puffs. Each dose was taken by four subjects in random order.

The order of measurements throughout each 6-hour trial run was as follows. The treatments were administered at zero hour. The tracer aerosol was given before treatments (zero–10 minutes) so that any change in airway calibre caused by treatment would not affect the penetration of the tracer particles and hence their subsequent rate of clearance. Immediately after inhalation the topographical distribution of the deposited particles in the right lung was obtained by rectilinear gamma scanning. Whole lung gamma counts were also made at half-hourly intervals up to 6 hours to assess the rate of removal of the particles from the lungs. Airway resistance measurements were done 20 minutes before and at 1, 2, 3 and 6 hours. The three runs were one week apart.

Table 1. Physical characteristics, tobacco consumption and ventilatory capacity in 12 subjects

Characteristics	Results (mean \pm SD)
Age (years)	67 ± 9.8
Height (m)	1.68 ± 0.84
Smoking history* (packet years)	25 ± 20
FEV ₁ observed (litres)	2.62 ± 1.02
FEV ₁ % predicted†	98 ± 24
FEV ₁ observed (litres)	3.55 ± 1.19
FEV ₁ % predicted†	105 ± 20
FEV ₁ /FVC % observed	72 ± 13
FEV ₁ /FVC % predicted†	104 ± 19

* 3 non-smokers, 6 current smokers, 3 ex-smokers.

† Predicted values from Cotes (1968).

The technique for assessing mucociliary clearance has been fully described by Thomson and co-workers (Thomson & Short 1969; Thomson et al. 1973). The $5 \mu\text{m}$ unit density tracer particles were inhaled before medication with a tidal volume of 500 ml without breath holding. The radioactive lung burden did not exceed $30 \mu\text{Ci}$ of ^{99m}Tc in any one trial run.

The forced expiratory volume in 1 second (FEV₁) and forced vital capacity (FVC) were measured by dry bellows spirometer (Drew & Hughes 1969). Airways resistance was also measured by constant volume body plethysmograph (Dubois et al. 1956) as specific resistance (SR_{aw}).

RESULTS

Lung clearance

Fig. 1 shows the mean initial distribution of the inhaled tracer particles across the right lung for all 12 subjects. For each subject the counts for each vertical traverse were first expressed as percentages of the total traverse counts. The height of the columns in the figure are the means of these percentages. In none of the five individual traverses (Fig. 1) was the effect of the treatment significant as shown by analysis of variance. An alternative index of penetration (Thomson & Pavia 1974) may be obtained from the ratio of the sums of the means of traverses 4 and 5 and to those of 1 and 2. The ratios were 0.45, 0.45 and 0.63 for control, drug and placebo treatments respectively; again these differences were not significant.

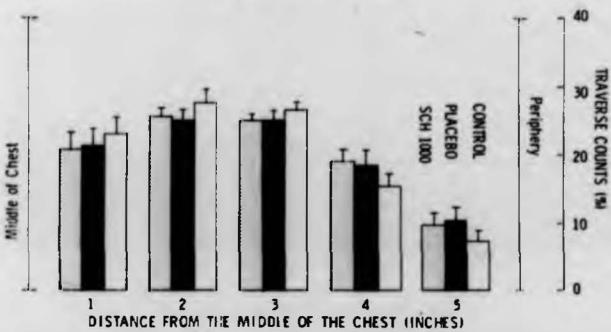


Fig. 1. Initial lateral distribution (mean \pm SE) of inhaled particles across the right lung of the 12 subjects for the drug, placebo and control runs. Sch 1000 = ipratropium bromide

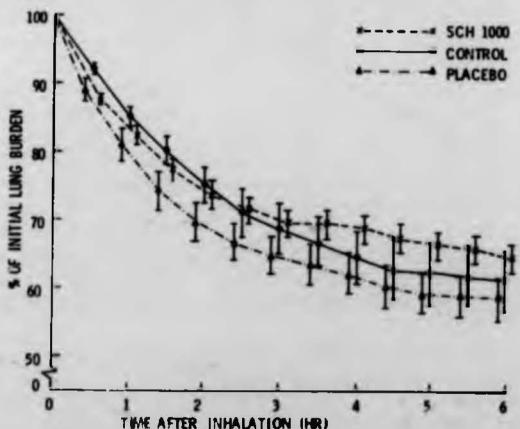


Fig. 2. Whole lung clearance curves (mean \pm SE) for the 12 subjects in drug, control and placebo runs

Fig. 2 shows the mean whole lung clearances for the 12 subjects after normalizing the initial counts to 100%. The percentages of the initial lung burden cleared in six hours were 35, 38 and 41% for the drug, control and placebo runs respectively. Analysis of variance showed that there was no significant difference between the three treatments. There was no evidence of dose effects on penetration or clearance.

Specific airway resistance

Since random variations in pulmonary function values between days is much greater

than between different times on the same day experimental measurements have been analysed, in line with most other authors, as differences from the pre-treatment values.

For all doses and treatments there was a fall in SR_{aw} over the first three hours after treatment with a rise towards pre-treatment levels at six hours. This general trend has been attributed to diurnal rhythm (McDermott 1966). Fig. 3 shows the mean changes in SR_{aw} expressed as percentages of the pre-treatment values for the 12 subjects for the drug, placebo and control runs. For the drug run but not for the control and placebo the mean SR_{aw} at all post-treatment measurements was significantly less by the non-parametric sign test than the pre-treatment value ($P < 0.02$). Separate analyses of variance for each dose level showed that the differences between the three doses were not significant, probably due to the small numbers of subjects in each group.

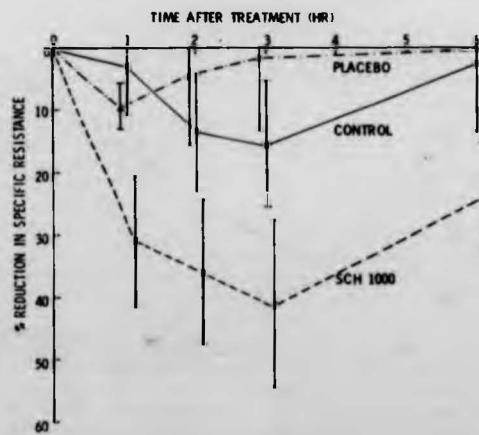


Fig. 3. Percentage changes from pre-treatment values (mean \pm SE) in specific resistance against time for the 12 subjects in the placebo, control and drug runs

DISCUSSION

Intravenous atropine is followed after about 40 minutes by virtual cessation of mucociliary clearance (Lippman et al. 1975). It is not known whether this effect is due to direct ciliastasis for which atropine has been incriminated (Corrsen & Allen 1959) or to change in the rheological characteristics of the mucus, for example by increased viscosity of the periciliary layer. The delay of 40 minutes suggests that a change in secretions rather than ciliastasis may have been responsible. In a previous study on 12 humans given cumulative doses of up to 1.2 mg of ipratropium bromide by inhalation there were no differences in salivation or heart rate attributable to the drug (Bleichert 1975). Since secretagogue drugs frequently have parallel effects on salivary and other mucous gland secretions it is perhaps not surprising that the present study has shown no evidence of mucociliary impair-

ments have been treatment values. three hours after general trend has mean changes subjects for the and placebo the y the non-paras es of variance were not signifi-

ment. It is difficult to compare the doses of atropine in the above experiments or those of atropine and ipratropium bromide. Blackwell et al. (1974) have shown that only 10% of tritium-labelled isoprenaline administered from pressure-packed canisters was absorbed via the lung, and it is unlikely that the lung absorption of ipratropium bromide was any greater in the present trial.

The results from the body plethysmograph have been expressed as SR_{aw} ($= R_{aw} \times TGV$) because the reciprocal conversion is grossly non-linear at the low R_{aw} found in normal persons. Although the curve for the drug in Fig. 3 rose towards pre-treatment level after reaching a minimum at three hours there was little, if any, change between three and six hours in relation to the control curve, especially if the changes in both curves are expressed as percentages of the three-hour SR_{aw} values. This indicates that the apparent rise in the drug curve (fall in SR_{aw}) during this period may be due to diurnal rhythm and if so its bronchodilating effect may still be present at six hours. This requires confirmation since there was considerable scatter in the data.

Sterling and Batten (1969) found up to 25% bronchoconstriction in normal subjects after inhaling a dichlorodifluoroethane propellant with added detergents. The action was believed to be mainly due to surfactant but the propellant itself may have had a slight constrictive action. However caused, bronchoconstriction via the propellants could lead to underestimation of the bronchodilator effects of the drug if the drug were to be compared only with the control run. A further difference between the two preparations was the marked coughing in Sterling and Batten's subjects in response to inhalation of the propellant which was not present in our subjects. Coughing implies stimulation of the epithelial irritant receptors which would normally cause bronchoconstriction. Until the effect of propellants is more clearly defined it appears to be desirable, in trials of this nature, to include a placebo of propellant only in addition to a control as has been done in this experiment. A stronger reason for doing so is that without a placebo aerosol the experiment would not be blind.

In conclusion, ipratropium bromide did not impair mucociliary clearance significantly when administered by aerosol to 12 normal subjects in doses up to four times the therapeutic dose. Subjectively there were no adverse effects of the drug. The drug produced significant bronchodilation even in these normal subjects, reducing the mean specific airway resistance by 40% at two to three hours after administration.

ACKNOWLEDGEMENTS

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APPENDIX

Pavia and Robert B. Douglas

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Original data is listed by patient and drug giving details of physical characteristics and smoking history. The 24 columns are read in pairs (12 pairs) the first column in each pair giving baseline data and the second response data. The three rows contain the replicate data. Reading from left to right the 12 pairs of columns have data for:- MEF 25 (Air), MEF 25 (He), MEF 40 (Air), MEF 40 (He). Excess helium flowrate at 25% VC, excess helium flow rate at 40% VC, closing volume, slow vital capacity (on CV manoeuvre), FEV₁, forced vital capacity CV/SVC %, and FEV₁/FVC%. Missing values are indicated by * and arise if the manoeuvre was not performed or was not technically competent when performed. Drug 1 indicates Acetylcholine, Drug 2, Prostaglandin F₂ , Drug 3, Ventolin and Drug 4, Ipratropium bromide (Sch 1000).

The data is listed as deflection in cm. on the X-Y plotter.

From the replicates calculation of the mean, standard deviation and standard error of the baseline data (columns 1, 3, 5, 7, 9, and 11 and rows 1, 2, and 3). Available format did not allow the sign of the maximum change to be shown but the data was stored in the computer memory with sign so that calculations made by the computer took sign into account.

The indices are listed in the same pairs of columns as in the original data. 1FVP values are not listed but were calculated from the appropriate raw data columns by a Fortran program, (17/05/77) Ruperts' statistics (IVV Location).

ACCORDING TO YOUR INPUT FORMAT, VARIABLES ARE TO BE READ AS FOLLOWS

VARIABLE	FORMAT	RECORD	COLUMNS
SVC2	F 5. 2	2	55- 59
FEV1	F 5. 2	2	60- 64
FEV2	F 5. 2	2	65- 69
FVC1	F 5. 2	2	70- 74
FVC2	F 5. 2	2	75- 79

THE INPUT FORMAT PROVIDES FOR 26 VARIABLES. 26 WILL BE READ.
IT PROVIDES FOR 2 RECORDS (*CARDS*) PER CASE. A MAXIMUM OF 79 COLUMNS* ARE USED ON A RECORD.

```

MISSING VALUES V25AIR1 TO CV2 (99.00)/CIGTOT,CIGH0(-1,0)/SVC1 TO FVC2(0.0,99.0)
COMPUTE V25FRAC1=(V25HE1/V25AIH1)-1
COMPUTE V25FRAC2=(V25HE2/V25AIH2)-1
COMPUTE V40FRAC1=(V40HE1/V40AIH1)-1
COMPUTE V40FRAC2=(V40HE2/V40AIH2)-1
COMPUTE CVPSVC1=(CV1/SVC1)*100
COMPUTE CVPSVC2=(CV2/SVC2)*100
COMPUTE FEVPPVC1=(FEV1/FVC1)*100
COMPUTE FEVPPVC2=(FEV2/FVC2)*100
COMPUTE STATHS=S*HCAT+7*SEX
IF (V25FRAC1 GT 0) HED0=0
IF (V25FRAC1 LT 0) HED0=-1
IF (V40FRAC1 LT 0 AND V25FRAC1 LT 0) HED2=2
IF (V40FRAC1 LT 0 AND V25FRAC1 GT 0) HED2=1
ASSIGN MISSING V25FRAC1 TO FEVPPVC2 (99.00)/STATHS, HED2(9)
PCODE V25AIR1(9,00#99.00)
DO REPEAT V1=V25AIR1 V25HE1 V40AIR1 V40HE1 CV1 SVC1 FEV1 FVC1
      V25FRAC1 V40FRAC1 CVPSVC1 FEVPPVC1
      /V2=V25AIR2 V25HE2 V40AIR2 V40HE2 CV2 SVC2 FEV2 FVC2
      V25FRAC2 V40FRAC2 CVPSVC2 FEVPPVC2
      /V3=V25AIR3 V25HE3 V40AIR3 V40HE3 CV3 SVC3 FEV3 FVC3
      V25FRAC3 V40FRAC3 CVPSVC3 FEVPPVC3
      V3=V2-V1
COMPUTE
END REPEAT
ASSIGN MISSING V25AIR1 TO FEVPPVC3(999.0)
COMPUTE INTA1=V25AIR1-V25HE1
COMPUTE INTA2=V25AIR2-V25HE2
COMPUTE INTB1=(V40AIR1-V25AIR1)-(V40HE1-V25HE1)
COMPUTE INTB2=(V40AIR2-V25AIR2)-(V40HE2-V25HE2)
ASSIGN MISSING INTA1 TO INTB2(999.0)
IF (INTA1 EQ 0 OR INTB1 EQ 0) IVV1=999.0
IF (INTA2 EQ 0 OR INTB2 EQ 0) IVV2=999.0
IF (INTA1 NE 0 AND INTB1 NE 0) IVV1=15.0*INTA1/INTB1
IF (INTA2 NE 0 AND INTB2 NE 0) IVV2=15.0*INTA2/INTB2
IF (IVV1 LT -75.0 OR GT 25.0) IVV1=999.0
IF (IVV2 LT -75.0 OR GT 25.0) IVV2=999.0
ASSIGN MISSING IVV1 IVV2(999.0)
COMPUTE IVVP1=IVV2-IVV1
COMPUTE IVFP1=25.0-IVV1
COMPUTE IVFP2=25.0-IVV2
COMPUTE IVFP3=IVFP2-IVFP1
ASSIGN MISSING IVV3 TO IVFP3(999.0)
PCODE CIGTOT(-1,0)

```

HIPERT'S STATISTICS (IVV LOCATION)

17/05/77 PAGE 3

VALUE LABELS STATUS(0)FEMALE NON SMOKC(1)FEMALE SMOKING(2)MALE NON SMOKING
L(MALE . S(MOKING/HFD2(0))IVV < 25 (1)25 < IVV < 40
(2)IVV >40 (3)*****
PRINT FORMATS I(17A1.1)/SFX TO TEST(0)/V25AIR1 TO FEVPFVC2(2)
/HEIGHT V25AIR3 TO IVV3(2)/HEIGHT TO CIGNO(1)
/IVVPI TO IVFP3(2)
PAGE SIZE NOEJECT
COMPUTE CIGNO=CIGNO*1000
COMPUTE CIGTOT=CIGTOT*1000

17536(DKC/PAL) CM NEEDED FOR WRITE CASES

WRITE CASES (A3,F1.0,F2.0,F3.2,F3.1,F1.0,F5.0,F6.0/12F8.2/12F8.2)
INITIAL SEX AGE HEIGHT WEIGHT SMOKCAT CIGNO CIGTOT V25AIR1 TO
V40HR2 V25FRAC1 TO V40FRAC2 CV1 TO FVC2 CVPSVC1 TO FEVPFVC2

FINISH

HIPERT'S STATISTICS (IVV LOCATION)

17/05/77 PAGE 4

291 CASES WERE WRITTEN ON ALTERNATE OUTPUT FILE TAPE1.

EACH CASE CONTAINS 32 VARIABLES

PUR COMPLETED

NUMBER OF CONTROL CARDS READ 64
NUMBER OF ERRORS DETECTED 0

AVERAGED DATA

ACCORDING TO YOUR INPUT FORMAT, VARIABLES ARE TO BE READ AS FOLLOWS

VARIABLE	FORMAT	RECORD	COLUMNS
FEFV1SD	F 8, 2	5	89- 96
FEFV3MN	F 8, 2	5	97- 104
IVFP1SE	F 8, 2	6	1- 8
IVFP3SE	F 8, 2	6	9- 16
V25AH1SE	F 8, 2	6	17- 24
V25AH2SE	F 8, 2	6	25- 32
V25HE1SE	F 8, 2	6	33- 40
V25HE2SE	F 8, 2	6	41- 48
V40AR1SE	F 8, 2	6	49- 56
V40AR2SE	F 8, 2	6	57- 64
V40HF1SE	F 8, 2	6	65- 72
V40HF2SE	F 8, 2	6	73- 80
V25FH1SE	F 8, 2	6	81- 88
V25FH2SE	F 8, 2	6	89- 96
V40FH1SE	F 8, 2	6	97- 104
V40FR3SE	F 8, 2	7	1- 8
CVISE	F 8, 2	7	9- 16
CVSSE	F 8, 2	7	17- 24
SVC1SE	F 8, 2	7	25- 32
SVC2SE	F 8, 2	7	33- 40
FEVISE	F 8, 2	7	41- 48
FEVISE	F 8, 2	7	49- 56
FVC1SE	F 8, 2	7	57- 64
FVC2SE	F 8, 2	7	65- 72
CVSVISE	F 8, 2	7	73- 80
CVSVISE	F 8, 2	7	81- 88
FEFVISE	F 8, 2	7	89- 96
FEFVISE	F 8, 2	7	97- 104

THE INPUT FORMAT PROVIDES FOR 87 VARIABLES. 87 WILL BE READ
IT PROVIDES FOR 7 RECORDS (*CARDS*) PER CASE. A MAXIMUM OF 104 *COLUMNS* ARE USED ON A RECORD.

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INPUT MEDIUM   DISK
N OF CASES    97
MISSING VALUES CIGNO CIGTOT(0)/IVFP1MN TO FEFV3SE(999)
              (SMGCAT FQ 0) CIGNO = 0
              (SMGCAT FQ 0) CIGTOT = 0
IF             V1=IVFP1MN TO FEFV3MX
DO REPEAT      /V2=IVFP1SD TO FEFV3MN
                /V3=IVFP1SE TO FEFV3SE
                (V1 EQ 0) V3=994
                (V1 EQ 0) V2=999
                (V1 EQ 0) V1=999
END REPEAT
COMPUTE
VALUE LABELS  STATUS=SMGCAT+2*SEX
              STATUS(0)=MALE  NON SMKG(1)FEMALE  SMOKING(2)MALE  NON SMOKING
              (1)MALE  SMOKING
PRINT FORMATS INIT(A)/WEIGHT(1)/HEIGHT IVFP1MN TO FEFV3SE(2)
TASK NAME
SELECT IF
DISCRIMINANT  DRUG ED 1
              GROUPS=SMGCAT(0 1)/VARIABLES=V25AR1MN V25HE1MN V40AR1MN V40HE1MN
              CV1NN V25FR1MN V40PR1MN IVFP1MN V25AR3MX V25HE3MX V40AR3MX

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PATIENT : PG SEX : F AGE : 22 HEIGHT : 1.65M WEIGHT : 60.4KG SMOKER? : Y CIGARETTES SMOKED : 7000 TOTAL : 43000

PATIENT : PG SEX : M AGE : 31 HEIGHT : 1.70M WEIGHT : 60.4KG SMOKER? : Y CIGARETTES SMOKED : 7000 TOTAL : 43000

7-12	2.70	3.09	2.31	1.70	3.42	4.19	3.42	.28	*.05	*.11	*.00	*.55	*.55
7-13	2.64	3.86	4.08	3.96	3.09	4.56	4.61	.60	*.28	.50	*.35	*.34	
7-12	2.26	4.40	3.31	3.09	3.20	5.29	4.85	.78	*.50	*.71	*.52	*.66	*.44
0916	2												
7-31	1.89	2.68	2.71	3.76	2.66	3.65	3.65	.14	*.18	*.03	*.37	*.06	*.78
7-14	1.77	3.32	3.21	3.21	3.21	5.43	4.76	.36	*.81	.29	.48	*.55	
2-43	2.71	3.76	2.44	3.54	2.44	4.87	3.65	.61	*.10	*.38	*.50	*.66	*.66
PATIENT	3												
7-11	2.71	3.21	3.54	3.43	3.54	4.76	4.87	.17	*.28	*.19	*.38	*.78	*.42
7-48	2.88	3.90	3.71	3.54	4.21	5.43	5.43	.39	*.65	.53	*.29	*.49	*.42
7-33	2.66	3.65	2.99	3.43	3.13	5.12	4.54	.32	*.85	.37	*.66	*.66	*.78
0916	4												
1-75	1.86	2.48	3.73	3.51	3.29	3.62	4.72	.63	1.01	.03	*.43	*.65	*.14
7-13	2.19	2.98	2.63	3.40	3.40	3.95	4.83	.15	*.20	.16	*.42	*.47	*.25
1-97	2.08	3.29	3.73	3.40	3.29	5.15	5.81	.67	*.79	.51	*.77	*.66	*.66

PATIENT : PG SEX : M AGE : 36 HEIGHT : 1.83M WEIGHT : 70.4KG SMOKER? : Y CIGARETTES SMOKED : 7000 TOTAL : 43000

PATIENT : PG SEX : M AGE : 36 HEIGHT : 1.83M WEIGHT : 70.4KG SMOKER? : Y CIGARETTES SMOKED : 7000 TOTAL : 43000

2-47	2.98	3.64	2.98	5.29	5.07	6.72	5.07	.27	*.00	*.27	*.00	2.76	2.53
4-09	2.20	3.09	2.20	5.40	3.06	5.45	3.86	.00	*.00	*.00	*.00	1.98	
2-76	2.64	3.75	2.64	4.85	4.74	7.71	4.74	.36	*.00	*.59	*.00	2.68	2.76
PATIENT	2												
7-62	2.07	2.95	2.29	4.58	1.71	6.11	4.80	.13	*.11	*.18	*.29	1.64	2.07
3-76	1.96	3.48	2.51	5.56	4.34	7.31	5.49	.10	*.28	.31	*.46	2.40	2.07
2-71	2.40	3.49	2.73	4.58	4.80	7.31	6.00	.28	*.14	*.60	*.28	1.53	1.96
0916	3												
4-07	2.74	3.29	3.62	4.81	4.91	6.58	7.35	.07	*.32	*.16	*.49	2.30	12.94
3-95	3.13	3.07	3.73	4.72	4.50	6.69	7.68	*.22	*.00	*.42	*.71	2.19	13.16
4-06	2.52	3.18	3.40	4.81	4.28	6.80	6.80	*.27	*.35	*.41	*.87	2.19	12.72
0916	4												
7-51	1.31	2.95	4.15	4.69	5.02	6.44	6.98	.18	*.12	*.37	*.39	1.05	8.40
3-16	3.71	3.16	3.71	4.91	4.91	6.55	7.20	.00	*.00	*.33	*.47	2.18	7.07
2-05	3.16	3.38	3.71	4.47	4.91	6.55	7.20	.15	*.17	*.47	*.36	9.46	10.04

PATIENT : G+C SEX : M AGE : 42 HEIGHT : 1.74M WEIGHT : 72.0KG SMOKER? : Y CIGARETTES SMOKED : 7000 TOTAL : 43000

PATIENT : PG SEX : M AGE : 42 HEIGHT : 1.70M WEIGHT : 60.4KG SMOKER? : Y CIGARETTES SMOKED : 7000 TOTAL : 43000

4-08	1.88	1.32	.98	1.43	1.32	2.53	1.54	.50	*.00	*.77	*.17	1.81	1.65
-49	.66	1.32	.77	1.76	1.21	2.31	1.54	.13	*.17	.31	*.27	1.21	1.65
-48	.33	1.21	.77	1.43	1.10	2.31	1.76	.38	*.33	.62	*.60	1.43	1.65
PATIENT	2												
-49	1.65	1.32	1.21	1.54	2.98	2.53	2.64	.13	*.27	*.64	*.11	1.31	1.43
-48	1.43	1.21	.99	1.54	2.76	2.42	2.42	.38	*.31	*.79	*.17	1.34	1.43
4-08	1.43	1.42	1.10	1.76	2.42	2.64	2.53	.50	*.23	.50	*.05	1.21	1.32
PATIENT	3												
1-01	1.38	1.24	1.57	1.80	2.25	2.47	1.26	.23	*.16	*.37	*.45	1.57	1.78
1-01	1.15	1.24	1.46	1.68	2.02	2.36	2.81	.23	*.08	*.10	*.37	1.57	1.77
-49	1.01	1.44	1.46	1.68	2.02	2.88	2.81	.89	*.45	*.49	*.39	1.57	1.45
PATIENT	4												
1-12	1.57	1.79	1.24	2.02	3.03	2.02	3.14	*.29	*.21	*.00	*.04	1.35	10.13
1-12	1.57	1.24	1.46	2.02	2.58	2.36	2.70	.11	*.07	*.17	*.05	1.68	10.67
1-12	1.24	1.35	1.91	1.68	2.25	2.47	1.59	.21	*.54	*.47	*.60	1.35	10.44

PATIENT : G+C SEX : M AGE : 42 HEIGHT : 1.74M WEIGHT : 72.0KG SMOKER? : Y CIGARETTES SMOKED : 7000 TOTAL : 43000

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PATIENT : #	SEX : M	AGE : 55	WEIGHT : 1,820	HEIGHT : 76,1KG	SUMMARY : N CIGARETTES SMOKED : ANNUAL : *** : TOTAL : ***
1-005	1				
2-38 2-17 2-17 2-28 4-78 8-12					
1-85 2-17 2-17 4-90 5-78					
2-30 2-06 2-30 2-39 5-41 5-32					
1-007	2				
2-42 3-17 3-18 2-93 4-88 6-01					
2-10 3-15 2-50 2-71 6-44 6-01					
2-81 3-18 2-71 2-61 6-44 6-01					
1-008	3				
2-18 1-78 2-29 2-07 3-02 3-09					
1-84 2-18 1-98 2-51 3-29 3-82					
1-85 2-07 2-18 2-29 2-95 3-49					
2-10 2-30 2-64 2-52 4-17 4-06					
2-30 2-37 2-74 3-05 4-17 5-17					
2-30 2-40 3-07 2-74 4-17 4-28					
PATIENT : HG SEX : F AGE : 22 WEIGHT : 1,568 HEIGHT : 66,1KG SUMMARY : N CIGARETTES SMOKED : ANNUAL : 6800 : TOTAL : 27400					
1-009	1				
1-48 1-16 1-74 1-09 3-04 2-17					
1-63 1-98 1-95 1-41 2-93 2-17					
1-30 1-30 1-44 1-41 2-87 2-50					
1-010	2				
1-44 1-51 2-16 2-48 2-92 2-16					
1-62 1-19 2-16 2-38 2-92 2-27					
1-49 1-30 2-46 1-94 2-27 2-48					
1-011	3				
1-48 2-28 1-19 1-63 2-19 3-58					
1-40 1-85 1-40 2-19 2-61 3-15					
1-30 1-85 1-41 1-74 2-81 3-26					
1-012	4				
1-41 1-52 1-95 2-19 2-50 3-15					
1-40 1-63 1-82 2-85 2-92 3-04					
1-36 1-63 1-82 1-95 2-81 2-93					
PATIENT : M SEX : M AGE : 41 HEIGHT : 1,908 WEIGHT : 74,2KG SUMMARY : N CIGARETTES SMOKED : ANNUAL : *** : TOTAL : ***					
1-013	1				
1-94 1-98 1-98 2-42 4-19 3-64					
2-89 1-87 2-98 2-09 4-04 3-64					
2-20 2-20 2-31 1-87 4-41 4-08					
1-014	2				
2-86 2-21 2-21 2-66 4-10 4-98					
2-21 1-85 2-33 4-10 4-76 4-32					
1-93 2-10 2-99 4-10 6-09 4-87					
1-015	3				
1-85 1-81 1-64 1-78 3-19 2-95					
1-83 1-83 1-64 1-75 3-27 3-38					
1-78 1-85 2-07 2-62 3-00 3-27					

PATIENT : JS SEX : F AGE : 23 HEIGHT : 1.62kg WEIGHT : 52.1kg SHOEsz : 4 CIGARETTES SMOKED : ANNUAL : *** : TOTAL :

height 1.62kg 2.28 1.95 2.02 2.03 3.60 1.47 .50 .38 *31 *18 *67 *65
 1.52 1.41 2.28 1.95 2.02 2.03 3.58 1.47 .50 .38 *31 *18 *67 *65
 1.74 1.52 2.28 1.95 2.02 2.03 3.58 1.47 .50 .38 *31 *18 *67 *65
 1.63 1.52 2.39 2.28 2.03 3.15 4.02 3.69 .47 .50 .37 *31 *18 *67 *54
 height 2 2.03 1.91 2.69 2.58 2.58 1.83 4.73 3.76 .25 *67 *35 *64
 2.15 1.93 2.69 3.22 2.79 4.41 3.44 .34 *64 .39 *66 .64
 1.41 1.91 2.88 3.01 3.12 2.47 4.41 3.44 .34 *64 .39 *66 .64
 height 3 2.17 2.71 7.82 3.08 3.69 4.02 4.23 *17 *35 *67 *35 *75
 1.48 2.78 2.28 2.78 3.08 3.69 4.02 4.23 *17 *35 *67 *35 *75
 1.63 1.68 2.28 2.71 2.93 3.15 4.56 4.89 *40 *46 *56 *55 *75
 height 4 1.75 1.85 2.40 1.75 2.95 3.42 3.27 *37 *08 *35 *11 *65
 1.75 1.63 2.40 2.29 2.40 2.84 4.15 3.71 *37 *40 *73 *31 *65
 1.42 1.63 2.29 2.18 2.51 2.51 4.15 3.71 *31 *65 *46 *87
 height 5 1.75 1.86 2.40 1.09 4.49 4.05 4.23 *17 *35 *67 *35 *75

PATIENT : ET SEX : F AGE : 26 HEIGHT : 1.69kg WEIGHT : 52.1kg SHOEsz : 4 CIGARETTES SMOKED : ANNUAL : *** : TOTAL :

height 1 1.75 2.18 1.85 2.05 2.05 3.60 1.47 .50 .38 *31 *18 *67 *65
 1.75 1.20 2.67 1.53 2.05 2.05 3.58 1.47 .50 .38 *31 *18 *67 *65
 1.63 1.53 2.40 1.09 4.49 4.05 4.23 *17 *35 *67 *35 *75
 height 2 2.05 2.62 2.62 2.62 3.00 3.93 4.58 4.91 *71 *42 *27 *25 *64
 1.81 1.68 2.18 2.18 3.16 3.82 4.47 5.07 *25 *55 *41 *31 *65
 1.76 2.01 2.07 2.18 3.49 4.04 4.36 5.07 *35 *55 *41 *25 *65
 height 3 1.75 2.07 2.18 3.71 3.19 3.92 4.04 *18 *25 *03 *16 *65
 1.75 1.85 2.29 3.60 3.49 4.53 4.67 *06 *17 *09 *28 *65
 1.48 1.65 2.18 2.07 3.60 3.49 4.67 4.36 *18 *12 *24 *25 *65
 height 4 1.75 1.86 2.40 1.62 2.74 3.40 3.62 *08 *39 *24 *37 *65

PATIENT : PE SEX : M AGE : 64 HEIGHT : 1.71kg WEIGHT : 77.4kg SHOEsz : 7 CIGARETTES SMOKED : ANNUAL : *** : TOTAL :

height 1 1.75 1.65 1.47 1.39 2.18 3.06 4.36 *40 1.58 *09 1.00 *100 *100
 1.49 1.68 1.69 1.64 1.66 2.07 3.91 4.36 *27 1.16 *26 1.11 *11
 1.47 1.68 1.69 1.64 1.66 2.07 3.91 4.36 *27 1.16 *26 1.11 *11
 1.26 1.55 1.67 1.64 1.27 2.51 3.60 4.04 *27 1.58 *30 *61 *61
 height 2 1.42 1.10 1.21 1.25 3.19 1.86 3.73 4.06 *08 *59 *17 1.18 *18
 1.48 1.10 1.21 1.25 3.19 1.86 3.73 4.06 *08 *59 *17 1.18 *18
 1.48 1.10 1.21 1.25 3.19 1.86 3.73 4.06 *08 *59 *17 1.18 *18
 height 3 1.75 1.99 2.09 2.09 2.94 2.94 2.87 1.98 *10 *10 *19 *34 1.76 *84
 1.49 1.71 1.10 1.21 1.37 2.11 3.69 4.32 *18 *00 *83 *40 1.54 *54
 1.21 1.66 1.43 1.66 1.87 2.20 3.42 3.69 *18 *00 *83 *40 1.54 *54
 height 4 1.10 1.10 1.21 1.21 1.53 2.31 2.74 3.40 3.62 *08 *39 *24 *37 *65
 1.10 1.10 1.21 1.21 1.53 2.31 2.74 3.40 3.62 *08 *39 *24 *37 *65
 height 5 1.75 1.99 2.09 2.09 2.94 2.94 2.87 1.98 *10 *10 *19 *34 1.76 *84
 1.49 1.71 1.10 1.21 1.37 2.11 3.69 4.32 *18 *00 *83 *40 1.54 *54
 1.21 1.66 1.43 1.66 1.87 2.20 3.42 3.69 *18 *00 *83 *40 1.54 *54
 height 6 1.10 1.10 1.21 1.21 1.53 2.31 2.74 3.40 3.62 *08 *39 *24 *37 *65
 1.10 1.10 1.21 1.21 1.53 2.31 2.74 3.40 3.62 *08 *39 *24 *37 *65
 height 7 1.75 1.99 2.09 2.09 2.94 2.94 2.87 1.98 *10 *10 *19 *34 1.76 *84
 1.49 1.71 1.10 1.21 1.37 2.11 3.69 4.32 *18 *00 *83 *40 1.54 *54
 1.21 1.66 1.43 1.66 1.87 2.20 3.42 3.69 *18 *00 *83 *40 1.54 *54
 height 8 1.10 1.10 1.21 1.21 1.53 2.31 2.74 3.40 3.62 *08 *39 *24 *37 *65
 1.10 1.10 1.21 1.21 1.53 2.31 2.74 3.40 3.62 *08 *39 *24 *37 *65

PATIENT : 55 SEX : F AGE : 24 HEIGHT : 1.72M WEIGHT : 55.0KG Smoker? : N CIGARETTES SMOKED : ANNUAL : *** : TOTAL : ***

100G 1	1.44	3.65	3.10	3.54	3.32	4.38	4.34	.47	.27	.41	.37	.66	.55
100G 2	2.47	3.43	3.21	3.51	3.54	4.43	4.43	.16	.11	.25	.25	.55	.44
100G 3	1.71	2.58	3.41	3.45	3.45	4.87	4.87	.50	.61	.47	.67	.55	.55
100G 4	2.11	2.13	3.05	3.76	3.12	3.21	4.87	.50	.50	.61	.47	.67	.55
100G 5	2.09	2.21	3.54	2.77	3.68	3.21	5.32	4.12	.18	.25	.37	.35	1.00
100G 6	2.48	2.71	3.65	2.88	3.54	4.32	4.32	.27	.10	.41	.39	.44	.49
100G 7	2.21	2.21	3.81	2.77	3.68	4.32	4.32	.10	.28	.28	.21	.55	.78
100G 8	2.05	2.43	3.21	2.68	3.65	4.34	4.34	.26	.24	.24	.34	.44	.44
100G 9	2.55	2.43	3.21	2.68	3.65	4.34	4.34	.12	.12	.12	.10	.44	.44
100G 10	2.41	2.73	3.21	3.10	3.54	4.54	4.76	.20	.32	.23	.50	.50	.50
100G 11	2.31	2.34	3.65	2.99	3.54	4.10	5.32	4.87	.50	.50	.10	.44	.44
100G 12	2.71	2.66	2.77	3.10	3.37	4.10	4.12	4.54	.25	.17	.30	.11	.33
100G 13	2.55	2.77	3.10	3.50	3.10	4.54	4.98	.09	.12	.28	.21	.33	.33
100G 14	2.44	2.68	3.32	3.54	3.32	4.21	4.87	.09	.36	.23	.47	.11	.44
100G 15	2.44	2.68	3.32	3.54	3.32	4.21	4.87	.09	.36	.23	.47	.11	.44

PATIENT : JC SEX : M AGE : 55 HEIGHT : 1.79M WEIGHT : 71.6KG Smoker? : N CIGARETTES SMOKED : ANNUAL : *** : TOTAL : ***														
100G 1	1.82	4.15	3.16	3.70	3.44	4.80	4.80	.03	.17	.03	.03	1.53	.87	
100G 2	2.47	3.60	3.46	3.56	3.46	4.86	4.86	.89	.89	.89	.89	1.53	.87	
100G 3	1.58	4.58	4.47	3.71	5.87	4.44	5.67	.02	.19	.28	.28	1.42	.87	
100G 4	1.83	5.29	5.73	6.04	6.95	6.64	6.64	.00	.52	.44	.44	1.54	1.10	
100G 5	1.81	5.06	5.51	5.38	7.38	6.69	6.69	.22	.00	.00	.00	1.54	1.21	
100G 6	1.81	4.28	7.87	6.61	6.61	6.44	6.44	.05	.82	.44	.44	1.54	.99	
100G 7	1.41	6.69	6.03	7.46	7.46	9.43	9.43	.22	.17	.28	.26	1.54	1.64	
100G 8	1.40	5.20	7.02	6.31	6.91	7.47	11.59	9.05	.28	.54	.48	.48	1.54	1.64
100G 9	1.46	5.49	6.69	4.50	7.68	7.24	11.40	9.31	.05	.18	.15	.15	1.54	1.71
100G 10	1.18	6.11	5.82	6.11	5.89	8.40	8.40	.06	.39	.04	.42	.20	.29	
100G 11	1.93	5.29	4.91	4.91	4.47	5.02	7.09	10.47	.59	.17	.09	.20	.17	
100G 12	1.26	6.66	6.98	4.91	7.20	7.08	10.15	.56	.39	.44	.41	.20	.09	
100G 13	1.44	5.49	5.49	5.49	5.49	5.49	5.49	.00	.00	.00	.00	1.40	.07	
100G 14	1.44	5.49	5.49	5.49	5.49	5.49	5.49	.00	.00	.00	.00	1.40	.07	
100G 15	1.76	2.42	2.64	3.42	2.76	4.10	4.10	.10	.50	.26	.68	.88	.88	
100G 16	1.76	2.42	2.64	3.42	2.76	4.10	4.10	.10	.50	.26	.68	.88	.88	
100G 17	2.31	2.09	2.84	1.87	3.75	3.64	4.41	3.54	.14	.11	.18	.03	1.21	
100G 18	2.31	1.87	2.53	2.84	5.63	2.98	4.63	6.63	.10	.41	.27	.53	.97	
100G 19	2.44	1.87	2.42	2.20	6.10	7.98	4.85	4.19	.08	.18	.13	.41	1.21	
100G 20	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 21	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 22	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 23	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 24	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 25	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 26	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 27	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 28	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 29	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 30	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 31	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 32	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 33	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 34	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 35	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 36	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 37	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 38	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 39	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 40	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 41	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 42	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 43	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 44	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 45	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 46	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 47	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 48	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 49	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 50	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 51	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 52	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 53	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 54	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 55	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 56	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 57	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 58	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 59	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 60	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 61	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 62	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 63	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 64	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 65	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 66	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 67	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 68	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 69	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 70	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 71	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 72	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 73	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 74	2.44	1.87	2.42	2.20	6.10	7.98	4.85	.00	.00	.00	.00	1.21	.98	
100G 75	2.44	1.87	2.42	2.20	6.10	7.98	4							

WEIGHT : 1.65H HEIGHT : 61.98G SNIPPERZ

PATIENT : LO SEX : F AGE : 25 HEIGHT : 1.698 WEIGHT : 63.9KG SMOKE# : N CIGARETTES SMOKED : ANNUAL : *** : TOTAL : ***

OPNG 1	2.84	.87	2.94	1.03	4.48	.40	4.52	1.43	.30	.19	.12	.17	1.21	.46
	.20	.48	.39	.52	.28	.11	.30	.10	.10	.08	0	.11	.11	.59
	.11	.10	.23	.26	.16	.19	.29	.28	.07	.11	0	.03	.03	.07
OPNG 2	1.18	1.04	5.29	1.07	5.11	.88	7.40	1.07	.67	.57	.46	.34	1.51	.40
	.39	.63	.39	.89	.23	.85	.68	.22	.09	.51	.08	.25	.17	.29
	.23	.21	.23	.32	.04	.39	.45	.05	.03	.05	.07	.10	.06	.07
OPNG 3	3.13	1.18	4.49	1.32	4.89	1.06	6.68	1.47	.55	.17	.37	.11	1.54	.11
	.23	.74	.66	.81	.17	.92	.54	.88	.14	.03	.12	.07	.00	.29
	.13	.10	.50	.32	.10	.15	.31	.35	.10	.07	.03	.00	.05	.27
OPNG 4	3.46	.66	0	0	6.50	1.10	0	0	0	0	0	0	0	.11
	.11	.26	0	0	.81	.70	0	0	0	0	0	0	0	.07
	.06	.20	0	0	.35	.40	0	0	0	0	0	0	0	.03

PATIENT : LO SEX : M AGE : 39 HEIGHT : 1.944 WEIGHT : 75.0KG SMOKE# : N CIGARETTES SMOKED : ANNUAL : *** : TOTAL : ***

OPNG 1	7.94	1.62	3.27	.96	6.76	2.68	7.05	2.53	.17	.65	.13	43	1.61	.29
	.28	.44	.34	.67	.28	.60	.69	1.39	.06	.26	.06	.23	.17	.26
	.18	.13	.19	.24	.16	.07	.40	.64	.04	.06	.04	.13	.10	.04
OPNG 2	2.48	2.29	2.72	.19	4.34	.26	6.28	.77	.26	.22	.46	.24	1.72	.51
	.08	.07	.17	.11	.32	.0	.79	.40	.09	.00	.30	.10	.17	.04
	.04	.11	.10	.08	.18	.13	.46	.22	.05	.11	.17	.07	.10	.04
OPNG 3	1.40	1.12	2.32	.49	3.63	1.99	6.10	.75	.46	.37	.69	.53	2.14	.63
	.12	.75	.64	.51	.23	.68	.97	.48	.42	.42	.46	.40	.11	.27
	.07	.37	.08	.14	.25	.56	.29	.34	.05	.19	.02	.17	.08	.04
OPNG 4	1.92	1.18	2.45	.22	4.34	.31	5.90	.78	.35	.18	.36	.31	2.15	.05
	.16	.15	.40	.00	.22	.04	.67	.15	.24	.10	.17	.03	.07	.07
	.04	.23	.11	.11	.19	.39	.32	.14	.08	.10	.14	.04	.06	.11
OPNG 5	2.42	.85	3.31	1.44	3.20	.66	3.93	.48	.39	.21	.24	.21	.59	.15
	.14	.33	.19	.15	.31	.37	.23	.15	.27	.21	.13	.09	.13	.15
	.22	.17	.11	.16	.19	.15	.13	.28	.14	0	.07	.07	0	
OPNG 6	3.28	1.15	4.40	2.07	4.12	.89	5.21	1.78	.12	0	.38	.24	.85	.22
	.13	1.03	.45	.66	.29	.78	.88	.152	.07	0	.08	.24	.11	.11
	.07	.11	.26	.22	.17	.06	.51	.16	.05	0	.06	0	.06	
OPNG 7	2.52	.37	2.92	.52	4.01	.44	5.04	.86	.21	.17	.26	.17	6.0	.67
	.13	.22	.16	.26	.20	.30	.39	.11	.16	.08	.11	.01	.13	.04
	.07	.10	.21	.14	.11	.10	.23	.17	.09	.07	.07	.04	.14	.07
OPNG 8	1.83	.15	2.05	.15	3.80	.18	4.19	.55	.18	.07	.25	.16	.59	.04
	.06	.11	.23	.15	.21	.13	.11	.11	.08	.02	.04	.04	.11	.05
	.03	.14	.11	.14	.06	.19	.24	.06	.05	.04	.04	.04	.06	.04

PATIENT : LO SEX : F AGE : 32 HEIGHT : 1.59M WEIGHT : 54.1KG SMOKE# : N CIGARETTES SMOKED : ANNUAL : *** : TOTAL : ***

OPNG 1	2.42	.85	3.31	1.44	3.20	.66	3.93	.48	.39	.21	.24	.21	.59	.15
	.14	.33	.19	.15	.31	.37	.23	.15	.27	.21	.13	.09	.13	.15
	.22	.17	.11	.16	.19	.15	.13	.28	.14	0	.07	.07	0	
OPNG 2	3.28	1.15	4.40	2.07	4.12	.89	5.21	1.78	.12	0	.38	.24	.85	.22
	.13	1.03	.45	.66	.29	.78	.88	.152	.07	0	.08	.24	.11	.11
	.07	.11	.26	.22	.17	.06	.51	.16	.05	0	.06	0	.06	
OPNG 3	2.52	.37	2.92	.52	4.01	.44	5.04	.86	.21	.17	.26	.17	6.0	.67
	.13	.22	.16	.26	.20	.30	.39	.11	.16	.08	.11	.01	.13	.04
	.07	.10	.21	.14	.11	.10	.23	.17	.09	.07	.07	.04	.14	.07
OPNG 4	1.83	.15	2.05	.15	3.80	.18	4.19	.55	.18	.07	.25	.16	.59	.04
	.06	.11	.23	.15	.21	.13	.11	.11	.08	.02	.04	.04	.11	.05
	.03	.14	.11	.14	.06	.19	.24	.06	.05	.04	.04	.04	.06	.04

PATIENT : LO SEX : M AGE : 32 HEIGHT : 1.59M WEIGHT : 54.1KG SMOKE# : N CIGARETTES SMOKED : ANNUAL : *** : TOTAL : ***

OPNG 1	2.42	.85	3.31	1.44	3.20	.66	3.93	.48	.39	.21	.24	.21	.59	.15
	.14	.33	.19	.15	.31	.37	.23	.15	.27	.21	.13	.09	.13	.15
	.22	.17	.11	.16	.19	.15	.13	.28	.14	0	.07	.07	0	
OPNG 2	3.28	1.15	4.40	2.07	4.12	.89	5.21	1.78	.12	0	.38	.24	.85	.22
	.13	1.03	.45	.66	.29	.78	.88	.152	.07	0	.08	.24	.11	.11
	.07	.11	.26	.22	.17	.06	.51	.16	.05	0	.06	0	.06	
OPNG 3	2.52	.37	2.92	.52	4.01	.44	5.04	.86	.21	.17	.26	.17	6.0	.67
	.13	.22	.16	.26	.20	.30	.39	.11	.16	.08	.11	.01	.13	.04
	.07	.10	.21	.14	.11	.10	.23	.17	.09	.07	.07	.04	.14	.07
OPNG 4	1.83	.15	2.05	.15	3.80	.18	4.19	.55	.18	.07	.25	.16	.59	.04
	.06	.11	.23	.15	.21	.13	.11	.11	.08	.02	.04	.04	.11	.05
	.03	.14	.11	.14	.06	.19	.24	.06	.05	.04	.04	.04	.06	.04

PATIENT : LO SEX : F AGE : 32 HEIGHT : 1.59M WEIGHT : 54.1KG SMOKE# : N CIGARETTES SMOKED : ANNUAL : *** : TOTAL : ***

OPNG 1	2.42	.85	3.31	1.44	3.20	.66	3.93	.48	.39	.21	.24	.21	.59	.15
	.14	.33	.19	.15	.31	.37	.23	.15	.27	.21	.13	.09	.13	.15
	.22	.17	.11	.16	.19	.15	.13	.28	.14	0	.07	.07	0	
OPNG 2	3.28	1.15	4.40	2.07	4.12	.89	5.21	1.78	.12	0	.38	.24	.85	.22
	.13	1.03	.45	.66	.29	.78	.88	.152	.07	0	.08	.24	.11	.11
	.07	.11	.26	.22	.17	.06	.51	.16	.05	0	.06	0	.06	
OPNG 3	2.52	.37	2.92	.52	4.01	.44	5.04	.86	.21	.17	.26	.17	6.0	.67
	.13	.22	.16	.26	.20	.30	.39	.11	.16	.08	.11	.01	.13	.04
	.07	.10	.21	.14	.11	.10	.23	.17	.09	.07	.07	.04	.14	.07
OPNG 4	1.83	.15	2.05	.15	3.80	.18	4.19	.55	.18	.07	.25	.16	.59	.04
	.06	.11	.23	.15	.21	.13	.11	.11	.08	.02	.04	.04	.11	.05
	.03	.14	.11	.14	.06	.19	.24	.06	.05	.04	.04	.04	.06	.04

PATIENT : LO SEX : M AGE : 32 HEIGHT : 1.59M WEIGHT : 54.1KG SMOKE# : N CIGARETTES SMOKED : ANNUAL : *** : TOTAL : ***

OPNG 1	2.42	.85	3.31	1.44	3.20	.66	3.93	.48	.39	.21	.24	.21	.59	.15
	.14	.33	.19	.15	.31	.37	.23	.15	.27	.21	.13	.09	.13	.15
	.22	.17	.11	.16	.19	.15	.13	.28	.14	0	.07	.07	0	
OPNG 2	3.28	1.15	4.40	2.07	4.12	.89	5.21	1.78	.12	0	.38	.24	.85	.22
	.13	1.03	.45	.66	.29	.78	.88	.152	.07	0	.08	.24	.11	.11
	.07	.11	.26	.22	.17	.06	.51	.16	.05	0	.06	0	.06	
OPNG 3	2.52	.37	2.92	.52	4.01	.44	5.04	.86	.21	.17	.26	.17	6.0	.67
	.13	.22	.16	.26	.20	.30	.39	.11	.16	.08	.11	.01	.13	.04
	.07	.10	.21	.14	.11	.10	.23	.17	.09	.07	.07	.04	.14	.07
OPNG 4	1.83	.15	2.05	.15	3.80	.18	4.19	.55	.18	.07	.25	.16	.59	.04
	.06	.11	.23	.15	.21	.13	.11	.11	.08					

FATIGUE : JIFC		SEX : M		AGE : 55		WEIGHT : 1,82kg		SMOKER : 76,1kg		NON-SMOKER : 71,8kg		CIGARETTES SMOKED : ANNUAL : ***		TOTAL : ***					
1000G 1																			
2,10	*.07	2,24	*.15	5,07	*.68	0	0	*.13	*.08	0	0	1,90	*.40						
*.17	*.04	*.11	*.04	*.13	*.40	0	0	*.04	*.03	0	0	*.06	*.47						
*.10	*.04	*.07	*.06	*.19	*.14	0	0	*.03	*.06	0	0	*.04	*.04						
1000G 2																			
2,61	*.86	2,79	*.18	0	0	0	0	*.07	0	0	0	0	2,02	*.91					
*.27	*.85	*.33	*.04	0	0	0	0	*.04	0	0	0	*.16	*.43						
*.13	*.11	*.19	*.04	0	0	0	0	*.03	0	0	0	*.04	*.06						
1000G 3																			
*.29	2,14	3,37	3,35	*.41	4,62	*.84	*.14	*.04	*.40	*.16	2,07	*.54	*.79	1,34	*.84	2,40			
*.11	*.17	*.15	*.14	*.25	*.23	*.55	*.08	*.06	*.28	*.03	*.22	*.92	*.06	*.51	*.14	*.30			
*.13	*.10	*.13	*.25	*.11	*.13	*.29	*.05	*.02	*.14	*.07	*.11	*.53	*.04	*.19	*.68	2,12			
1000G 4																*.64			
*.40	2,97	4,40	4,10	*.18	5,26	*.87	*.25	*.15	*.29	*.21	2,01	*.79	11,66	2,01	*.51	17,22	*.61		
*.17	*.11	*.25	*.26	*.13	*.07	*.07	*.07	*.10	*.14	*.15	*.06	*.12	*.86	*.06	*.72	*.51	76,98		
*.15	*.15	*.07	*.07	*.06	*.27	*.22	*.04	*.05	*.08	*.03	*.04	*.18	*.15	*.04	*.15	*.29	1,00	0	
1000G 5																2,26			
FATIGUE : M	AGE : F	SEX : F	AGE : 22	WEIGHT : 1,59kg	WEIGHT : 66,1kg	SMOKER : 66,1kg	NON-SMOKER : 66,1kg	CIGARETTES SMOKED : ANNUAL : ***	SMOKER : 66,1kg	NON-SMOKER : 66,1kg	CIGARETTES SMOKED : ANNUAL : ***	SMOKER : 66,1kg	NON-SMOKER : 66,1kg	CIGARETTES SMOKED : ANNUAL : ***	SMOKER : 66,1kg	NON-SMOKER : 66,1kg	TOTAL : ***		
1,20	*.72	1,48	*.76	2,93	*.76	3,48	1,70	*.20	*.24	*.17	*.59	1,30	*.21						
*.13	*.55	*.15	*.54	*.11	*.65	*.10	*.40	0	*.06	*.13	*.19	0	*.04						
*.13	*.09	*.10	*.06	*.11	*.21	*.44	0	*.18	*.10	*.21	0	*.03							
1000G 6																			
*.28	*.46	2,27	*.31	2,10	*.54	*.42	0	*.40	*.60	*.29	0	*.06	*.31						
*.27	*.32	*.18	0	*.38	*.40	*.27	0	*.34	*.31	*.28	0	*.04	*.09						
*.16	*.09	*.11	*.17	*.22	*.09	*.13	0	*.19	*.15	*.18	0	*.04	*.09						
1000G 7																			
*.16	*.09	*.11	*.17	*.22	*.09	*.13	0	*.19	*.15	*.18	0	*.04	*.09						
1000G 8																			
1,19	*.99	1,40	1,08	2,54	1,04	3,01	1,77	*.14	*.14	*.18	*.34	*.17	*.72	*.26	*.81	*.19	*.45	*.40	
*.18	*.06	*.11	*.62	*.13	*.79	*.18	1,30	*.09	*.14	*.10	*.13	*.04	*.36	*.06	*.34	*.24	*.19	*.86	
*.11	*.06	*.24	*.07	*.13	*.22	*.32	*.07	0	*.06	*.14	*.10	*.04	*.24	*.23	*.19	*.10	*.56	1,42	
1000G 9																			
*.16	*.33	1,52	*.43	2,61	*.43	3,29	1,05	*.17	*.14	*.27	*.16	*.65	*.22	*.43	*.16	*.25	*.13	*.38	
*.07	*.26	0	*.36	*.72	*.22	*.16	*.58	*.01	*.04	*.05	*.11	*.11	*.15	*.44	*.04	*.17	*.13	*.16	
0	*.07	0	*.03	*.17	*.16	*.09	*.25	*.00	*.05	*.03	*.04	*.06	*.04	*.25	*.17	*.07	*.17	*.02	
P 1000T 1																			
1000G 1																			
2,00	*.22	2,42	*.55	4,23	*.59	5,47	*.71	*.24	*.12	*.30	*.14	*.61	*.26						
*.11	*.07	*.51	*.30	*.17	*.44	*.63	*.59	*.27	*.07	*.20	*.06	*.16	*.15						
*.16	*.10	*.29	*.16	*.10	*.15	*.36	*.07	*.19	*.05	*.11	*.07	*.04	*.11						
1000G 3																			
*.29	*.74	7,51	1,59	4,98	*.66	5,87	2,66	*.28	*.18	*.35	*.62	1,66	*.11	*.81	*.50	*.45	*.10	*.50	
*.14	*.33	*.42	1,11	1,01	*.26	*.29	2,22	*.32	*.66	*.07	*.37	*.16	*.11	*.08	*.51	*.19	*.73	*.94	
*.20	*.24	*.48	*.59	*.20	*.17	*.38	*.22	*.42	*.05	*.14	*.11	*.05	*.05	*.05	*.48	*.19	*.77	*.49	
*.06	*.4	*.40	1,78	*.84	3,42	*.47	*.84	1,02	*.13	*.20	*.29	*.38	*.07	*.62	*.71	*.10	*.33	*.10	
*.71	*.18	*.25	*.26	*.17	*.22	*.61	*.33	*.08	*.17	*.11	*.20	*.06	*.18	*.17	*.48	*.19	*.44	*.47	0
*.18	*.18	*.25	*.26	*.17	*.22	*.61	*.33	*.08	*.17	*.11	*.20	*.06	*.18	*.17	*.48	*.19	*.44	*.47	0
*.07	*.14	*.29	*.10	*.13	*.16	*.09	*.05	*.05	*.06	*.06	*.11	*.08	*.05	*.05	*.11	*.10	*.19	*.10	

— 4.0 190 STANDARD : V CIGARETTES Smokes : Auskin : 2000

case : F BACK : 11 UPRIGHT : 1.7MM WEIGHT : 66.7KG Dimensions :

Age : 41 Height : 1.75m Weight : 64kg

STATION : 35 SEX : F AGE : 23 HEIGHT : 5'2" WEIGHT : 162# GENDER : MALE STATUS : SINGLE : ANNUAL : 000 : TOTAL : 000

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Table 1. Summary of the results of the simulation study. The first column shows the sample size, the second column shows the true parameter value, the third column shows the estimated parameter value, the fourth column shows the standard error of the estimate, and the fifth column shows the absolute bias.

FIG. 2. The effect of the concentration of the polymer solution on the viscosity of the polymer solution.

Table 1. Summary of the results of the experiments on the effect of the addition of organic acids on the properties of the polyacrylate gel.

DATA : ET SEX : F AGE : 26 HEIGHT : 1.69# WEIGHT : 49.8KG SURVEY : V CHARTS SURVEY : V

THEORY OF THE CLOUDS OF VARIOUS TYPES

Table 1. Summary of the results of the two experiments.

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PIGMENTED SENSITIZED AURAT. I TOTAL: 76800

Table 1. Summary of the results of the simulation study.

THE INFLUENCE OF THE ENVIRONMENT ON THE GROWTH OF COTTON 11

Table 1. Summary of the results of the simulation study.

100 -44 1 -14 -68 1 -76 77 3 -16 -47 -14 -43 -87 -61 1 -65 -11 0 11 0 0 0 0 0 0 0 0 0 0 0 0 0

C H A P T E R 4 ■ THE NATURE OF POLYMERIZATION

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PATIENT : 56 SEX : F AGE : 24 HEIGHT : 1.72m WEIGHT : 55.0kg SMOKE : N CIGARETTES : 0 ANNUAL : *** TOTAL : ***

PATIENT	56	SEX	F	AGE	24	HEIGHT	1.72m	WEIGHT	55.0kg	SMOKE	N	CIGARETTES	0	ANNUAL	***	TOTAL	***	
1	2.462	29	1.50	.40	3.47	.26	4.76	.44	.44	.37	.48	.24	.59	.18				
2	.117	-.01	-.25	-.15	-.13	-.11	-.20	-.01	-.17	-.01	-.11	-.04	-.06	-.01				
3	.110	-.17	-.15	-.20	-.07	-.10	-.17	-.24	-.10	-.15	-.07	-.11	-.04					
4	PATIENT	57	SEX	F	AGE	24	HEIGHT	1.72m	WEIGHT	55.0kg	SMOKE	N	CIGARETTES	0	ANNUAL	***	TOTAL	***
5	2.458	42	1.58	.81	3.77	.67	5.09	.77	.49	.12	.35	.14	.50	.50				
6	.106	-.17	-.06	-.17	-.06	-.17	-.20	-.05	-.17	-.08	-.07	-.04	-.05					
7	.110	-.14	-.04	-.11	-.11	-.11	-.05	-.03	-.04	-.05	-.05	-.06	-.06					
8	PATIENT	58	SEX	M	AGE	55	HEIGHT	1.79m	WEIGHT	71.6kg	SMOKE	N	CIGARETTES	0	ANNUAL	***	TOTAL	***
9	4.466	94	4.07	.91	7.01	0	7.82	3.02	.03	0	.25	0	1.49	.62				
10	.266	-.17	-.44	-.13	-.13	0	-.02	-.23	-.17	0	0	0	-.06	-.02				
11	.18	-.24	-.25	-.35	-.17	0	-.47	-.33	0	0	0	0	0					
12	PATIENT	59	SEX	M	AGE	55	HEIGHT	1.79m	WEIGHT	71.6kg	SMOKE	N	CIGARETTES	0	ANNUAL	***	TOTAL	***
13	4.466	94	4.07	.91	7.01	0	7.82	3.02	.03	0	.25	0	1.49	.62				
14	.266	-.19	-.49	-.19	-.19	0	-.02	-.23	-.17	0	0	0	-.06	-.02				
15	.111	-.61	-.63	-.55	-.49	-.31	0	-.12	-.08	0	0	0	-.06	-.02				
16	.156	-.07	-.36	-.81	-.27	-.41	0	0	-.08	0	0	0	0	0				
17	PATIENT	60	SEX	M	AGE	55	HEIGHT	1.79m	WEIGHT	71.6kg	SMOKE	N	CIGARETTES	0	ANNUAL	***	TOTAL	***
18	5.77	1.27	6.80	2.40	7.35	.88	10.71	2.38	.18	.16	.46	.11	1.43	.48				
19	.81	-.86	-.19	-.99	-.40	-.29	1.02	-.51	.12	.15	.17	-.16	.19	.21				
20	.279	-.31	.11	-.70	-.23	-.30	-.59	-.41	.07	.21	.10	-.11	0	.07				
21	PATIENT	61	SEX	M	AGE	55	HEIGHT	1.79m	WEIGHT	71.6kg	SMOKE	N	CIGARETTES	0	ANNUAL	***	TOTAL	***
22	5.111	2.00	5.13	1.85	5.09	1.31	7.45	3.02	.40	.01	.47	.62	1.27	.48				
23	.117	1.56	1.43	1.87	1.73	1.78	.63	2.33	.22	-.01	.10	.04	1.13	.07				
24	.111	-.33	.83	.60	.47	.99	.36	.53	.46	0	.06	.31	.07	.20				

PATIENT : 62 SEX : F AGE : 32 HEIGHT : 1.67m WEIGHT : 58.1kg SMOKE : N CIGARETTES : 0 ANNUAL : *** TOTAL : ***

PATIENT	62	SEX	F	AGE	32	HEIGHT	1.67m	WEIGHT	58.1kg	SMOKE	N	CIGARETTES	0	ANNUAL	***	TOTAL	***	
1	7.432	55	2.53	6.66	3.90	.92	4.63	1.10	.12	.29	.19	.36	1.14	.17				
2	.114	-.48	-.11	-.29	-.38	-.20	-.22	-.51	-.01	-.17	-.07	-.29	-.13	-.29				
3	.111	-.07	.08	-.22	-.22	-.20	-.22	-.13	-.12	-.02	-.11	-.04	-.07	.04				
4	PATIENT	63	SEX	F	AGE	32	HEIGHT	1.67m	WEIGHT	58.1kg	SMOKE	N	CIGARETTES	0	ANNUAL	***	TOTAL	***
5	7.432	55	2.53	6.66	3.90	.92	4.63	1.10	.12	.29	.19	.36	1.14	.17				
6	.114	-.48	-.11	-.29	-.38	-.20	-.22	-.51	-.01	-.17	-.07	-.29	-.13	-.29				
7	.111	-.07	.08	-.22	-.22	-.20	-.22	-.13	-.12	-.02	-.11	-.04	-.07	.04				
8	PATIENT	64	SEX	M	AGE	32	HEIGHT	1.67m	WEIGHT	58.1kg	SMOKE	N	CIGARETTES	0	ANNUAL	***	TOTAL	***
9	7.432	55	2.53	6.66	3.90	.92	4.63	1.10	.12	.29	.19	.36	1.14	.17				
10	.114	-.48	-.11	-.29	-.38	-.20	-.22	-.51	-.01	-.17	-.07	-.29	-.13	-.29				
11	.111	-.07	.08	-.22	-.22	-.20	-.22	-.13	-.12	-.02	-.11	-.04	-.07	.04				
12	PATIENT	65	SEX	M	AGE	32	HEIGHT	1.67m	WEIGHT	58.1kg	SMOKE	N	CIGARETTES	0	ANNUAL	***	TOTAL	***
13	7.432	55	2.53	6.66	3.90	.92	4.63	1.10	.12	.29	.19	.36	1.14	.17				
14	.114	-.48	-.11	-.29	-.38	-.20	-.22	-.51	-.01	-.17	-.07	-.29	-.13	-.29				
15	.111	-.07	.08	-.22	-.22	-.20	-.22	-.13	-.12	-.02	-.11	-.04	-.07	.04				
16	PATIENT	66	SEX	M	AGE	32	HEIGHT	1.67m	WEIGHT	58.1kg	SMOKE	N	CIGARETTES	0	ANNUAL	***	TOTAL	***
17	7.432	55	2.53	6.66	3.90	.92	4.63	1.10	.12	.29	.19	.36	1.14	.17				
18	.114	-.48	-.11	-.29	-.38	-.20	-.22	-.51	-.01	-.17	-.07	-.29	-.13	-.29				
19	.111	-.07	.08	-.22	-.22	-.20	-.22	-.13	-.12	-.02	-.11	-.04	-.07	.04				
20	PATIENT	67	SEX	M	AGE	32	HEIGHT	1.67m	WEIGHT	58.1kg	SMOKE	N	CIGARETTES	0	ANNUAL	***	TOTAL	***

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PATIENT #	AGE	SEX	WEIGHT	HEIGHT	1.8M WEIGHT	1.8M HEIGHT	Y	CHARTS DOWN
1001G 1	3.24	♂	6.02	2.93	4.41	0	0	0
	.94	♂	6.62	3.41	6.69	0	0	0
	.47	♂	1.19	4.64	4.44	0	0	0
1001G 2	4.34	♀	9.92	0	7.42	0	0	0
	.26	♀	3.80	0	5.54	0	0	0
	.26	♀	3.80	0	3.11	0	0	0
1001G 3	3.89	♂	6.67	0	6.56	1.17	0	0
	.12	♂	2.26	0	6.82	.71	0	0
	.17	♂	2.21	0	6.16	.51	0	0
1001G 4	4.15	♀	5.50	1.14	9.2	2.20	9.66	0
	.35	♀	3.34	2.26	4.7	.37	6.3	0
	.23	♀	2.20	2.26	4.18	.13	.36	0

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AVERAGED DATA

FEMALES

FILE NODNAME (CREATION DATE = 31/05/77)

PEARSON CORRELATION COEFFICIENTS

	V25AR3MX	V25HE3MX	V40AR3MX	V40HE3MX	CV3MX	SVC3MX	FEV3MX	FVC3MX	V25FR3MX	V40FR3MX
V25AR3MX	1.0000 (0) S= .001	.6995 (-38) S= .001	.7910 (-38) S= .001	.6551 (-38) S= .001	.0999 (-38) S= .001	.0022 (20) S= .275	.3369 (20) S= .071	.2251 (20) S= .170	.0129 (-38) S= .469	.1326 (-38) S= .214
V25HE3MX	.6995 (-38) S= .001	1.0000 (0) S= .001	.5352 (-38) S= .001	.7659 (-38) S= .001	.2606 (-38) S= .057	.0381 (20) S= .437	.3245 (20) S= .081	.1628 (20) S= .246	.3034 (-38) S= .032	.1245 (-38) S= .228
V40AR3MX	.7910 (-38) S= .001	.5352 (-38) S= .001	1.0000 (0) S= .001	.7112 (-38) S= .001	.0206 (-38) S= .001	.1286 (20) S= .451	.6276 (20) S= .295	.0118 (20) S= .480	.1047 (-38) S= .266	.2800 (-38) S= .044
V40HE3MX	.6551 (-38) S= .001	.7659 (-38) S= .001	.7112 (-38) S= .001	1.0000 (0) S= .001	.2902 (-38) S= .039	.2003 (20) S= .199	.2657 (20) S= .129	.2966 (20) S= .102	.2322 (-38) S= .080	.1361 (-38) S= .208
CV3MX	.0999 (-38) S= .275	.2606 (-38) S= .057	.0206 (-38) S= .451	.2902 (-38) S= .039	1.0000 (0) S= .001	.5891 (20) S= .003	.2649 (20) S= .130	.0405 (20) S= .433	.3915 (-38) S= .008	.5252 (-38) S= .001
SVC3MX	.0022 (20) S= .496	.0381 (20) S= .437	.1286 (20) S= .295	.2003 (20) S= .199	.5891 (20) S= .003	1.0000 (0) S= .001	.1730 (20) S= .233	.1617 (20) S= .248	.2021 (-20) S= .196	.3858 (-20) S= .046
FEV3MX	.3369 (20) S= .073	.3245 (20) S= .081	.6276 (20) S= .002	.2657 (20) S= .129	.2649 (20) S= .130	.1730 (20) S= .233	1.0000 (0) S= .001	.5105 (20) S= .011	.0069 (20) S= .488	.1803 (20) S= .223
FVC3MX	.2251 (20) S= .170	.1628 (20) S= .246	.0118 (20) S= .480	.2966 (20) S= .102	.0405 (20) S= .433	.1617 (20) S= .248	.5105 (0) S= .011	1.0000 (20) S= .001	.0397 (20) S= .434	.0764 (20) S= .374
V25FR3MX	.0129 (-38) S= .469	.3034 (-38) S= .032	.1047 (-38) S= .266	.2322 (-38) S= .080	.3915 (-38) S= .008	.2021 (-38) S= .196	.0069 (-38) S= .488	.0397 (-38) S= .434	.10000 (-38) S= .001	.6634 (-38) S= .001
V40FP3MX	.1326 (-38) S= .214	.1245 (-38) S= .228	.2800 (-38) S= .044	.1361 (-38) S= .208	.5252 (-38) S= .001	.3858 (-38) S= .046	.1803 (-38) S= .223	.0764 (-38) S= .374	.6634 (-38) S= .001	.0000 (-38) S= .001
CVSV3MX	.5746 (20) S= .004	.3818 (20) S= .048	.2849 (20) S= .112	.2973 (20) S= .102	.7730 (20) S= .001	.3497 (20) S= .065	.3683 (20) S= .055	.0479 (20) S= .420	.1388 (20) S= .280	.2199 (20) S= .176
FEFV3MX	.5438 (20) S= .007	.2160 (20) S= .180	.3725 (20) S= .053	.1956 (20) S= .204	.1125 (20) S= .150	.2438 (20) S= .279	.1393 (20) S= .004	.5717 (20) S= .251	.1596 (20) S= .409	.0517 (20) S= .409
IVFP3MX	.0481 (-38) S= .387	.2051 (-38) S= .108	.0962 (-38) S= .283	.1317 (-38) S= .215	.1883 (-38) S= .129	.3194 (-38) S= .085	.3356 (-38) S= .074	.4006 (-38) S= .040	.0402 (-38) S= .405	.2193 (-38) S= .093

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AVERAGED DATA
FEMALE (CREATION DATE = 31/05/77)
FILE: NORMAN

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	CVS3MX	FEV3MX	IVFP3MX
V25AR3MX	-.5746 (.20) S= .004	.5438 (.20) S= .007	.0481 (.38) S= .387
V25HE3MX	.3818 (.20) S= .048	.2160 (.20) S= .180	.2051 (.38) S= .108
V40AR3MX	.2849 (.20) S= .112	.3725 (.20) S= .053	.0962 (.38) S= .283
V40HE3MX	.2973 (.20) S= .102	.1956 (.20) S= .204	-.1317 (.38) S= .215
CV3MX	.7710 (.20) S= .001	.1125 (.20) S= .318	-.1983 (.38) S= .129
SVC3MX	.3497 (.20) S= .065	.2438 (.20) S= .150	-.3194 (.20) S= .085
FEV3MX	.3683 (.20) S= .055	.1193 (.20) S= .279	.3356 (.20) S= .074
FVC3MX	-.0479 (.20) S= .420	-.5717 (.20) S= .004	.4026 (.20) S= .040
V25FR3MX	-.1388 (.20) S= .280	-.1586 (.20) S= .251	-.0402 (.38) S= .405
V40FR3MX	.2199 (.20) S= .176	-.0547 (.20) S= .409	-.2193 (.38) S= .093
CV53MX	1.0000 (.00) S= .001	.3024 (.00) S= .098	-.0494 (.20) S= .418
FEV3MX	.3024 (.00) S= .098	1.0000 (.00) S= .001	-.0306 (.20) S= .449
IVFP3MX	-.0194 (.20) S= .118	-.0306 (.20) S= .449	1.0000 (.00) S= .001

AVERAGED DATA
FILE: NONAME (CREATION DATE: 31/05/77)

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PEARSON CORRELATION COEFFICIENTS

	V25AR3MX	V25WF3MX	V40AR3MX	V40WF3MX	CV3MX	SVC3MX	PEV3MX	PFC3MX	V25FR3MX	WASH3MX
V25AR3MX	1.0000	.4566	.4008	.4982	.0374	.0673	.0831	.0271	.0459	.0948
Ss	.001 (.001)	.581 (.001)	.591 (.001)	.591 (.001)	.301 (.301)	.301 (.301)	.301 (.301)	.301 (.301)	.591 (.591)	.591 (.591)
V25WF3MX	.4566	1.0000	.4549	.6875	.1232	.2112	.4166	.4155	.3023	.4465
Ss	.001 (.001)	.581 (.001)	.591 (.001)	.581 (.001)	.291 (.291)	.291 (.291)	.291 (.291)	.291 (.291)	.581 (.581)	.581 (.581)
V40AR3MX	.4008	.4549	1.0000	.5208	.1299	.0118	.0399	.0627	.0065	.0146
Ss	.001 (.001)	.581 (.001)	.591 (.001)	.591 (.001)	.136 (.136)	.136 (.136)	.1013 (.1013)	.1013 (.1013)	.591 (.591)	.591 (.591)
V40WF3MX	.4982	.6875	.5208	1.0000	.1494	.1849	.1943	.3646	.3469	.5844
Ss	.001 (.001)	.581 (.001)	.591 (.001)	.591 (.001)	.301 (.301)	.301 (.301)	.301 (.301)	.301 (.301)	.591 (.591)	.591 (.591)
CV3MX	.0374	.1232	.1299	.1494	1.0000	.6632	.2072	.1582	.1231	.2192
Ss	.591 (.591)	.581 (.581)	.591 (.591)	.591 (.591)	.301 (.301)	.301 (.301)	.301 (.301)	.301 (.301)	.591 (.591)	.591 (.591)
SVC3MX	.0673	.2112	.0118	.1849	.6632	1.0000	.2445	.1805	.3271	.3140
Ss	.301 (.301)	.291 (.291)	.301 (.301)	.301 (.301)	.301 (.301)	.301 (.301)	.301 (.301)	.301 (.301)	.301 (.301)	.301 (.301)
PEV3MX	.0831	.1466	.0399	.1943	.2072	.2445	1.0000	.2072	.1129	.0783
Ss	.301 (.301)	.291 (.291)	.301 (.301)	.301 (.301)	.301 (.301)	.301 (.301)	.301 (.301)	.301 (.301)	.301 (.301)	.301 (.301)
PFC3MX	.0271	.0155	.0627	.3646	.1582	.1805	.2072	1.0000	.2419	.0728
Ss	.443 (.443)	.0113 (.0113)	.371 (.371)	.024 (.024)	.202 (.202)	.170 (.170)	.136 (.136)	.001 (.001)	.099 (.099)	.160 (.160)
V25FR3MX	.0459	.3823	.0065	.1469	.2211	.2271	.1129	.2419	1.0000	.7162
Ss	.365 (.365)	.581 (.581)	.591 (.591)	.591 (.591)	.301 (.301)	.301 (.301)	.301 (.301)	.301 (.301)	.591 (.591)	.591 (.591)
WASH3MX	.0948	.4445	.0146	.5844	.2192	.3190	.0683	.0728	.7162	1.0000
Ss	.237 (.237)	.581 (.581)	.591 (.591)	.591 (.591)	.301 (.301)	.301 (.301)	.301 (.301)	.301 (.301)	.591 (.591)	.591 (.591)
CVS3MX	.0538	.1094	.0997	.0169	.0443	.3039	.3169	.0287	.2170	.1259
Ss	.301 (.301)	.291 (.291)	.301 (.301)	.301 (.301)	.301 (.301)	.301 (.301)	.301 (.301)	.301 (.301)	.301 (.301)	.301 (.301)
PEFV3MX	.2013	.2830	.2166	.2446	.0689	.0689	.1722	.8186	.3617	.1923
Ss	.143 (.143)	.286 (.286)	.300 (.300)	.465 (.465)	.301 (.301)	.301 (.301)	.301 (.301)	.301 (.301)	.591 (.591)	.591 (.591)
WFPP3MX	.1363	.2855	.1582	.2852	.2215	.2879	.0844	.1512	.1169	
Ss	.152 (.152)	.581 (.581)	.591 (.591)	.591 (.591)	.301 (.301)	.301 (.301)	.301 (.301)	.301 (.301)	.591 (.591)	.591 (.591)

AVERAGED DATA

MALES
FILE N0NAME (CREATION DATE = 31/05/77)

PEARSON CORRELATION COEFFICIENTS-----

	CVSV3MX	FFV3MX	IVFP3MX
V25AR3MX	.0538 (30) Sx .389	-.2013 (30) Sx .143	.1363 (59) Sx .152
V25HE3MX	-.1094 (29) Sx .286	.2230 (29) Sx .068	-.2855 (58) Sx .015
V40AR3MX	.0997 (30) Sx .300	-.2166 (30) Sx .125	.1582 (59) Sx .116
V40HE3MX	-.0169 (30) Sx .465	.2446 (30) Sx .096	-.2852 (59) Sx .014
CV3MX	.8043 (30) Sx .001	-.0689 (30) Sx .359	-.2215 (59) Sx .046
SVC3MX	.3039 (30) Sx .051	-.0689 (30) Sx .359	-.2879 (30) Sx .061
FFV3MX	.3369 (30) Sx .034	-.1722 (30) Sx .181	-.0844 (30) Sx .329
FVC3MX	.0787 (30) Sx .340	-.8386 (30) Sx .001	.1512 (30) Sx .213
V25FR3MX	.2370 (30) Sx .104	.3617 (30) Sx .025	-.0962 (59) Sx .234
V40FR3MX	.1259 (30) Sx .254	.1923 (30) Sx .154	-.1169 (59) Sx .189
CVSV3MX	1.0000 (0) Sx .001	.1313 (30) Sx .245	.0432 (30) Sx .410
FFFV3MX	.1313 (30) Sx .245	1.0000 (0) Sx .001	-.2491 (30) Sx .092
IVFP3MX	.0432 (30) Sx .410	-.2491 (30) Sx .092	1.0000 (0) Sx .001

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AVERAGED DATA
NON-SUMERS (CREATION DATE = 31/05/77)
FILE: NONAME

PAGE: 15

31/05/77
----- PEASON CORRELATION COEFFICIENTS -----

	CVS3MX	FEF3MX	TFP3MX
V25AR3MX	+.1260 (.28) Sz .261	+.2053 (.28) Sz .147	+.0013 (.55) Sz .496
V25MF3MX	+.070 (.26) Sz .026	+.2638 (.28) Sz .087	+.0234 (.55) Sz .185
V40AR3MX	+.2510 (.28) Sz .099	+.2106 (.28) Sz .141	+.0988 (.55) Sz .234
V40MF3MX	+.0048 (.28) Sz .490	+.2800 (.28) Sz .068	+.2347 (.55) Sz .042
CV3MX	+.8658 (.28) Sz .001	+.0365 (.28) Sz .427	+.1599 (.55) Sz .172
SV3MX	+.2964 (.28) Sz .076	+.1375 (.28) Sz .243	+.2120 (.55) Sz .117
FFV3MX	+.1815 (.28) Sz .023	+.1659 (.28) Sz .199	+.0186 (.28) Sz .463
FVC3MX	+.0022 (.28) Sz .496	+.8576 (.28) Sz .001	+.2584 (.55) Sz .092
V25FR3MX	+.0557 (.28) Sz .389	+.5372 (.28) Sz .002	+.0019 (.55) Sz .383
V40FR3MX	+.0347 (.28) Sz .471	+.2180 (.28) Sz .133	+.0163 (.55) Sz .453
CV5V3MX	1.00000 (.00) Sz .001	.1227 (.28) Sz .267	.0312 (.28) Sz .431
FFF3MX	+.1227 (.28) Sz .267	1.00000 (.00) Sz .001	-.3168 (.28) Sz .050
TFP3MX	+.0142 (.28) Sz .431	-.3168 (.28) Sz .050	1.00000 (.00) Sz .001

AVERAGED DATA		PEARSON CORRELATION COEFFICIENTS											
SINKERS	FILE	NAME	(CREATION DATE = 31/05/77)	V25AR3MX	V25HE3MX	V40AR3MX	V40HE3MX	CV3MX	SVC3MX	FEP3MX	FVC3MX	V25FR3MX	V40FR3MX
V25AR3MX	{	1,0000		.6855	.8170	.5358	.0373	.0194	.1560	.0642	.0337	.1001	
		{ .001	{ .41	{ .42	{ .42	{ .42	{ .42	{ .22	{ .22	{ .42	{ .42	{ .42	
	S#	.001	S#	.001	S#	.001	S#	.007	S#	.044	S#	.044	S#
V25HE3MX	{	.6855	1,0000	.5519	.8320	.2389	.1736	.3111	.1687	.2984	.3012		
		{ .41	{ .001	{ .41	{ .41	{ .41	{ .41	{ .21	{ .21	{ .41	{ .41	{ .41	
	S#	.001	S#	.001	S#	.001	S#	.066	S#	.226	S#	.226	S#
V40AR3MX	{	.8170	.5519	1,0000	.4754	.0099	.1733	.1207	.0152	.0257	.0245		
		{ .42	{ .41	{ .001	{ .42	{ .42	{ .42	{ .22	{ .22	{ .42	{ .42	{ .42	
	S#	.001	S#	.001	S#	.001	S#	.475	S#	.220	S#	.220	S#
V40HE3MX	{	.5358	.8328	.4754	1,0000	.2819	.1500	.2360	.1939	.3795	.4010		
		{ .42	{ .41	{ .42	{ .001	{ .42	{ .42	{ .22	{ .22	{ .42	{ .42	{ .42	
	S#	.001	S#	.001	S#	.001	S#	.035	S#	.253	S#	.253	S#
CV3MX	{	.0373	.2389	.0099	.2819	1,0000	.7424	.2979	.3534	.5117	.5140		
		{ .42	{ .41	{ .42	{ .42	{ .42	{ .42	{ .22	{ .22	{ .42	{ .42	{ .42	
	S#	.007	S#	.006	S#	.035	S#	.001	S#	.089	S#	.089	S#
SVC3MX	{	.0194	.1736	.1733	.1500	.7424	1,0000	.2976	.2946	.3162	.3177		
		{ .22	{ .21	{ .22	{ .22	{ .22	{ .22	{ .00	{ .00	{ .22	{ .22	{ .22	
	S#	.466	S#	.226	S#	.220	S#	.053	S#	.001	S#	.001	S#
FEP3MX	{	.1560	.3311	.1202	.7360	.2979	.2976	1,0000	.6720	.0427	.1565		
		{ .22	{ .21	{ .22	{ .22	{ .22	{ .22	{ .00	{ .00	{ .22	{ .22	{ .22	
	S#	.244	S#	.071	S#	.297	S#	.145	S#	.089	S#	.089	S#
FVC3MX	{	.0687	.0952	.1939	.3534	.2946	.2946	.6720	1,0000	.1398	.3114		
		{ .22	{ .21	{ .22	{ .22	{ .22	{ .22	{ .00	{ .00	.22	.22	.22	
	S#	.388	S#	.384	S#	.337	S#	.194	S#	.062	S#	.062	S#
V25FR3MX	{	.0337	.2984	.0257	.3195	.5117	.3162	.0427	.1398	1,0000	.427		
		{ .42	{ .41	{ .42	{ .42	{ .42	{ .42	{ .22	{ .22	{ .00	{ .42	{ .42	
	S#	.416	S#	.079	S#	.436	S#	.007	S#	.076	S#	.076	S#
V40FR3MX	{	.1001	.3012	.0261	.4010	.510	.3895	.1565	.3114	.5980	.1,0000		
		{ .42	{ .41	{ .42	{ .42	{ .42	{ .42	{ .22	{ .22	{ .42	{ .42	{ .42	
	S#	.264	S#	.028	S#	.059	S#	.004	S#	.031	S#	.031	S#
CV5V3MX	{	.2620	.2088	.0104	.1476	.7891	.4127	.2694	.2923	.1770	.3053		
		{ .22	{ .21	{ .22	{ .22	{ .22	{ .22	{ .22	{ .22	{ .22	{ .22	{ .22	
	S#	.119	S#	.182	S#	.482	S#	.256	S#	.028	S#	.028	S#
FEP3MX	{	.5256	.1129	.3883	.2179	.1635	.1441	.2792	.2075	.0289	.0092		
		{ .22	{ .21	{ .22	{ .22	{ .22	{ .22	{ .22	{ .22	{ .22	{ .22	{ .22	
	S#	.006	S#	.313	S#	.037	S#	.165	S#	.234	S#	.261	S#
IVFP3MX	{	.2562	.0045	.4050	.2067	.3505	.0315	.1866	.1149	.2881			
		{ .42	{ .41	{ .42	{ .42	{ .42	{ .42	{ .22	{ .22	{ .42	{ .42	{ .42	
	S#	.051	S#	.489	S#	.004	S#	.130	S#	.094	S#	.045	S#

AVERAGED DATA
SMOKERS
FILE: NONAME (CREATION DATE = 31/05/77)

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----- PEARSON CORRELATION COEFFICIENTS -----

	CVS3MX	FFV3MX	IVFP3MX
V25AR3MX	.2620 (.22) S= .119	.5256 (.22) S= .006	.2562 (.42) S= .051
V25HE3MX	.2088 (.21) S= .182	.1129 (.21) S= .313	.0045 (.41) S= .489
V40AR3MX	-.0104 (.22) S= .482	.3883 (.22) S= .037	.4050 (.42) S= .004
V40HE3MX	.1476 (.22) S= .256	-.2179 (.22) S= .165	-.1779 (.42) S= .130
CV3MX	.7891 (.22) S= .001	.1635 (.22) S= .234	-.2067 (.42) S= .094
SVC3MX	-.4127 (.22) S= .028	-.1441 (.22) S= .261	-.3505 (.22) S= .055
FFV3MX	.2694 (.22) S= .113	.2792 (.22) S= .104	.0315 (.22) S= .445
IVFC3MX	.2923 (.22) S= .093	-.2075 (.22) S= .177	.1866 (.22) S= .203
V25FR3MX	.1770 (.22) S= .215	.0289 (.22) S= .449	-.1149 (.42) S= .234
V40FR3MX	.3053 (.22) S= .084	-.0092 (.22) S= .484	-.2881 (.42) S= .032
CVS3MX	1.0000 (.01) S= .001	.3060 (.22) S= .083	.0296 (.22) S= .448
FFFV3MX	.3060 (.22) S= .083	1.0000 (.01) S= .001	.0595 (.22) S= .396
IVFP3MX	.0296 (.22) S= .446	.0595 (.22) S= .396	1.0000 (.01) S= .001

AVERAGED DATA
DRUG ONE FILE: N0NAME (CREATION DATE = 01/06/77)

GROUP COUNTS		TOTAL				TOTAL	
NUMBER	NPANS	GROUP 1	GROUP 2	GROUP 3	GROUP 4	GROUP 1	GROUP 4
V25AP1**N	2,32600	1.62667	2.25714	2.38500	2.20316		
V25H1**N	2,85200	2.17000	2.56857	2.81500	2.61632		
VADAF1**N	3,52600	2.45000	4.31000	4.30000	3.90154		
VADHF1**N	4,22700	3.47333	5.41286	5.51750	4.84664		
CV1**N	90400	94000	1.85286	1.30750	1.34471		
V25PP1**N	27000	14667	1.36714	2.27500	3.0195		
V40F1**N	23800	2R383	3.80000	2.00000	3.0632		
IVPP1**N	314000	40.11667	13.96141	16.17250	23.35158		
V25AP1**X	~29000	~16667	~5.0143	~5.51750	~4.0174		
V25HF3**X	~44800	~31667	~3.4290	~6.60000	~6.21648		
V40AP3**X	~21400	~21333	~8.1143	~9.9750	~6.1842		
V40HF3**X	~03600	~1.12000	~7.7571	~3.8500	~5.31721		
CV1**X	~17400	~1.37667	~7.2000	~3.3750	~6.03568		
V25FR3**X	~14200	~14333	~8.10000	~2.7750	~25.189		
V40FR3**X	~14000	~03000	~6.2429	~27000	~11895		
IVPP3**X	~7,37000	~12.41000	~15.10143	~14750	~1.69763		

STANDARD DEVIATIONS

GROUP 1		TOTAL			
GROUP 1	GROUP 2	GROUP 3	GROUP 4	GROUP 1	GROUP 4
V25AP1**N	39416	83243	1.35266	4.1605	90411
V25H1**N	61267	1.44679	1.01769	6.9431	90401
VADAF1**N	69640	57420	2.06778	4.91851	4.45504
VADHF1**N	61913	1.05136	1.95892	1.02812	1.49713
CV1**N	29716	35511	4.2680	4.4131	5.55472
V25PP1**N	10949	18175	2.26487	1.14442	1.14477
V40PP1**N	09612	13317	1.17436	1.13745	
IVPP1**N	30.66124	41.63206	7.95593	3.47880	21.26222
V25AP3**N	56720	58184	4.65644	7.13330	6.0471
V25HF3**X	91087	87157	3.4287	7.0356	6.3555
V40AP3**X	74127	57292	9.1140	1.26907	9.14275
V40HF3**X	1.09742	11765	2.50057	1.46439	1.70166
CV1**X	55649	34106	1.7758	4.4607	4.0654
V25FP3**X	31124	52291	1.58023	1.3788	1.06245
V40FP3**X	21304	68700	1.68972	2.02071	1.09434
IVPP3**X	38.09492	31.50287	20.21311	9.77180	26.68191

APPENDIX DATA
COPIC ONE

WITHIN GROUPS COVARIANCE MATRIX

	V25AP1MN	V25AR1MN	V25HE1MN	V40AP1MN	V40HE1MN	CY1MN	V25FP1MN	V40FP1MN	IVFP1MN	V25AR3MX
V25AP1MN	.4907	.91335								
V25AR1MN	.81357	1.07472	2.05746							
V40AP1MN	1.22677	1.12676	1.92061	1.99612						
V40AR1MN	1.27474	1.15407	1.09157	1.11726	1.15777					
CY1MN	-1.17224	-1.10047	-0.9864	-1.15976	-1.12766	1.01418	1.0326	1.02022	1.01842	
V25FP1MN	-1.10047	0.6957	0.3697	1.4717	1.0872	0.0306	1.18722	1.45775	1.50946358	
V40FP1MN	0.6957	6.72162	10.72103	3.8364	6.20123	2.19905	1.18722	1.35611	1.41390	
IVFP1MN	6.72162	-2.16096	-2.27830	-1.44155	-1.2004	1.12041	-0.0216	-0.00281	-0.00281	
V25AP3MX	-2.16096	-2.16832	-3.60334	-2.21232	-3.36167	1.11507	-0.00053	-0.126272	-0.126272	
V25AR3MX	-2.16832	-3.67726	-3.4757	-2.76163	-2.73188	0.08664	0.0677	5.66157	5.66157	
V40AP3MX	-3.67726	-1.04472	-1.90185	-1.44175	-1.545482	1.12708	1.2547	1.0104	1.01745	
V40AR3MX	-1.04472	-1.1892	-1.12334	-0.21009	-0.21014	0.00381	0.01579	0.00567	0.01359	
CY3MN	-1.1892	-0.57093	-0.48975	-0.6184	-0.76568	-0.06184	-0.02910	-0.02260	-0.05374	0.05374
V25FP3MX	-0.57093	-0.60397	-0.44667	-0.51676	-0.67114	0.06915	0.01457	-0.39593	-0.39593	
V40FP3MX	-0.60397	-2.41863	-7.54255	5.51630	0.93677	1.18896	-1.78242	-1.17499	-4.48573	-4.48573
IVFP3MX	-2.41863	V40AR3MX	V40HE3MX	V40HE3MX	CY3MN	V25FP3MX	V40FP3MX	IVFP3MX		
V25HE3MX	-1.48861	-3.0614	-0.85489							
V40HE3MX	-3.0614	-1.10644	3.26663							
CY3MN	-0.85489	5.6241	-0.07193	-0.11518	-1.9315					
V25FP3MX	5.6241	-0.07193	-1.15157	-0.44712	-1.15960	1.08253				
V40FP3MX	-0.07193	-1.15157	-0.06468	1.20670	0.40097	1.93676	1.25276			
IVFP3MX	-1.15157	-0.06468	-12.49920	-3.23352	-1.95475	10.43996	11.30334	102.50068		

VARIABLES IN THE ANALYSIS •

VARIABLE	ENTRY CRITERION	F TO REMOVE
V25AP1MN	+1.41327	.56125
V25SF1MN	+2.27031	.53992
V40AP1MN	-1.93047	2.06502
V40HF1MN	+3.47254	1.07415
CV1MN	-4.10284	1.40426
V25SF1MN	-1.52833	.91617
V40AP1MN	+2.91336	4.80114
IVFP1MN	+1.42128	.88657

• VARIABLES NOT IN THE ANALYSIS •

NUMBER OF VARIOIDS	EIGENVALUE	CANONICAL CORRELATION	PERCENT OF TRACE	WILKS LAMBDA	CHI-SQUARE	D.F.	SIGNIFICANCE
0	9.47394	.95107	90.5	.04329	40.81771	24	.017
1	.66696	.61254	6.4	.45344	10.28164	14	.741
2	.32299	.49410	3.1	.475587	3.63859	6	.725

3 FUNCTIONS WILL BE USED IN REMAINING ANALYSES

STANDARDIZED DISCRIMINANT FUNCTION COEFFICIENTS

1	2	3
V25AP1MN	2.45091	.07861
V25SF1MN	-2.10352	3.34118
V40AP1MN	+5.65337	2.54588
V40HF1MN	1.79741	-5.33511
CV1MN	-1.15307	1.18464
V25SF1MN	1.60888	-.30116
V40AP1MN	+3.40499	.52017
IVFP1MN	-4.47900	-.99230

UNSTANDARDIZED DISCRIMINANT FUNCTION COEFFICIENTS

1	2	3
V25AP1MN	2.71084	.869479E-01
V25SF1MN	-.114258	3.68781
V40AP1MN	+3.43106	1.77298
V40HF1MN	1.20460	+3.57550
CV1MN	-2.0791	2.13171
V25SF1MN	8.54462	-1.60216
V40AP1MN	+2.76009	3.77351
IVFP1MN	-.192158E-01	-.426571E-01
CONSTANT	12.00992	-2.03157

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AVERAGED DATA
DRUG ONE

CENTROIDS OF GROUPS IN REDUCED SPACE

GROUP 1	3.14468	.88023	.04031
GROUP 2	2.61086	-1.19218	-.66278
GROUP 3	-3.20490	.23123	-.24710
GROUP 4	-.28043	-.61080	.87912

PREDICTION RESULTS =

ACTUAL GROUP NAME	GROUP CODE	N OF CASES	PREDICTED GROUP MEMBERSHIP			
			GROUP 1	GROUP 2	GROUP 3	GROUP 4
GROUP 1	2	6	5. 26.3 PCT	1. 5.3 PCT	0 0 PCT	0 0 PCT
GROUP 2	3	4	1. 5.3 PCT	2. 10.5 PCT	0 0 PCT	1. 5.3 PCT
GROUP 3	4	8	0 0 PCT	0 0 PCT	7. 36.8 PCT	1. 5.3 PCT
GROUP 4	5	7	0 0 PCT	2. 10.5 PCT	1. 5.3 PCT	4. 21.1 PCT

94.7 PERCENT OF KNOWN CASES CORRECTLY CLASSIFIED

CHI-SQUARE = 49.281 SIGNIFICANCE = .000

APPENDIX DATA
PRINTING D.F.

VARIABLES IN THE ANALYSIS		
VARIABLE	ENTRY CRITERION	F TO REMOVE
V75AR3MX	-2.42227	*.69872
V75HE3MX	-2.58444	1.12678
V40P3MX	-6.36666	1.48815
V40R3MX	-3.93536	3.68199
V40E3MX	-.80295	.65367
CV3MX	-2.45794	.59552
V25F3MX	-2.94830	1.19079
V40P3MX	-2.78914	.19510
V40F3MX		

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VARIABLES NOT IN THE ANALYSIS

VARIABLE	ENTER	TOLERANCE	F TO ENTER	CRITERION
V75AR3MX				
V75HE3MX				
V40P3MX				
V40R3MX				
V40E3MX				
CV3MX				
V25F3MX				
V40P3MX				
V40F3MX				

FUNCTIONS WILL BE USED IN REMAINING ANALYSES

STANDARDIZED DISCRIMINANT FUNCTION COEFFICIENTS

	1	2	3
V75AR3MX	1.34584	1.40164	2.36103
V75HF3MX	*1.50263	-.91131	*.70770
V40AR3MX	*2.41162	.07110	*2.31101
V40RF3MX	3.12797	-.51644	*.21755
CV3MX	*.59905	-.63306	-.04080
V25F3MX	1.77148	.16048	.66967
V40FR3MX	*3.81883	-.13760	*.23613
V40F3MX	*.66700	.83257	*.57301

UNSTANDARDIZED DISCRIMINANT FUNCTION COEFFICIENTS

	1	2	3
V75AR3MX	2.20169	2.31781	3.69410
V75HT3MX	-.1.36129	-.1.41189	-.1.11195
V40AR3MX	*2.61282	*.76092E-02	*2.5432
V40RF3MX	1.93819	*.303491	*.124907
CV3MX	1.23124	-.1.30114	-.936605E-01
V25F3MX	1.66873	*.151042	*.630306
V40FR3MX	-.1.48861	-.398879	-.215770
V40F3MX	*.124439E-01	*.312038E-01	*.216758E-01
CONSTANT	*1.27966	*.213759	*.580361

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AVERAGED DATA
DRUG ONE

CENTROIDS OF GROUPS IN REDUCED SPACE

GROUP 1	-.57104	-.76879	-.50152
GROUP 2	-.296097	.71140	.27136
GROUP 3	1.25710	.79025	-.07259
GROUP 4	.73459	-.95700	.55042

PREDICTION RESULTS -

ACTUAL GROUP NAME	GROUP CODE	N OF CASES	PREDICTED GROUP MEMBERSHIP			
			GROUP 1	GROUP 2	GROUP 3	GROUP 4
GROUP 1	2	6	3. 15.8 PCT	0 0 PCT	2. 10.5 PCT	1. 5.3 PCT
GROUP 2	3	4	0 0 PCT	3. 15.8 PCT	0 0 PCT	1. 5.3 PCT
GROUP 3	4	8	1. 5.3 PCT	0 0 PCT	7. 36.8 PCT	0 0 PCT
GROUP 4	5	7	1. 5.3 PCT	1. 5.3 PCT	1. 5.3 PCT	4. 21.1 PCT

89.5 PERCENT OF KNOWN CASES CORRECTLY CLASSIFIED

CHI-SQUARE = 42.123 SIGNIFICANCE = .000

01/06/77

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AVERAGED DATA
 DRING TWO
 FILE: ROMEA (CREATION DATE = 01/06/77.)

GROUP COUNTS

	NUMBER	GROUP 1	GROUP 2	GROUP 3	GROUP 4	TOTAL
	4.	2.	5.	4.	15.	

MEANS

	GROUP 1	GROUP 2	GROUP 3	GROUP 4	TOTAL
V25AR1MN	2.355750	2.010000	1.720000	1.70500	1.97800
V25MF1MN	3.07750	2.76000	2.11600	2.05000	2.44067
V25AR1MN	3.56500	3.27000	3.22600	3.56250	3.41200
V25MF1MN	4.14600	4.03500	4.69700	5.11500	4.74067
CV1MN	65750	.77000	1.59000	1.40500	1.18533
V25FR1MN	26250	38500	29600	27750	279667
V25FR1MN	40250	31500	44000	44500	44000
IVPP1MN	15.16500	67.44500	18.13000	19.69750	24.51933
V25AP3MX	61000	51000	25000	21000	311933
V25MF3MX	69750	68500	69400	72100	73660
V25MF3MX	71000	67000	66600	52500	555667
V25MF3MX	1.20750	1.20750	1.55000	1.17500	1.92267
CY1MN	16500	50000	40200	28250	306533
V25FR3MX	39000	52000	66000	14250	22113
V25FR3MX	40400	40400	424000	424000	42933
IVPP3MX	20.10000	9.48000	15.61800	16.00000	13.57533

STANDARD DEVIATIONS

	GROUP 1	GROUP 2	GROUP 3	GROUP 4	TOTAL
V25AR1MN	.68066	.50912	.75802	.60379	.70799
V25MF1MN	.48210	.69246	.67059	.63451	.73311
V25AR1MN	.60534	.80610	.1.27500	.1.01501	.1.01501
V25AR1MN	.44714	.86974	.1.84735	.1.79896	.1.24137
CY1MN	1.70695	2.26217	.23156	.49346	.50623
V25FR1MN	.14569	.04121	.0.04949	.1.0342	.1.11976
V25FR1MN	.04573	.03536	.0.04949	.1.0342	.1.11976
IVPP1MN	5.15523	44.11639	2.89776	3.84426	21.35577
V25AR1MN	.45395	.08890	.63106	.71680	.57485
V25FR3MX	1.00705	.50705	.61897	.1.28243	.90131
V25AP3MX	.20116	.60H11	1.05099	.88776	.79123
V25MF3MX	.45666	.32521	.99440	1.41503	.92576
CY1MN	.31670	.32527	.41017	.40665	.39815
V25FR3MX	.96336	.11314	.28991	.2R441	.93669
V25FR3MX	.26211	.31113	.52675	.30854	.39913
IVPP3MX	39.54302	13.10976	33.811979	17.01320	28.76585

AVERAGED DATA
DURING TWO

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VARIABLES IN THE ANALYSIS

VARIABLE	ENTRY CRITERION	F TO REMOVE
V75AP1MN	*.9235	1.78804
V75MF1MN	*1.8194	2.48912
V40R1MN	-.86038	2.49515
V40E1MN	-1.37238	2.92223
CV1MN	-.3.46312	1.08706
V25P1MN	-.05943	.56187
V40F1MN	-.99028	.79018
IVP1PM	-.216442	.795262

VARIABLES NOT IN THE ANALYSIS

VARIABLE	TOLERANCE	F TO ENTER	ENTRY CRITERION
V75AP1MN	.00008	47.52034	.74
V75MF1MN	.00508	20.07662	14
V40R1MN	.01741	.548046	.484
V40E1MN	.54393	6	

3 FUNCTIONS WILL BE USED IN REMAINING ANALYSES

STANDARDIZED DISCRIMINANT FUNCTION COEFFICIENTS

	1	2	3
V75AP1MN	*5.62607	*8.09688	-1.09621
V75MF1MN	7.16674	3.92580	-1.01119
V40R1MN	7.12664	6.25221	.12557
V40E1MN	*8.16657	*2.77226	2.16318
CV1MN	*3.3579	*.30344	-2.56276
V25P1MN	*1.0113	*1.0164	.38814
V40F1MN	.05113	.54980	1.52349
IVP1PM	.36683	1.31617	.69304
IVFP1PM	1.51646		

UNSTANDARDIZED DISCRIMINANT FUNCTION COEFFICIENTS

	1	2	3
V75AP1MN	*7.94657	*11.4365	*1.54835
V75MF1MN	9.26287	5.07402	*1.30721
V40R1MN	7.79552	6.16024	.12712
V40E1MN	*6.80939	*2.23020	1.67900
CV1MN	*6.63309	*5.94409	-5.0144
V25P1MN	4.94370	*16.4734	*3.75412
V40F1MN	3.06297	12.9173	12.7210
IVP1PM	*7.8189E-01	*6.03109E-01	*32.9527E-01
CONSTANT	*5.29751	*1.81471	*1.44995

AVERAGED DATA
DRUG TWO

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CENTROIDS OF GROUPS IN REDUCED SPACE

GROUP 1	1.09804	-2.68064	.39771
GROUP 2	8.95586	1.77554	-.10505
GROUP 3	-2.13921	.20507	-1.01528
GROUP 4	-2.90195	1.53653	.92392

PREDICTION RESULTS -

ACTUAL GROUP NAME	GROUP CODE	N OF CASES	PREDICTED GROUP MEMBERSHIP			
			GROUP 1	GROUP 2	GROUP 3	GROUP 4
GROUP 1	2	6	4. 26.7 PCT	1. 6.7 PCT	1. 6.7 PCT	0 0 PCT
GROUP 2	3	2	0 0 PCT	2. 13.3 PCT	0 0 PCT	0 0 PCT
GROUP 3	4	7	0 0 PCT	1. 6.7 PCT	6. 40.0 PCT	0 0 PCT
GROUP 4	5	7	0 0 PCT	1. 6.7 PCT	3. 20.0 PCT	3. 20.0 PCT

100.0 PERCENT OF KNOWN CASES CORRECTLY CLASSIFIED

CHI-SQUARE = 45.000 SIGNIFICANCE = .000

AVERAGED DATA
DURING TWO

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***** VARIABLES IN THE ANALYSIS *****

VARIABLE	ENTRY CRITERION	F TO REMOVE	VARIANCE	TOLERANCE	F TO ENTER	ENTRY CRITERION
V25AR3MX	+1.25449	1.41060				
V25HF3MX	-3.29544	.54518				
V40AP3MX	+2.13008	.47275				
V40HE3MX	+2.93096	.22637				
CY34X	+2.63028	.98975				
V25FR3MX	+3.82379	1.56802				
V40FR3MX	+5.00000	1.74187				
IVFP3MX	+2.23944	1.05309				

3 FUNCTIONS WILL BE USED IN REMAINING ANALYSES

STANDARDIZED DISCRIMINANT FUNCTION COEFFICIENTS

	1	2	3
V25AR3MX	+1.60558	+2.36795	+36395
V25HF3MX	+2.26113	2.00243	1.36458
V40AP3MX	+35889	1.48806	+44055
V40HE3MX	+86766	+11078	+60094
CY34X	+93108	+26514	+12310
V25FR3MX	+1.48518	+11058	+99031
V40FR3MX	1.77138	+73021	+40666
IVFP3MX	1.20353	+30429	+22565

UNSTANDARDIZED DISCRIMINANT FUNCTION COEFFICIENTS

	1	2	3
V25AR3MX	+2.79306	+4.11927	+631127
V25HF3MX	+2.26191	2.22169	1.51400
V40AP3MX	+448527	1.88669	+566794
V40HE3MX	+937235	+118662	+755442
CY34X	+2.38449	+665915	+39183
V25FR3MX	+1.58557	+118051	+1.05724
V40FR3MX	4.48809	+1.92948	+1.01887
IVFP3MX	+418389E-01	+10578E-01	+74446E-02
CONSTANT	+195732	+306833	+557122

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AVERAGED DATA
DRUG TWO

CENTROIDS OF GROUPS IN REDUCED SPACE

GROUP 1	2.51644	.10105	.63916
GROUP 2	1.04719	-1.65411	-1.14804
GROUP 3	-.56080	1.03922	-.52171
GROUP 4	-2.33903	-.57301	.58700

PREDICTION RESULTS -

ACTUAL GROUP NAME	GROUP CODE	N OF CASES	PREDICTED GROUP MEMBERSHIP			
			GROUP 1	GROUP 2	GROUP 3	GROUP 4
GROUP 1	2	6	4. 26.7 PCT	1. 6.7 PCT	0 0 PCT	1. 6.7 PCT
GROUP 2	3	2	0 0 PCT	2. 13.3 PCT	0 0 PCT	0 0 PCT
GROUP 3	4	7	0 0 PCT	0 0 PCT	5. 33.3 PCT	2. 13.3 PCT
GROUP 4	5	7	2. 11.1 PCT	0 0 PCT	0 0 PCT	5. 33.3 PCT

100.0 PERCENT OF KNOWN CASES CORRECTLY CLASSIFIED

CHI-SQUARE = 53.356 SIGNIFICANCE = .000

AVERAGED DATA
DRUG THREE
FILE: N0NAME (CREATION DATE: 31/05/77)

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GROUP COUNTS

NUMBER	GROUP 1	GROUP 2	GROUP 3	GROUP 4	TOTAL
5.	4.	7.	4.	20.	

MEANS

	GROUP 1	GROUP 2	GROUP 3	GROUP 4	TOTAL
V25AB1NN	2.17000	1.65750	1.83286	1.82000	1.87950
V25HE1NN	2.58660	2.06000	2.10429	2.10500	2.21100
V46AB1NN	3.47800	2.90500	3.32786	3.78750	3.71100
V40HE1NN	4.48800	3.82000	4.52143	4.80000	4.44700
CV1NN	*.98600	1.02500	1.90571	1.56750	1.31350
SVC1NN	7.70000	7.06750	10.11246	9.16250	8.85050
FFV1NN	6.84400	6.42250	9.00571	8.11500	7.91050
FVC1NN	R.31200	7.91250	12.76571	11.24500	10.38250
V25FP1NN	*.27000	*.38750	*.23429	*.29000	*.28500
V40FP1NN	*.28600	*.34250	*.42571	*.37750	*.38450
CVSV1NN	12.80000	14.37250	19.15286	16.19750	16.01750
FFFP1NN	82.62200	81.91750	71.58571	79.10750	77.91550
TFP1NN	18.67400	17.36750	22.36714	16.34000	19.31150
V25AB3NN	*.30000	*.45250	*.01857	*.69500	*.11100
V25HE3NN	*.63400	*.43250	*.52714	*.07000	*.44350
V40AB3NN	*.67200	*.87500	*.36087	*.99000	*.44850
V40HE3NN	*.95000	*.67000	1.11857	*.17500	*.79750
CF3NN	*.16200	*.04250	*.35143	*.28250	*.12650
SVC3NN	*.41400	*.41250	*.03571	*.24750	*.01600
FFV3NN	*.30000	*.53000	*.14143	*.18500	*.09550
FVC3NN	*.17600	*.12500	*.1272616	*.12000	*.08700
V25FP3NN	*.07600	*.07250	*.17714	*.44750	*.15600
V40FP3NN	*.01200	*.05500	*.00000	*.13500	*.00350
CVSV3NN	1.85800	1.34250	*.64571	*.68500	*.67500
FFFV3NN	1.39000	*.71750	31.44429	4.26750	14.16300
TFP3NN	13.40200	40.37000	-8.53857	26.24250	13.58450

STANDARD DEVIATIONS

	GROUP 1	GROUP 2	GROUP 3	GROUP 4	TOTAL
V25AB1NN	*.65718	*.72491	*.95777	*.43871	*.72653
V25HE1NN	*.02080	*.127260	*.74405	*.90164	*.89390
V46AB1NN	1.02356	*.64003	1.25114	*.92408	*.96298
V40HE1NN	1.49001	1.19861	1.74596	.90377	1.18450
CV1NN	*.16329	*.36546	*.31416	*.20419	*.50903
SVC1NN	2.87716	*.94013	1.78411	1.62046	2.76647
FFV1NN	1.70485	*.52506	1.37483	*.90091	1.66942
FVC1NN	2.22500	1.00500	2.21450	2.00557	2.03292
V25FP1NN	*.12219	*.20119	*.12660	*.18813	*.16343
FFFP1NN	*.08114	*.13099	*.12687	*.14198	*.12479
CVSV1NN	*.97576	3.28479	3.31161	3.44166	3.76382

WITHIN GROUPS COVARIANCE MATRIX

	V25AR1HN	V25HE1HN	V40AH1HN	V40HF1HN	V40V1HN	V40V2HN	V40FR1HN	V40FR3HN
V25HE1HN	.54629	.8786	.9905	2.12787	.10287	3.93844	1.67124	4.02023
V40AP1HN	.64886	.76400	.96506	.16506	.14375	.43728	.217994	.191760
CY1HN	1.00264	1.22743	1.36520	1.36520	1.3713	.53713	1.91524	.02789
SUC14K	.08613	.09616	.09616	.09616	.09616	.09616	.09616	.09616
FIC1HN	.57761	.69428	.53713	.1.4375	.1.4375	.1.4375	.1.4375	.1.4375
FIC2HN	.42274	.56943	.56943	.56943	.56943	.56943	.56943	.56943
V25FR1HN	-.24433	-.03778	-.26221	-.07772	-.07772	.21023	.21023	.21023
V25FR3HN	.01110	-.05773	.03616	-.07578	-.07578	.01848	.01848	.01848
V40FR1HN	-.02244	-.04713	-.03414	-.05533	-.05533	.01023	.01023	.01023
CYSV1HN	-.18706	-.45196	-.07182	-.79068	-.79068	.43231	.43231	.43231
FIFV1HN	5.11299	5.52748	5.4531	9.69071	9.69071	-.38393	-.38393	-.38393
IVP1HN	-.21761	-.21418	-.61853	-.96521	-.96521	4.15604	4.15604	4.15604
V25AR3HN	-.25994	-.10655	-.22699	-.46406	-.46406	-.0367	-.0367	-.0367
V25HF1HN	-.02271	-.16133	-.10958	-.05996	-.05996	.24300	.24300	.24300
V25AR3HN	.04680	-.01948	-.06160	.04231	.04231	.01890	.01890	.01890
V40HE3HN	-.07384	-.27954	-.08702	-.01708	-.01708	.00281	.00281	.00281
CY3HN	-.02133	-.02700	-.07631	-.08488	-.08488	-.015171	-.015171	-.015171
SUC34K	-.11460	-.26981	-.18052	-.39119	-.39119	-.07092	-.07092	-.07092
FIF3HN	-.03128	-.09229	-.01905	-.00631	-.00631	.015446	.015446	.015446
FIC3HN	-.07053	-.25181	-.81210	-.70135	-.70135	-.17978	-.17978	-.17978
V25FP3HN	.0R18A	-.03084	-.17004	-.14175	-.14175	-.03218	-.03218	-.03218
V40FP3HN	.02752	-.03527	-.02429	-.02255	-.02255	-.02117	-.02117	-.02117
CYSV3HN	-.02706	-.21957	-.87967	-.91441	-.91441	-.542786	-.542786	-.542786
FIFV3HN	7.28410	9.12049	29.57158	24.82685	24.82685	-.415060	-.415060	-.415060
IVPF3HN	6.95176	11.36805	6.97283	11.17131	11.17131	3.07648	3.07648	3.07648
V40FR1HN	CY5V1HN	FEV1HN	I4VP1HN	V25AR3HN	V40FR3HN	CY3HN	V25HE3HN	V40HE3HN
V40FR1HN	.01577	-.02201	8.60204	8.60204	8.60204	.04718	.04718	.04718
CYSV1HN	-.02263	4.69711	72.3844	72.3844	72.3844	-.02023	-.02023	-.02023
FFV1HN	-.30601	5.61893	13.74636	13.74636	13.74636	-.21792	-.21792	-.21792
IVP1HN	-.00786	.38004	-.67828	-.67828	-.67828	-.09827	-.09827	-.09827
V25AP1HN	-.03122	-.17070	-.55481	-.55481	-.55481	-.12111	-.12111	-.12111
V55HE1HN	-.03394	-.37474	-.76014	-.76014	-.76014	-.05049	-.05049	-.05049
V40AP1HN	-.03394	-.37474	-.76014	-.76014	-.76014	-.05714	-.05714	-.05714

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			PAGE: 17	PAGE: 17	
AVERAGED DATA					
DATAC TIMEF	*.07256	*.71866	*.73467	*.09515	*.72060
V4NIE 1#X	*.01028	*.45149	*.21019	*.00849	*.601075
CV34Y	*.04852	*.41696	*.14903	*.11327	*.15742
SVC34X	*.04852	*.22269	*.01506	*.11820	*.01788
FV34Y	*.007201	*.23860	*.313975	*.05052	*.05846
FVC34X	*.01524	*.07879	*.41082	*.05052	*.05846
V25FR34X	*.02855	*.16082	*.31621	*.06816	*.16138
V40FR34X	*.043860	*.02404	*.10871	*.152213	*.16920
CVC34Y	*.1.17220	*.317595	*.161.5933	*.102.28768	*.10.77555
FFPV34X	1.03869	23.48084	49.75939	*11.00361	*5.18428
TVPF34X					
SVC34X	1.01910	*.50321	4.30108	*.39436	
FVC34X	*.08450	*.53961	*.55083	*.39436	
FPC34X	*.20044	*.05829	*.17361	*.07415	*.11122
V25FR34X	*.18539	*.01876	*.15337	*.36640	*.10444
V40FR34X	*.14095	*.74611	*.122.40695	*.18.30107	*.7.9001
CVSV34X	1.08311	*.10.37655	*11.14998	*4.20764	*4.94905
FFPV34X	3.49990	*.96841	*.31956		
TVPF34X	*8.96841				
					1VPF34X
					CVSV34X
					V25FR34X
					FVC34X
					FV34Y
					V40FR34X
					V4NIE 1#X
					DATAC TIMEF

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AVERAGED DATA
DRUG THREE

STANDARDIZED DISCRIMINANT FUNCTION COEFFICIENTS

	1	2	3
V25AP1MN	5.02165	2.63451	3.22711
V25HE1MN	-6.66352	1.30381	-2.19162
V40AP1MN	1.16220	-4.07932	*-0.01236
V40HE1MN	2.38947	2.32081	1.11162
CV1MN	*-11.80457	3.78131	*9.59366
SC1MN	7.24682	-6.74291	5.51044
FEV1MN	1.02751	*-0.04832	2.20517
FVC1=MV	*-2.20576	5.02342	*3.01135
V25SP1MN	-1.23499	1.11462	*-1.77560
V40FP1MN	1.49292	-1.21281	.43690
CV5VMN	8.18173	-1.61524	4.95444
FFV1MN	*-5.71265	.77250	*3.46244
TFP1MN	*-2.32482	1.62220	*.46273

UNSTANDARDIZED DISCRIMINANT FUNCTION COEFFICIENTS

	1	2	3
V25AP1MN	8.01569	3.62615	4.44180
V25HE1MN	*-7.45441	1.45811	*2.45175
V40AP1MN	1.89995	-5.06712	*-1.20355E=01
V40HE1MN	1.72586	1.82080	.80897
CV1MN	*-23.1905	7.42851	*18.0471
SC1MN	3.41801	-2.97507	2.43129
FEV1MN	6.15488	*-5.6054	1.37092
FVC1=MV	*-7.77817	1.77323	*1.2081
V25FP1MN	*10.6159	6.87205	*4.75169
V40FR1MN	15.6197	*9.71904	3.50100
CV5VMN	2.12278	*-429148	1.31633
FFV1MN	*-6.18836	*-836255E=01	*-375077
TFP1MN	*-31.9124	*-222676	*-6351846E=01
CONSTANT	10.7672	*1.66554	10.3756

CONSTRAINTS OF GROUPS IN REDUCED SPACE

	1	2	3
GROUP 1	-3.47603	*-48663	*97168
GROUP 2	-5.26295	*-55973	*98149
GROUP 3	4.73853	1.10145	*-10179
GROUP 4	1.31031	*-0.09305	*-0.05222

AVERAGED DATA
DRUG THREE
PREDICTION RESULTS =

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ACTUAL GROUP NAME	CODE	N OF CASES	PREDICTED GROUP MEMBERSHIP		
			GROUP 1	GROUP 2	GROUP 3
GROUP 1	2	6	4*	2*	0
			20.0 PCT	10.0 PCT	0 PCT
GROUP 2	3	4	1*	3*	0
			5.0 PCT	15.0 PCT	0 PCT
GROUP 3	4	8	0	0	8*
			0 PCT	0 PCT	40.0 PCT
GROUP 4	5	7	0	0	2*
			0 PCT	0 PCT	10.0 PCT
					5*
					25.0 PCT

100.0 PERCENT OF KNOWN CASES CORRECTLY CLASSIFIED

CHI-SQUARE = 60.300 SIGNIFICANCE = .000

AVERAGED DATA
DRUG THREE

STANDARDIZED DISCRIMINANT FUNCTION COEFFICIENTS

	1	2	3
V25AP3MX	2.34427	.90429	*.97201
V25HF3MX	8.41186	*1.51495	3.19351
V40AB3MX	-7.35756	-.81067	-.60042
V40HF3MX	-9.21879	3.01384	*3.27980
CV3MX	*11.13361	2.00072	*28320
SVC3MX	4.26748	-.22833	.53179
FFV3MX	3.36733	*.78845	.80928
FVC3MX	*15.68435	*1.13173	*3.09260
V25PF3MX	7.15231	*1.47907	.58711
V40PF3MX	*3.16337	*.66606	*.79403
CVS3MX	10.57209	-1.02531	*24776
FFF3MX	*17.90855	*.08877	*3.86776
IVFP3MX	3.11708	2.05408	*1.07361

UNSTANDARDIZED DISCRIMINANT FUNCTION COEFFICIENTS

	1	2	3
V25AP3MX	4.16072	1.60499	*1.72516
V25HF3MX	14.88884	*2.6507	5.63906
V40AP3MX	-11.46885	-1.20362	*.95891
V40HF3MX	*8.86738	2.97754	*3.13199
CV3MX	-26.44606	4.79022	*6.2552
SVC3MX	4.88605	*23.990	.546504
FFV3MX	4.98460	*665511	1.13080
FVC3MX	-7.82111	666888E-01	*1.54210
V25PF3MX	11.8340	*2.45551	*97698
V40PF3MX	*9.94694	2.04416	*2.49676
CVS3MX	2.32164	*229158	*544082E-01
FFF3MX	*12.99892	*144812E-03	*6.617659E-01
IVFP3MX	*102895	*876116E-01	*354401E-01
CONSTANT	1.57953	-1.10510	1.65487

CENTROIDS OF GROUPS IN REDUCED SPACE

GROUP	1	2	3
GROUP 2	2.80260	*67707	1.62950
GROUP 3	*2.45881	2.68364	*.95112
GROUP 4	-4.59805	*1.49576	*.01434
	7.00389	*1.03490	*1.12686

AVERAGED DATA
DRUG THREE

PREDICTION RESULTS *

ACTUAL GROUP NAME	GROUP CODE	N OF CASES	PREDICTED GROUP MEMBERSHIP			
			GROUP 1	GROUP 2	GROUP 3	GROUP 4
GROUP 1	2	6	5. 25.0 PCT	0 0 PCT	1. 5.0 PCT	0 0 PCT
GROUP 2	3	4	0 0 PCT	4. 20.0 PCT	0 0 PCT	0 0 PCT
GROUP 3	4	8	0 0 PCT	0 0 PCT	8. 40.0 PCT	0 0 PCT
GROUP 4	5	7	0 0 PCT	0 0 PCT	3. 15.0 PCT	4. 20.0 PCT

100.0 PERCENT OF KNOWN CASES CORRECTLY CLASSIFIED

CHI-SQUARE = 68.267 SIGNIFICANCE = .000

APPENDIX DATA
INPUT FROM
FILE NUMBER CREATION DATE = 31/05/77

GROUP COUNTS

NUMBER	Group 1	Group 2	Group 3	Group 4	TOTAL
4.	4.	4.	4.	14.	

HEADS

	Group 1	Group 2	Group 3	Group 4	TOTAL
V258H10H	1,87600	1,41333	1,00000	1,16667	1,74887
V258H10H	2,37150	1,27000	2,10000	1,21111	7,14001
V258H10H	3,16000	2,08333	3,50000	3,75000	3,45071
V400H10H	4,20000	3,50000	4,78250	5,00000	4,40571
C41H	1,10500	1,11333	1,08750	1,21111	1,24487
AV13H	6,19250	6,29133	10,31500	9,17667	8,21979
FEV13H	6,55250	6,29133	9,00000	8,40000	7,62643
FUV13H	7,18150	8,16333	12,21250	10,96667	9,85214
V25P10H	10,0000	1,15000	10,20500	1,30667	1,24443
V40P10H	1,37250	2,35000	2,40000	3,33333	3,23866
C45P10H	11,24500	16,00000	18,46000	11,90000	14,92643
FPEV13H	84,411750	75,28687	74,02750	71,30000	78,06387
FVEV13H	11,140000	21,169000	20,135000	16,03000	17,06186
V25P10H	11,00000	11,00000	12,26750	11,00000	11,39279
V25H10H	11,06750	1,09667	10,7500	1,20000	1,16443
V40H10H	1,08750	1,16667	1,12500	1,14667	1,07511
V40H10H	1,10250	1,01750	1,05000	1,11333	1,02011
C45H	1,01750	1,03333	1,01750	1,03333	1,01443
SVC10H	4,80000	2,22000	1,0250	4,4333	7,14379
FEV10H	2,22000	1,01000	1,18750	1,01000	1,03000
FVC10H	1,18250	1,02333	1,00500	1,11333	1,12286
V25P10H	1,02670	1,18667	1,12500	1,05333	1,03000
V40P10H	1,26750	1,07000	1,26750	1,10000	1,05114
C45H	1,26750	1,14000	1,11333	1,13111	1,13114
FPEV13H	1,92750	1,66000	1,52750	1,89333	2,04279
FVEV13H	19,58000	9,30667	1,52750	1,51667	2,91000

STANDARD DEVIATIONS

	Group 1	Group 2	Group 3	Group 4	TOTAL
V25H10H	36652	32517	7800	25384	46399
V25H10H	40294	41714	87289	45938	61619
V40H10H	44195	46365	1,24148	45061	78151
V40H10H	26721	49217	1,48006	1,10001	1,19045
C45H	36928	41304	2,39136	1,7502	61307
FUV13H	1,27182	1,14568	1,31721	1,44661	2,10586
FUV13H	1,69416	1,65760	1,12502	1,14240	1,76342
FVEV13H	1,66997	1,6289	1,60622	1,70845	2,32113
FVEV13H	1,62028	1,0132	1,09339	1,0893	1,0893
V40P10H	1,4431	1,04008	1,0499	1,03117	1,2065
V40P10H	5,32038	8,46107	3,69057	5,76466	

STANDARD DATA

NAME	PERIOD, SEC.	PERIOD, SEC.	PERIOD, SEC.
Upsilon Andromedae	10.05051	9.34644	7.54927
51 Pegasi	5.17584	5.92768	4.23574
5555 Pegasi	3.13707	5.50298	3.95233
1755 Pegasi	6.16705	2.07211	3.65288
1754 Pegasi	2.07211	4.48444	3.36729
1754 Pegasi	4.48444	6.49366	5.80000
1754 Pegasi	6.49366	6.44644	5.77443
1754 Pegasi	6.44644	7.31069	6.08135
1754 Pegasi	7.31069	4.08135	2.77449
4409 Eri	2.8779	4.07885	2.25595
4409 Eri	4.07885	4.28779	1.01139
4409 Eri	4.28779	4.77516	0.95677
4409 Eri	4.77516	4.69560	1.05677
4409 Eri	4.69560	4.24331	0.89310
4409 Eri	4.24331	6.10403	0.81189
4409 Eri	6.10403	6.09365	0.02517
4409 Eri	6.09365	1.33899	0.15588
4409 Eri	1.33899	4.06377	0.54244
4409 Eri	4.06377	4.98971	0.54244
4409 Eri	4.98971	7.49971	1.67311
4409 Eri	7.49971	3.45707	3.07551

WITHIN GROUPS COVARIANCE MATRIX

AVERAGED DATA				31/05/77		PAGE 24	
SPIC FOUR							
V40FR3MX	-0.6530	=1.65081	=1.34673	* 50274	.08241	.14645	* 59203
CV3MX	-0.04955	-1.89441	-0.07618	* 62276	.1106	.14940	* 19167
SVC4MX	-0.0492	-1.01470	-0.68103	* 13512	.18162	.28479	-0.0492
FV3MX	-0.0139	* 0.02751	1.70112	* 14395	* 12337	* 10705	-0.0716
FVC4MX	-0.0111	1.60723	2.25127	* 31099	* 12462	* 18760	* 0.9792
W25P3MX	-0.00459	* 0.0501	* 16900	* 37174	* 0.02055	* 0.0189	* 0.0174
V40FR14X	-0.01841	* 0.0919	* 43595	* 0.0501	* 0.02490	* 0.00945	* 0.0110
CV513MX	-0.10815	* 02.56058	4.42122	5.14561	1.39011	1.20926	* 0.06715
FFP3MX	* 10213	* 16.17529	* 11.31110	* 8.86453	1.66116	2.77766	1.66663
TFP3MX	* 14035	21.30575	1.54854	* 38.45867	* 2.00746	* 2.43860	* 3.06070
SVC4MX		FEV3MX	FVC3MX	V25FR3MX	V40FR3MX	FEFP3MX	TFPP3MX
SPIC3MX							
FFP3MX							
V25FR3MX							
W40FR3MX							
CV513MX							
FFP3MX							
TFP3MX							

AVERAGED DATA
DRUG FOUR

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VARIABLES IN THE ANALYSIS

VARIABLE	ENTRY CRITERION	F TO REMOVE
V40F1MN	+1.19626	71.11275
V40H1MN	-0.50004	126.51957
CV1MN	+1.41176	1.07698
SVC1MN	+1.78263	5.61242
FVC1MN	+3.44928	34.59466
V25F1MN	+0.71382	69.26335
V40F1MN	+0.64447	141.53766
CVS1MN	+0.26779	182.02686
FFV1MN	+1.68069	330.71862
TVF1MN	+2.33650	

VARIABLES NOT IN THE ANALYSIS

VARIABLE	TOLERANCE	F TO ENTER	F TO CRITERION
V25A1MN	.00000	0	*.39159
V25H1MN	.00000	0	*.41753
FEV1MN	*.00000	0	*.59163

FUNCTIONS WILL BE USED IN REMAINING ANALYSES

STANDARDIZED DISCRIMINANT FUNCTION COEFFICIENTS

	1	2	3
V40A1MN	-148.97832	8.11685	*6.97375
V40H1MN	209.52969	*10.75946	6.28019
CV1MN	-4.44907	11.46115	8.17953
SVC1MN	20.66412	--	*5.81214
FVC1MN	-43.39498	2.26543	*.01064
V25F1MN	29.34663	-1.27425	*3.11612
V40F1MN	-84.82074	3.45389	*6.34205
CVS1MN	4.04414	-8.27930	
FFV1MN	-47.28598	.48472	*.44073
TVF1MN	52.27710	*.47139	*.40611

UNSTANDARDIZED DISCRIMINANT FUNCTION COEFFICIENTS

	1	2	3
V40A1MN	*190.624	10.7687	*8.92597
V40H1MN	176.008	*9.31812	5.27796
CV1MN	+1.76804	19.3821	13.6905
SVC1MN	9.31411	*2.02878	*2.36416
FVC1MN	-18.7792	*.96001	*.466953E=02
V25F1MN	269.463	*11.2492	*.45109
V40F1MN	*703.027	30.2849	*25.4301

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AVERAGED DATA

DRUG FOUR			
CYSV1MN	.701540	-1.42754	-1.10016
FFPV1MN	-6.26371	.130441	.583804E-01
IVFP1MN	8.38181	-.140034	.651451E-01
CONSTANT	476.656	-1.88555	27.3963

CENTROIDS OF GROUPS IN REDUCED SPACE

GROUP 1	-81.01501	.33726	.59316
GROUP 2	56.02262	-.93008	2.52310
GROUP 3	20.37944	3.13428	-.84225
GROUP 4	24.82481	-2.79929	-2.19098

PREDICTION RESULTS -

ACTUAL GROUP NAME	GROUP CODE	N OF CASES	PREDICTED GROUP MEMBERSHIP			
			GROUP 1	GROUP 2	GROUP 3	GROUP 4
GROUP 1	2	6	6. 42.9 PCT	0 0 PCT	0 0 PCT	0 0 PCT
GROUP 2	3	4	1. 7.1 PCT	3. 21.4 PCT	0 0 PCT	0 0 PCT
GROUP 3	4	8	4. 28.6 PCT	0 0 PCT	4. 28.6 PCT	0 0 PCT
GROUP 4	5	7	0 0 PCT	3. 21.4 PCT	0 0 PCT	4. 28.6 PCT

121.4 PERCENT OF KNOWN CASES CORRECTLY CLASSIFIED

CHI-SQUARE = 69.429 SIGNIFICANCE = .000

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AVERAGED DATA
DRUG FOUR

----- VARIABLES IN THE ANALYSIS -----

VARIABLE	ENTRY CRITERION	F TO REMOVE
V25AR3MX	-.34855	37.87676
V25HF3MX	+3.53194	256.41211
V40AR3MX	-.83428	2.12536
V40HF3MX	-1.26130	268.84208
CV3MX	-2.09950	6.04701
SVC3MX	-.00981	37.25886
FFV3MX	-.58043	14.33494
V40F3MX	-4.48775	299.54704
CVSV3MX	-1.63094	18.29039
FEFV3MX	-.17551	43.47877

----- VARIABLES NOT IN THE ANALYSIS -----

VARIABLE	TOLERANCE	F TO ENTER	ENTRY CRITERION
FVC3MX	-.00000	0	-.15439
V25FR3MX	-.00000	0	-.03200
IVFP3MX	-.00000	0	-.01758

NUMBER REMOVED	EIGENVALUE	CANONICAL CORRELATION	PERCENT OF TRACE	WILKS LAMBDA	CHI-SQUARE	D.F.	SIGNIFICANCE
0	3980.57970	.99987	98.6	.00000	89.07661	30	.000
1	56.54529	.99127	1.4	.01185	31.05057	18	.028
2	.46701	.56422	.0	.68166	2.68256	8	.953

3 FUNCTIONS WILL BE USED IN REMAINING ANALYSES

STANDARDIZED DISCRIMINANT FUNCTION COEFFICIENTS

	1	2	3
V25AR3MX	22.33574	-2.26633	.57047
V25HF3MX	55.48624	.07274	.16666
V40AR3MX	.22951	5.51914	-1.57495
V40HE3MX	-77.14881	-7.32528	-.42414
CV3MX	32.64695	19.15055	1.83763
SVC3MX	-36.51901	-4.48233	-.25479
FFV3MX	-8.29687	4.01097	-.92462
V40F3MX	65.10437	11.76166	.05766
CVSV3MX	-55.52057	-15.32356	-1.31285
FEFV3MX	22.16275	-.28421	1.34015

UNSTANDARDIZED DISCRIMINANT FUNCTION COEFFICIENTS

	1	2	3
V25AR3MX	61.5508	-6.24534	1.57204
V25HF3MX	98.4052	.129003	.295577
V40AR3MX	-.414844	10.0195	-2.85918
V40HE3MX	-97.1385	-9.22331	-.534043
CV3MX	69.4245	40.7241	3.90777
SVC3MX	-44.0100	-5.40178	-.307052
FFV3MX	-14.5146	7.01680	-.1.61753

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AVERAGED DATA

DRUG FOUR			
V40FP3MX	210.424	38.0149	.186375
CVSV3MX	-11.3418	-3.13032	.268191
FFFV3MX	3.81345	-.489021E-01	.230595
CONSTANT	3.78711	2.93436	.288601E-03

CENTROIDS OF GROUPS IN REDUCED SPACE

GROUP 1	-55.77741	-7.24582	.18796
GROUP 2	6.03792	1.47200	-1.09587
GROUP 3	76.99925	-1.79594	.33424
GROUP 4	-34.33371	10.58368	.39961

PREDICTION RESULTS -

ACTUAL GROUP NAME	GROUP CODE	N OF CASES	PREDICTED GROUP MEMBERSHIP			
			GROUP 1	GROUP 2	GROUP 3	GROUP 4
GROUP 1	2	6	5. 35.7 PCT	0 0 PCT	0 0 PCT	1. 7.1 PCT
GROUP 2	3	4	0 0 PCT	3. 21.4 PCT	1. 7.1 PCT	0 0 PCT
GROUP 3	4	8	4. 28.6 PCT	0 0 PCT	4. 28.6 PCT	0 0 PCT
GROUP 4	5	7	1. 7.1 PCT	2. 14.3 PCT	1. 7.1 PCT	3. 21.4 PCT

100 PERCENT OF KNOWN CASES CORRECTLY CLASSIFIED

CHI-SQUARE = 50.381 SIGNIFICANCE = .000

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FACTOR ANALYSIS
FEMALE'S
FILE: RNAME (CREATION DATE = 31/05/77)

FACTOR MATRIX USING PRINCIPAL FACTOR WITH ITERATIONS

	FACTOR 1	FACTOR 2	FACTOR 3	FACTOR 4	FACTOR 5
V75A1P3	.8054	-.34807	-.04079	-.01766	.01670
V75H1P3	.75477	-.05717	.01545	.12073	.07470
V50A1P3	.75851	-.07218	.26338	.09862	.17444
VANIE1	.81985	-.04271	-.15372	.35988	.13612
CV1	.50821	.022601	-.00650	-.29390	.09412
SIC3	.21166	.61744	.00178	-.17645	.37705
FFV1	.52472	.05444	.61135	.08148	.09424
FVC1	.16339	.27790	.91931	-.00772	.06772
V75FPAC3	.16631	.57798	-.09119	.63150	.23296
VADPAC3	.17230	.73427	-.04824	.18967	.21174
CVSIC3	.69213	.25693	-.00637	.59633	.09669
FFUPFVC3	.13809	-.35765	-.35205	-.21754	.34061
TFVP3	.00624	-.23002	.52196	.10258	.34839

MORE THAN 5 ITERATIONS REQUIRED.

VARIABLE	COMMUNALITY	FACTOR	EIGENVALUE	PCT OF VAR	CUM PCT
V75A1P3	.82997	1	3.05563	38.5	38.5
V75H1P3	.68163	2	2.48291	26.8	61.3
V50A1P3	.90891	3	1.76336	17.6	81.0
VANIE1	.94054	4	1.21739	12.2	93.1
CV3	.98453	5	.68846	6.9	100.0
SIC3	.62572				
FFV1	.69213				
FVC1	.05429				
V75FPAC3	.82614				
VADPAC3	.65930				
CVSIC3	.91010				
FFUPFVC3	.61598				
TFVP3	.45727				

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FACTOR ANALYSIS
FFNALES NOME (CREATION DATE # 31/05/77)
FILE

VARIAX ROTATED FACTOR MATRIX
AFTER ROTATION WITH KAISER NORMALIZATION

	FACTOR 1	FACTOR 2	FACTOR 3	FACTOR 4	FACTOR 5
V25AIR3	.79485	.17438	-.12448	.07666	.38257
V25HE3	.73771	*.09211	.28110	.11583	-.19076
V25ATB3	*.87516	*.00978	-.25842	.25901	.10074
V40HHE3	*.91889	*.13078	*.18651	-.21615	*.03525
CV3	*.07962	*.45518	*.17956	-.07214	-.41167
SVC3	*.04819	*.62618	*.16771	-.18347	
FY3	*.40144	*.31998	*.01381	*.65108	*.07094
FNC2	*.15517	*.11144	*.00799	*.74851	*.59788
V25PAC3	*.11046	*.00407	*.89585	-.01583	*.10545
V40PAC3	-.07456	*.39054	*.70428	*.02645	-.06719
CVPVC3	*.27502	*.84758	-.10085	*.10446	.73437
FPYPVC3	*.26579	*.09075	*.11539	*.06446	*.08682
IPPP3	*.02705	*.22477	*.01974	*.63094	

TRANSFORMATION MATRIX

	FACTOR 1	FACTOR 2	FACTOR 3	FACTOR 4	FACTOR 5
FACTOR 1	*.84028	*.44945	*.11321	*.09808	*.26360
FACTOR 2	-.29212	*.61320	*.63458	*.04177	*.36316
FACTOR 3	*.05147	*.00642	*.14675	*.88366	*.44150
FACTOR 4	*.33081	*.66623	*.63982	*.02998	*.24106
FACTOR 5	*.30225	*.06573	*.39285	*.45336	*.73065

FACTOR ANALYSIS

FILE: RONNAME (CREATION DATE = 31/05/77)

FACTOR MATRIX USING PRINCIPAL FACTOR WITH ITERATIONS

	FACTOR 1	FACTOR 2	FACTOR 3	FACTOR 4	FACTOR 5
V7A1R3	.40704	-.40391	.66767	.74069	.06914
V25MF3	.71934	-.30167	.04415	-.07197	-.05817
V401TR3	-.44337	-.37047	.69549	.2294n	-.11004
V401TR3	-.44337	-.37047	.69549	.2294n	-.11004
V40MF3	.79844	-.20819	.11636	-.09788	.02015
CV3	.41705	.80336	.11398	-.35506	.21872
SVC3	.41705	.80336	.11398	-.35506	.21872
FRV3	-.10633	.24469	.05121	.58933	.39167
FIC3	-.44952	-.45037	.59453	-.28627	.26092
V25FRAC3	.62006	.21613	.24061	-.201349	.22876
V25FRAC3	.62006	.21613	.24061	-.201349	.22876
CPSVC3	.26001	.64860	.14573	.37424	.50570
PEFFNFC3	.44027	-.07730	-.77716	.43406	.02028
IVFP3	-.26120	-.30957	.20461	.04692	.13797

3 ITERATIONS BECAUSE COMMUNALITIES EXCEED ONE.

ITERATIVE PROCEDURE STOPPED AFTER

C	VARIABLE	COMMUNALITY	FACTOR	EIGENVALUE	PCT OF VAR	CUM PCT
C	V7A1P3	.84173	1	3.35479	34.6	34.6
C	V25MF3	.64935	2	2.20492	22.0	57.4
C	V401TR3	.88226	3	2.09327	21.6	79.0
C	V40HE3	.74170	4	1.20419	13.4	92.4
C	CV3	1.00348	5	.73757	7.6	100.0
C	SVC3	.73580				
C	FRV3	.51600				
C	FVC3	.90168				
C	V25PAC3	.58282				
C	V40PAC3	.91144				
C	CPSVC3	.69936				
C	PEFFNFC3	.99261				
C	IVFP3	.14334				

FACTOR ANALYSIS
MALES
FILE NODNAME (CREATION DATE = 31/05/77)

VARMAX ROTATED FACTOR MATRIX
AFTER ROTATION WITH KAISER NORMALIZATION

	FACTOR 1	FACTOR 2	FACTOR 3	FACTOR 4	FACTOR 5
V25AIR3	*.91214	*.04615	*.10550	*.02059	*.11699
V25MF3	*.51378	*.04822	*.34993	*.41428	*.23927
V4DAIR3	*.93205	*.06784	*.07595	*.05665	*.01059
V4DME3	*.60826	*.05511	*.28853	*.49884	*.21218
CY3	*.07827	*.97511	*.00876	*.14021	*.15665
SNC1	*.02799	*.70846	*.09156	*.25668	*.41045
FK3	*.02011	*.10017	*.01394	*.06170	*.74943
FVC3	*.13919	*.15523	*.92161	*.01477	*.11765
V25FRAC3	*.00481	*.23837	*.22570	*.68777	*.04487
V4DFRAC3	*.09162	*.10320	*.02688	*.94112	*.01172
CIPS4C3	*.03697	*.20577	*.01579	*.04368	*.44483
PPPF4C3	*.14369	*.03632	*.94015	*.20128	*.21519
TWPP3	*.05266	*.19823	*.24670	*.15133	*.13525

TRANSFORMATION MATRIX

	- - - FACTOR 1 - - -	- - - FACTOR 2 - - -	- - - FACTOR 3 - - -	- - - FACTOR 4 - - -	- - - FACTOR 5 - - -
FACTOR 1	*.52140	*.36929	*.42562	*.62988	*.111733
FACTOR 2	*.46834	*.80614	*.26144	*.13113	*.21197
FACTOR 3	*.66563	*.17779	*.70274	*.17190	*.05211
FACTOR 4	*.24542	*.15239	*.41594	*.43899	*.74459
FACTOR 5	*.07362	*.39826	*.28921	*.60354	*.02101

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FACTOR ANALYSIS
NIN=SMOKERS (CREATION DATE = 31/05/77)
FILE NURNAME
PRINCIPAL FACTOR WITH ITERATIONS

FACTOR MATRIX USING PRINCIPAL FACTOR WITH ITERATIONS

	FACTOR 1	FACTOR 2	FACTOR 3	FACTOR 4	FACTOR 5
V75A1R3	.44488	.53186	.57592	.17720	.11863
V75H3	.67176	-.02110	-.20114	-.05672	-.01988
V75A1R3	.50141	.60484	-.48760	-.16268	-.01360
V75H3	.83528	.05694	-.32183	-.06010	-.12435
V75H3	.32401	.59448	.66008	.10484	-.09100
CV3	.31512	.36296	.49292	-.48141	-.30980
SVC1	-.16659	.18961	.15101	.51883	.35171
FV3	-.63472	.56820	.05585	.36638	.23516
FV3	.66540	-.28343	.23963	-.28239	.28963
V75FAC3	.65635	-.08373	.11245	-.42963	.41634
V40FAC3	.19344	.54040	.53966	.43869	.07061
CIPSIC3	.52359	-.64041	.32105	-.48949	.01380
FEVPPVC3	-.24889	.06126	-.09148	-.10563	.32834

ITERATIVE PROCEDURE STOPPED AFTER 3 ITERATIONS BECAUSE COMMUNALITIES EXCEED ONE.

VARIABLE	COMMUNALITY	FACTOR	EIGENVALUE	PCT OF VAR	CUM PCT
V75A1R3	.85195	1	3.49527	35.4	35.4
V75H3	.49669	2	2.28868	23.3	58.7
V75A1R3	.88100	3	1.95706	19.9	78.6
V75H3	.82358	4	1.19346	14.4	93.0
V75H3	.97532	5	.68389	7.0	100.0
CV3	.80162				
SVC1	.47904				
FV3	.91984				
V75FAC3	.80206				
V40FAC3	.88337				
CIPSIC3	.81759				
FEVPPVC3	1.02714				
FEVPPVC3	.19303				
TVP3					

FACTOR ANALYSIS
NON-SMOKERS CREATION DATE = 31/05/77
FILE NOME

VARIMAX ROTATED FACTOR MATRIX
AFTER ROTATION WITH KAISER NORMALIZATION

	FACTOR 1	FACTOR 2	FACTOR 3	FACTOR 4	FACTOR 5
V25AIR3	.91405	-.08457	.00929	.01146	-.112287
V25WE3	.47631	.30846	-.01614	.38568	-.16019
V4AIR3	.92443	-.02873	.16291	-.03195	-.00471
V4WNE3	.68245	-.38171	.00776	.36847	-.21692
V4WNE3	.07549	.01255	.95160	.11155	.01835
CV3	-.02525	-.10384	.65316	.25149	.54801
SVC3	-.01147	-.04086	.18220	-.13248	.65796
FFV3	-.14252	-.91533	.15921	-.18810	.03112
FVC3	-.04507	.28508	.13557	.83164	.09152
V25PAC3	-.17642	.01863	.05125	.87230	.11557
V4OPAC3	.10843	.06782	.00346	-.03736	.39280
CPASC3	-.17905	.90889	.01129	.31364	.26552
PPVPPC3	-.06264	-.35531	-.15844	.05793	.10548

TRANSFORMATION MATRIX

	FACTOR 1	FACTOR 2	FACTOR 3	FACTOR 4	FACTOR 5
FACTOR 1	.54590	.51989	.22656	.59714	-.15461
FACTOR 2	.52104	-.54830	.61433	-.21923	.04916
FACTOR 3	-.63859	.12092	.72323	.22242	.07111
FACTOR 4	.15102	.51493	.11538	-.44729	.70616
FACTOR 5	-.00359	.38640	-.18662	.58807	.68584

FACTOR ANALYSIS
 STUDENTS FILE NODNAME (CREATION DATE = 31/05/77)
 FACTOR MATRIX USING PRINCIPAL FACTOR WITH ITERATIONS

	FACTOR 1	FACTOR 2	FACTOR 3	FACTOR 4
V25A1B3	.52718	.78735	.16169	-.11039
V25H3	.72809	.42697	-.31072	.01485
V40A1H3	.35316	.82718	-.08713	.01848
V40H3	.35316	.27440	-.61603	.11926
V40HE3	.73211	-.47587	.21807	-.33776
C11	.76123	-.44038	.14645	-.17423
SNC3	.53822	-.05865	.30146	.45239
FFV3	.49608	-.34103	.18049	.74959
FVC3	.42808	-.26678	-.24393	-.17233
V25PAC3	.47642	-.41905	-.26217	-.08107
V40PAC3	.53413	-.21093	-.39596	-.14194
CVPSC3	.58033	-.28459	.55668	-.33522
FEVPFVC3	.24567	-.36252	.26103	.28868
IVFP3	-.08146			

ITERATIVE PROCEDURE STOPPED AFTER 3 ITERATIONS BECAUSE COMMUNALITIES EXCEED ONE.

VARIABLE	COMMUNALITY	FACTOR	EIGENVALUE	PCT OF VAR	CUM PCT
V25A1B3	.93618	1	3.66656	.61.8	41.8
V25H3	.80918	2	2.62271	.29.8	71.6
V40A1P3	.81691	3	1.36509	.15.3	87.1
V40H3	1.00550	4	1.13880	12.9	100.0
C11	.91305				
SVC3	.51542				
FFV3	.56963				
FVC3	.89401				
V25PAC3	.39734				
V40PAC3	.55508				
CVPSC3	.55844				
FEVPFVC3	.56340				
IVFP3	.28953				

FACTOR ANALYSIS
 SNOFFS FILE NAME (CREATION DATE = 31/05/77)
 VARIMAX ROTATED FACTOR MATRIX
 AFTER ROTATION WITH KAISER NORMALIZATION

	FACTOR 1	FACTOR 2	FACTOR 3	FACTOR 4
V25AIR3	-.15911	.80604	.011917	.51099
V25ME3	-.25483	.85465	.11249	.03467
V26AIR3	-.37627	.76371	.01569	.15631
VOLUME1	.38257	.86856	.12214	-.39886
CV3	.82411	.02821	.30585	.37357
SVC3	.65149	-.05168	.23025	.23516
FFT3	.10172	.16305	.69648	.16616
FVC3	.15033	-.01854	.92158	-.14748
V25FAC3	.59163	.18050	.02924	.06226
V40FAC3	.70902	.10387	-.14937	-.13981
CV8NC3	.46375	.04355	.31391	.47289
FV8FVC3	-.02085	.12416	-.03066	.73933
IVP3	-.45255	.12237	.21481	.15365

TRANSFORMATION MATRIX

	FACTOR 1	FACTOR 2	FACTOR 3	FACTOR 4
FACTOR 1	.63013	.59859	.41645	.26681
FACTOR 2	-.63058	.70631	-.21463	.23984
FACTOR 3	-.22877	-.37520	.35749	.82407
FACTOR 4	-.39111	.04511	.80790	-.43885

FACTOR ANALYSIS
SMOKERS
FILE NONAME (CREATION DATE = 31/05/77)

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VARIMAX ROTATED FACTOR MATRIX
AFTER ROTATION WITH KAISER NORMALIZATION

	FACTOR 1	FACTOR 2	FACTOR 3	FACTOR 4
V25AIR3	-.15811	.00604	.01917	.51099
V25HE3	.25483	.05465	.11249	.03402
V40AIR3	-.32627	.76371	.01569	.35631
V40HE3	.38257	.06856	.12214	-.29886
CV3	.82411	.02821	.30585	.17357
SVC3	.65149	-.05168	.23025	.23516
FFV3	.10172	.16305	.69648	.16616
FVC3	.15033	-.01854	.92158	-.14748
V25FPAC3	.59163	.18050	.02924	-.06226
V40FPAC3	.70902	.10387	.14937	-.13881
CVPSPC3	.46375	.04355	.31391	.49289
FFVPPFVC3	-.02085	.12416	-.03066	.73933
IVFP3	-.45255	.12237	.21481	.15365

TRANSFORMATION MATRIX

	FACTOR 1	FACTOR 2	FACTOR 3	FACTOR 4
FACTOR 1	.63013	.59859	.41645	.26681
FACTOR 2	-.63058	.70631	-.21463	.23964
FACTOR 3	-.22877	-.37520	.35749	.82407
FACTOR 4	-.39111	.04511	.80790	-.43851

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AVERAGED DATA

FEMALE (CREATION DATE = 02/06/77)

***** MULTIPLE REGRESSION *****

DEPENDENT VARIABLE.. V25AR1MN

MEAN RESPONSE 2.07184 STD. DEV. .71178

VARIABLE(S) ENTERED ON STEP NUMBER 1.. HEIGHT
WEIGHT
AGE
CIGTOT

		ANALYSIS OF VARIANCE	DF	SUM OF SQUARES	MEAN SQUARE	F	SIGNIFICANCE
MULTIPLE R	.84748	REGRESSION	4.	2.18323	.54581	1.27446	.484
R SQUARE	.71822	RESIDUAL	7.	.85653	.42826		
STD DEVIATION	.65442						

COEFFICIENT OF VARIABILITY 32.367 PERCENT

----- VARIABLES IN THE EQUATION -----

VARIABLE	B	STD ERROR B	T	SIGNIFICANCE	
				ELASTICITY	
HEIGHT	4.1622103	6.1254440	.46171827	.2604017	3.41354
WEIGHT	-.28834695E-01	.48934730E-01	.34721329	-.2262556	-.83029
AGE	.43782380E-01	.77110066E-01	.37238631	.2533791	
CIGTOT	.77509954E-04	.36168554E-04	4.5925370	.9837709	
(CONSTANT)	-6.7296037	10.998369	.37438843	1.16679	
			.603		

ALL VARIABLES ARE IN THE EQUATION.

COEFFICIENTS AND CONFIDENCE INTERVALS.

VARIABLE	B	STD ERROR B	T	95.0 PCT CONFIDENCE INTERVAL	
HEIGHT	4.1622103	6.1254440	.67949855	-.22.193718	.30.518178
WEIGHT	-.28834695E-01	.48934730E-01	-.58924807	-.23938616	.18171677
AGE	.43782380E-01	.77110066E-01	.56779073	-.28799910	.37556386
CIGTOT	.77509954E-04	.36168554E-04	2.1430205	-.78112484E-04	.23313219E+03
CONSTANT	-6.7296037	10.998369	-.61187289	-.54.052285	.40.593078

64

MULTIPLE REGRESSION					
VARIABLE(S) ENTERED ON STEP NUMBER 1..					
MULTIPLE R	.86927	ANALYSIS OF VARIANCE	DF	SUM OF SQUARES	F
R SQUARE	.75563	REGRESSION	4..	4.21411	1.44610
STD DEVIATION	.82550	RESIDUAL	2.	1.36289	.429
COEFFICIENT OF VARIABILITY	32.814 PERCENT				
VARIABLES IN THE EQUATION					
VARIABLE	B	STD ERROR B	F	BETA	VARIABLE
WEIGHT	5.7238609	7.7267450	.54876171	.2641712	WEIGHT
WEIGHT	*.52077144E+01	*.61727147E+01	.71177365	.516	AGE
AGE	*21156028E+01	*.97268021E+01	.57657767E+01	.488	AGE
CIGTOT	*10053576E+03	*.45623663E+04	4.8558000	.833	CIGTOT
(CONSTANT)	*7.6272697	13.873540	*.20224815	1.21632	(CONSTANT)
				.638	

ALL VARIABLES ARE IN THE EQUATION.

COEFFICIENTS AND CONFIDENCE INTERVALS.

VARIABLE	B	STD ERROR B	T	95.0 % CT CONFIDENCE INTERVAL
WEIGHT	5.7238609	7.7267450	.7407653	*.5722005
WEIGHT	*.52077144E+01	*.61727147E+01	*.84166679	*.11767054
AGE	*21156028E+01	*.97268021E+01	.24012032	*.19515902
CIGTOT	*10053576E+03	*.45623663E+04	2.2035880	*.95769179E+04
CONSTANT	*7.6272697	13.873540	*.54497099	*.320949

MULTIPLE REGRESSION									
DEPENDENT VARIABLE..		VARIABLE(S) ENTERED ON STEP NUMBER 1..		ANALYSIS OF VARIANCE		SUM OF SQUARES		F SIGNIFICANCE	
MEAN RESPONSE	3.31243	STD. DEV.	.93617	DF		MEAN SQUARE		F	.29564
R MULTIPLE	.60957	HEIGHT	1.95380		4.	*48447			.862
R SQUARE	.37157	WEIGHT	1.65528		2.	3.30456			
STD DEVIATION	1.28541	AGE							
Coefficient of Variability	38.806 PERCENT	CIGTOT							
VARIABLES IN THE EQUATION									
VARIABLE	B	STD ERROR B		F		NETA		VARIABLE	
HEIGHT	5.0504117	12.031601		.17620034		.2402344		PARTIAL	
WEIGHT	*.10766565E-01	*.96117625E-01		.23608946E-01		.252817		TOLERANCE	
AGE	.34659652E-01	*.15145964		.521666623E-01		.1525058			
CIGTOT	*.71473787E-04	*.71042295E-04		1.0121844		.27949			
(CONSTANT)	-7.3032731	21.003002		.11428964		.689219			
				.768		.65672			

ALL VARIABLES ARE IN THE EQUATION.

COEFFICIENTS AND CONFIDENCE INTERVALS.			
VARIABLE	B	STD ERROR B	T
HEIGHT	5.0504117	12.031601	.41976725
WEIGHT	*.14768665E-01	*.96117625E-01	*.15365190
AGE	*.34659652E-01	*.15145964	.22883755
CIGTOT	*.71473787E-04	*.71042295E-04	1.0060737
CONSTANT	-7.3032731	21.003002	*.33806750

95.0 PCT CONFIDENCE INTERVAL			
VARIABLE	B	STD ERROR B	T
HEIGHT	56.818780	.46.7117956	
WEIGHT	*.19879665	*.42811396	
AGE	*.61010237	*.68614503	
CIGTOT	*.23419989E-03	*.37714747E-03	
CONSTANT	*.100425851	*.85.647965	

AVERAGED DATA

FILENAMES CREATION DATE = 02/06/77

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PDT:

***** MULTIPLE REGRESSION *****

DEFENDANT VARIABLE..*

VAGHEMIN

MEAN PESNPWF 4.16444 STD. DEV. 1.06482

VARIABLE(S) ENTERED ON STEP NUMBER 1..

HEIGHT

WEIGHT

AGE

CIGTOT

COEFFICIENT OF VARIABILITY

27.814 PERCENT

***** F SIGNIFICANCE *****

.76764 .633

MEAN SQUARE

1.02993

1.34168

1.34168

2.68336

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AVERAGED DATA

FILE NAME (CREATION DATE = 02/06/77)

FILE NUMBER

DEPENDENT VARIABLE = C1INN

DEPENENT VARIABLE = .90303 STD. DEV.

MEAN RESPONSE = .39271

VARIABLE(S) ENTERED ON STEP NUMBER 1..

HEIGHT

WEIGHT

AGE

CIGTOT

MULTIPLE REGRESSION

F SIGNIFICANCE

.676

MEAN SQUARE

.12502

R SQUAR

.18935

STD DEVIATION

.37870

ANALYSIS OF VARIANCE

.50009

DF

4.

SUM OF SQUARES

.37870

REGRESSION

2.

RESIDUAL

2.

COEFFICIENT OF VARIABILITY = 46.187 PERCENT

VARIABLES NOT IN THE EQUATION

F SIGNIFICANCE

.66027

.676

VARIABLES IN THE EQUATION

F SIGNIFICANCE

.676

VARIABLE B

F

SIGMA

VARIABLE

SIGMA

ANALYSIS OF VARIANCE

.1444015

FLASTICITY

HEIGHT

.928366015E-01

.1444015

WEIGHT

.227876

.227876

AGE

.6204619

.6204619

CIGTOT

.74102

.74102

(CONSTANT)

.0104716

.0104716

.30491194E-02

.08374

.08374

.961

.5384367

.5384367

.89955175

.76878

.76878

.443

.3131775

.3131775

.733

.39109222

.39109222

ALL VARIABLES ARE IN THE EQUATION.

COEFFICIENTS AND CONFIDENCE INTERVALS.

VARIABLE	B	STD ERROR B	T	95.0 PCT CONFIDENCE INTERVAL
HEIGHT	1.2410091	4.0130094	.31066499	-16.281928
WEIGHT	.4516184E-01	.32538313E-01	1.3066499	*.91486415E-01
AGE	-.2831236E-02	.51273021E-01	-.55218832E-01	*.22344367
CIGTOT	-.22809311E-04	.24049662E-04	*.94844702	*.12628831E-01
CONSTANT	-.2.8601268	7.3131775	-.39109222	*.34326536

AVERAGED DATA

FEMALE
FILE NODNAME (CREATION DATE = 02/06/77)

***** MULTIPLE REGRESSION *****

DEPENDENT VARIABLE.. SVC1MN

MEAN RESPONSE 7.00412 STD. DEV. 1.74090

VARIABLE(S) ENTERED ON STEP NUMBER 1.. HEIGHT
WEIGHT
AGE
CIGTOT

		ANALYSIS OF VARIANCE	DF	SUM OF SQUARES	MEAN SQUARE	F SIGNIFICANCE
MULTIPLE R	.62613	REGRESSION	4.	7.12904	1.78226	.32242
R SQUARED	.39204	RESIDUAL	2.	11.05539	5.52770	.846
STD DEVIATION	2.35111					

COEFFICIENT OF VARIABILITY 33.567 PERCENT

----- VARIABLES IN THE EQUATION -----

VARIABLE	B	STD ERROR B	F		BETA
			SIGNIFICANCE	ELASTICITY	
HEIGHT	18.609716	22.006644	.7151033	.4760220	
WEIGHT	.11644402	.17580590	.43869985	.3735690	
AGE	-.92870355E+01	.27703033	.11238273	.96789	
CIGTOT	-.23801784E-04	.12994136E-03	.33552449E-01	.2197446	
(CONSTANT)	-27.427876	39.513412	.48183163	-.1235138	
			.559	-.10343	

ALL VARIABLES ARE IN THE EQUATION.

COEFFICIENTS AND CONFIDENCE INTERVALS.

VARIABLE	B	STD ERROR B	T	95.0 PCT CONFIDENCE INTERVAL	
				L	U
HEIGHT	18.609716	22.006644	.84564078	-76.078273	111.29770
WEIGHT	.11644402	.17580590	.66234421	-.63999603	.87288408
AGE	-.92870355E+01	.27703033	-.33523533	-1.284888	1.0991081
CIGTOT	-.23801784E-04	.12994136E-03	-.18317328	-.58290046E-03	.53529690E-03
CONSTANT	-27.427876	39.513412	-.69414093	-197.44223	142.58648

AVERAGED DATA

FEMALE
FILE NONAME (CREATION DATE = 02/06/77)

***** MULTIPLE REGRESSION *****

DEPENDENT VARIABLE.. FEV1MM

MEAN RESPONSE 6.84421 STD. DEV. 1.46619

VARIABLE(S) ENTERED ON STEP NUMBER 1.. HEIGHT
WEIGHT
AGE
CIGTOT

	MULTIPLE R	.84811	ANALYSIS OF VARIANCE	DF	SUM OF SQUARES	MEAN SQUARE	F SIGNIFICANCE
R SQUARE	.71930		REGRESSION	4.	9.27771	2.31943	1.28125
STD DEVIATION	1.34547		RESIDUAL	2.	3.62058	1.81029	.483

COEFFICIENT OF VARIABILITY 19.659 PERCENT

VARIABLES IN THE EQUATION					VARIABLES NOT IN THE EQUATION			
VARIABLE	B	STD ERROR B	F SIGNIFICANCE	BETA	VARIABLE	PARTIAL	TOLERANCE	F SIGNIFICANCE
HEIGHT	17.362593	12.593767	1.9007189	.5273339				
			.302	4.20646				
WEIGHT	.57555168E-01	.10060864	.32726392	.2192410				
			.625	.48958				
AGE	-.20425615	.15853645	1.6599385	-.5738521				
			.327	-.79714				
CIGTOT	-.10735018E-03	.74361684E-04	2.0840451	-.6614434				
			.286	-.47738				
(CONSTANT)	-16.573413	22.612384	.53719407					
			.540					

ALL VARIABLES ARE IN THE EQUATION.

COEFFICIENTS AND CONFIDENCE INTERVALS.

VARIABLE	B	STD ERROR B	T	95.0 PCT CONFIDENCE INTERVAL	
HEIGHT	17.362593	12.593767	1.3786656	-36.824606	.71.549793
WEIGHT	.57555168E-01	.10060864	.57206985	-.37533361	.49044395
AGE	-.20425615	.15853645	-1.2883860	-.88639095	.47787865
CIGTOT	-.10735018E-03	.74361684E-04	-1.4436222	-.42730620E-03	.21260584E-03
CONSTANT	-16.573413	22.612384	-.73293524	-113.86772	.80.720890

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AVERAGED DATA

FF'ALFS

FILE NONAME (CREATION DATE = 02/06/77)

***** MULTIPLE REGRESSION *****

DEPENDENT VARIABLE.. FVCIMN

MEAN RESPONSE 8.38050 STD. DEV. 1.69663

VARIABLE(S) ENTERED ON STEP NUMBER 1.. HEIGHT
WEIGHT
AGE
CIGTOT

MULTIPLE R	1.00000	ANALYSIS OF VARIANCE	DF	SUM OF SQUARES	MEAN SQUARE	F	SIGNIFICANCE
R SQUARE	1.00000	REGRESSION	4.	17.27138	4.31784	R	0
STD DEVIATION	0	RESIDUAL	2.	0	0		

COEFFICIENT OF VARIABILITY 0 PERCENT

----- VARIABLES IN THE EQUATION -----

VARIABLE	B	STD ERROR B	F	BETA	
				SIGNIFICANCE	ELASTICITY
HEIGHT	14.213542	0	0	R .3730577	2.81228
WEIGHT	.16099471	0	0	R .5299704	1.11841
AGE	-.35593145	0	0	R -.8641592	-.13443
CIGTOT	-.26705199E-03	0	0	R -.1.4219629	-.96986
(CONSTANT)	-6.9256202	0	0	R	

----- VARIABLES NOT IN THE EQUATION -----

VARIABLE	PARTIAL	TOLERANCE	F
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MATRIX IS SINGULAR - UNIQUE ESTIMATES ARE IMPOSSIBLE.

COEFFICIENTS AND CONFIDENCE INTERVALS.

VARIABLE	B	STD ERROR B	T	95.0 PCT CONFIDENCE INTERVAL	
				14.213542	14.213542
HEIGHT	14.213542	0	.10000000E+76	.16099471	.16099471
WEIGHT	.16099471	0	.10000000E+76	-.35593145	-.35593145
AGE	-.35593145	0	.10000000E+76	-.26705199E-03	-.26705199E-03
CIGTOT	-.26705199E-03	0	.10000000E+76	-6.9256202	-6.9256202
CONSTANT	-6.9256202	0			

71

AVPARF01 DATA
 FINFILES (CREATION DATE = 02/06/77)
 FILE NODATE

 INDEPENDENT VARIABLE(S) ENTERED ON STEP NUMBER 1..
 MEAN RESPONSE *31973 STD. DFY.
 VARIABLE(S) V75FRHM HE

				MEAN SQUARE	F	SIGNIFICANCE
MULTIPLE R	.47291	ANALYSIS OF VARIANCE	4.	.03062	.00766	* .14403
R SQUARED	.22664	REGRESSION		* .10630	* .05115	
STD DEVIATION	.23054	RESIDUAL	2.			
Coefficient of Variability	72.104 PERCENT					

VARIABLE	B	STD. ERROR B
VARIABLES IN THE EQUATION	6	

***** VARIABLES NOT IN THE EQUATION *****							
F	BETA	ELASTICITY	VARIABLE	PARTIAL	TOLERANCE	F	SIGNIFICANCE
.9951	.2702514						
	.7112	4.75446					
	.144421E-02						
	.935	.053046					
	.28715						
	.8092E+02						
	.0456728						
	.936						
	.13992						
	.4027064						
	.00755						
	.6640						
	.644100						
	.744						

ALL VARIABLES ARE IN THE EQUATION.

COEFFICIENTS AND CONFIDENCE INTERVALS.

COEFFICIENTS AND CONFIDENCE INTERVALS.		95.0 PCT CONFIDENCE INTERVAL			
VARIABLE	B	STD ERROR B	T		
HFIGHT	.91676558	2.1578693	.42484759	=.3676888	10.201410
wFIGHT	.157698326E-02	.172387105E-01	.91479189E+01	=-.72566156E-01	.7549882E-01
AGE	-.167491905E-02	.27164308E-01	-.61658813E-01	-.11855719	.11520495
CIGOUT	.67157505E-05	.1414464E-04	.47446953E-04	=.4808422E-04	.61556417E-04
CONSTANT	.014524118	.387450516E-	.274469932	=.18E-12550	.15.718386

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RENAME NONAME (CREATION DATE = 02/06/77)

***** MULTIPLE REGRESSION *****

DEPENDENT VARIABLE.. V40FRMIN

MEAN RESPONSE .31750 STD. DEV. .10530

VARIABLE(S) ENTERED ON STEP NUMBER 1.. HEIGHT
WEIGHT
AGE
CIGTOT

		ANALYSIS OF VARIANCE	DF	SUM OF SQUARES	MEAN SQUARE	F	SIGNIFICANCE
MULTIPLE R	.64538	REGRESSION	4.	.02771	.00693	.35691	.827
R SQUARE	.41651	RESIDUAL	2.	.03882	.01941		
STD DEVIATION	.13932						

COEFFICIENT OF VARIABILITY 43.879 PERCENT

----- VARIABLES IN THE EQUATION -----

VARIABLE	B	STD ERROR B	F		BETA	PARTIAL	TOLERANCE	F
			SIGNIFICANCE	ELASTICITY				
HEIGHT	.53784175	1.3040143	.17011567	.2274533				
WEIGHT	.50893681E-02	.10417463E-01	.23867326	.2.80890				
AGE	.39554830E-02	.16415566E-01	.58061378E-01	.2699404				
CIGTOT	.83389701E-05	.76997377E-05	1.1729324	.93321				
(CONSTANT)	-.63748712	2.3413862	.74130463E-01	.1547359				
			.811	.33277				
				.7154329				
				.79938				

ALL VARIABLES ARE IN THE EQUATION.

COEFFICIENTS AND CONFIDENCE INTERVALS.

VARIABLE	B	STD ERROR B	T	95.0 PCT CONFIDENCE INTERVAL	
HEIGHT	.53784175	1.3040143	.41245081	-.5.0729406	.6.1486241
WEIGHT	.50893681E-02	.10417463E-01	-.48854197	-.49912587E-01	.39733851E-01
AGE	.39554830E-02	.16415566E-01	.24095929	-.66675772E-01	.74588738E-01
CIGTOT	.83389701E-05	.76997377E-05	1.0830200	-.24790691E-04	.41468631E-04
CONSTANT	-.63748712	2.3413862	-.27226910	-.10.711770	.9.4367954

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AVERAGED DATA
 FIELDS: NICKNAME (CREATION DATE = 02/06/77) * MULTIPLE REGRESSION
 FILE: C:\SUSY\INN
 * * * * *
 DEPENDENT VARIABLE: CIGS1MN
 MEAN RESPONSE: 13.12706 STD. DEV.: 4.58558
 VARIABLE(S) ENTERED ON STEP NUMBER 1:
 HEIGHT WEIGHT
 AGE CIGTOT
 MEAN SQUARE: 14.90439
 SUM OF SQUARES: 59.61756
 33.27398
 R SQUARF: .67241
 R SQUARF: .66.54795
 STD DEVIATION: 5.76836
 COEFFICIENT OF VARIABILITY: 43.943 PERCENT

VARIABLES IN THE EQUATION			VARIABLES NOT IN THE EQUATION		
VARIABLE	B	STD ERROR B	F	VARIABLE	PARTIAL TOLERANCE
WEIGHT	3.6727412	.53.092577	+46271388E+02	BETA	
WEIGHT	*37194567	*.43133399	*.74358676	SIGIFICANCE	ELASTICITY
HEIGHT	*61743599	*.67966840	*.82521812		*.0366662
AGE	*-21152854E-04	*.31880664E-03	*.44023308E-02		*.46393
CIGTOT	(CONSTANT)	96.944855	.98755221E-01		*.4510151
	*30.465254		*.783		*.64958

ALL VARIABLES ARE IN THE EQUATION.

COEFFICIENTS AND CONFIDENCE INTERVALS		95.0 PCT CONFIDENCE INTERVAL	
VARIABLE	B	STD ERR B	T
WEIGHT	3.6727412	.53.092577	*6002207E=01
HEIGHT	*37194567	*.43133399	*.8231477
AGE	*61743599	*.67966840	*.9841517
CIGTOT	*-21152854E-04	*.31880664E-03	*.66150063E=01
CONSTANT	*30.465254	96.944855	*.31425343

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AVERAGED DATA

PPVALSF FILE: NODNAME: (CREATION DATE: 02/06/77)

***** MULTIPLE REGRESSION *****

DEFINING VARIABLE: PEFYINN

MEAN RESPONSE 82.58421 STD. DEV. 6.96199

VARIABLE(S) ENTERED ON STEP NUMBER 1.

HEIGHT

WEIGHT

AGE

CIGTOT

COEFFICIENT OF VARIABILITY 6.281 PERCENT

VARIABLES IN THE EQUATION

R STD ERROR R

F SIGNIFICANCE

BETA

ELASTICITY

VARIABLE	R	STD ERROR R	F	SIGNIFICANCE	BETA	ELASTICITY
HEIGHT	.381450090	.485515680	.622174227	*.2459180		
WEIGHT	-.025148889	.387867055	4.53800108	*.511 .77202		
AGE	.54165656	.61119073	*.78540653	*.6621131		
CIGTOT	.67320584E-03	*.28667963E-03	5.5157593	*.17519		
(CONSTANT)	31.918533	.87175402	*.130405960	*.8716683		

ALL VARIABLES ARE IN THE EQUATION.

COEFFICIENTS AND CONFIDENCE INTERVALS.

VARIABLE	R	STD ERROR R	T	95.0 PCT CONFIDENCE INTERVAL
HEIGHT	.38450090	.48551568	*.79194332	*.17045274
WEIGHT	-.025148889	.38786705	*.21279110	*.4924244
AGE	.54165656	.61119073	*.88623164	*.08861118
CIGTOT	*.67320584E-03	*.28667963E-03	*.36856534	*.56021058E-03
CONSTANT	31.918533	.87175402	*.13614150	*.34317107

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AVERAGED DATA

FVALUES
FILE NODATE (CREATION DATE = 02/06/77)

***** MULTIPLE REGRESSION *****

DEPENDENT VARIABLE.. IVFP1MN

MEAN RESPONSE 25.68606 STD. DEV. 23.31861

VARIABLE(S) ENTERED ON STEP NUMBER 1.. HEIGHT
WEIGHT
AGE
CIGTOTMULTIPLE R .47117 ANALYSIS OF VARIANCE DF SUM OF SQUARES MEAN SQUARE F SIGNIFICANCE
R SQUARE .22200 REGRESSION 4 724.29166 181.07342 .14268 .951
STD DEVIATION 35.62480 RESIDUAL 2 2538.25235 1269.12617
COEFFICIENT OF VARIABILITY 138.693 PERCENT

----- VARIABLES IN THE EQUATION -----

VARIABLE	B	STD ERROR B	F	BETA
			SIGNIFICANCE	ELASTICITY
HEIGHT	-125.31814	331.45260	.14124063	-.2393166
WEIGHT	.96038341	2.6638743	.12997560	-.08988
AGE	-.23636856	4.1976634	.31707659E-02	.2300222
CIGTOT	.57370250E-03	.19689183E-02	.84902038E-01	.17675
(CONSTANT)	166.42380	598.72145	.77264720E-01	-.0417544
			.807	-.24580

ALL VARIABLES ARE IN THE EQUATION.

COEFFICIENTS AND CONFIDENCE INTERVALS.

VARIABLE	B	STD ERROR B	T	95.0 PCT CONFIDENCE INTERVAL	
				L	U
HEIGHT	-125.31814	331.45260	-.37581995	-1560.0646	1109.4284
WEIGHT	.96038341	2.6638743	.36052129	-10.501469	12.42236
AGE	-.23636856	4.1976634	-.56309554E-01	-18.297655	17.824918
CIGTOT	.57370250E-03	.19689183E-02	.29137954	-.78979621E-02	.90453671E-02
CONSTANT	166.42380	598.72145	.27796532	-2409.6950	2742.5426

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AVERAGED DATA

VALUES NUNAME (CREATION DATE = 02/06/77)

***** MULTIPLE REGRESSION *****

DEPENDENT VARIABLE.. V25ARIMN

MEAN RESPONSE 2.12017 STD. DEV. 1.15762

VARIABLE(S) ENTERED ON STEP NUMBER 1.. HEIGHT
WEIGHT
AGE
CIGTOT

		ANALYSIS OF VARIANCE	DF	SUM OF SQUARES	MEAN SQUARE	F SIGNIFICANCE
MULTIPLE R	.35609	REGRESSION	4.	1,52931	.38233	.18152 .938
R SQUARE	.12680	RESIDUAL	5.	10,53154	2,10631	
STD DEVIATION	1.45131					

COEFFICIENT OF VARIABILITY 68.453 PERCENT

----- VARIABLES IN THE EQUATION -----

VARIABLE	B	STD ERROR B	F		BETA ELASTICITY
			SIGNIFICANCE	P	
HEIGHT	.75906112	11.261185	.45414457E-02	.949	.0308117
HEIGHT	-.40164150E-01	.16999061	.55824816E-01	.823	.64528
AGE	-.19209798E-01	.55347158E-01	.12046336	.743	-.1562395
CIGTOT	-.10486297E-05	.68228376E-05	.23621869E-01	.884	-.141051
(CONSTANT)	4.7450140	23.578165	.40499997E-01	.848	-.1698183

ALL VARIABLES ARE IN THE EQUATION.

COEFFICIENTS AND CONFIDENCE INTERVALS.

VARIABLE	B	STD ERROR B	T	95.0 PCT CONFIDENCE INTERVAL	
				L	U
HEIGHT	.75906112	11.261185	.67405087E-01	-.28.188295	.29.706417
HEIGHT	-.40164150E-01	.16999061	-.23627276	-.47713228	.19680398
AGE	-.19209798E-01	.55347158E-01	.34707832	-.16148203	.17306244
CIGTOT	-.10486297E-05	.68228376E-05	.15369408	-.18587025E-04	.16489766E-04
CONSTANT	4.7450140	23.578165	.20124611	-.55.863668	.65.353696

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AVERAGED DATA

FILE: KINNAME (CREATION DATE = 02/06/77)

MULTIPLE REGRESSION

DEPENDENT VARIABLE.. VADARINN

MEAN RESPONSE 3.86174 STD. DEV. 1.62798

VARIABLE(S) ENTERED ON STEP NUMBER 1..

HEIGHT

WEIGHT

AGE

CIGTOT

ANALYSIS OF VARIANCE

DF 4.

SUM OF SQUARES 5.32921

18.52063

ANALYSIS OF VARIANCE

DF 5.

SUM OF SQUARES .

.

MEAN SQUARE

1.31210

3.70413

F SIGNIFICANCE

.35968

.828

VARIABLES NOT IN THE EQUATION

F

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SIGNIFICANCE

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VARIABLES IN THE EQUATION

F

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SIGNIFICANCE

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B

.

.

STD ERROR B

.

.

BETA

.

.

ELASTICITY

.

.

HEIGHT 5.5931341 14.933651 .13977126

.774

.1611620

WEIGHT .26909489E-01 .22542748 .17663711E-01

.900

.0827145

AGE -.46971423E-01 .73396822E-01 .404955565

.550

.2952852

CIGTOT -.38333715E-05 .90478827E-05 .18043916

.689

.553764

(CONSTANT) -5.8263399 31.267412 .34722448E-01

.860

.2510561

.

.

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AII. VARIABLES ARE IN THE EQUATION.

COEFFICIENTS AND CONFIDENCE INTERVALS.

VARIABLE	B	STD ERROR B	T	95.0 PCT CONFIDENCE INTERVAL
HEIGHT	5.5931341	14.933651	.37366263	.32.804455 .43.970723
WEIGHT	.26909489E-01	.22542748	.13267898	-.54956150 .60318048
AGE	-.46971423E-01	.73396822E-01	-.43966535	-.25566109 .14169425
CIGTOT	-.38333715E-05	.90478827E-05	-.02478131	-.2101341E-04 .19014598E-04
CONSTANT	-5.8263399	31.267412	-.18633905	-.86.200560 .74.547880

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AVERAGED DATA

MALES
FILE NONAME (CREATION DATE = 02/06/77)

***** MULTIPLE REGRESSION *****

DEPENDENT VARIABLE.. CVINN

MEAN RESPONSE 1.66434 STD. DEV. .39077

VARIABLE(S) ENTERED ON STEP NUMBER 1.. HEIGHT
HEIGHT
AGE
CIGTOT

MULTIPLE R	.64461	ANALYSIS OF VARIANCE	DF	SUM OF SQUARES	MEAN SQUARE	F	SIGNIFICANCE
R SQUARE	.41552	REGRESSION	4.	.57104	.14276	.88864	.532
STD DEVIATION	.40081	RESIDUAL	5.	.80324	.16065		

COEFFICIENT OF VARIABILITY 24.082 PERCENT

----- VARIABLES IN THE EQUATION -----

VARIABLE	B	STD ERROR B	F	BETA	-----	
					SIGNIFICANCE	ELASTICITY
HEIGHT	3.3008623	3.1100111	1.1264994	.3969324		
WEIGHT	-.45233991E-01	.46946454E-01	.92437671	3.57462		
AGE	.12622657E-01	.15285273E-01	.68195416	-.5212764		
CIGTOT	.30391493E-05	.18842690E-05	2.6014667	.3305700		
(CONSTANT)	-1.8470221	6.5116023	.80457859E-01	.447	.34797	
				.168	.21080	
				.788		

----- VARIABLES NOT IN THE EQUATION -----

VARIABLE	PARTIAL	TOLERANCE	F
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SIGNIFICANCE

ALL VARIABLES ARE IN THE EQUATION.

COEFFICIENTS AND CONFIDENCE INTERVALS.

VARIABLE	B	STD ERROR B	T	95.0 PCT CONFIDENCE INTERVAL		
				-4.6935542	11.295279	
HEIGHT	3.3008623	3.1100111	1.0613667	-.16591186	.75443876E+01	
WEIGHT	-.45233991E-01	.46946454E-01	-.96352307	-.26668790E-01	.51914104E+01	
AGE	.12622657E-01	.15285273E-01	.82580516			
CIGTOT	.30391493E-05	.18842690E-05	1.6129063	.18044447E-05	.78827432E+05	
CONSTANT	-1.8470221	6.5116023	-.28365095	-18.5H5374	14.8H1330	

AVERAGED DATA							
FILE NO NAME (CREATION DATE = 02/06/77) MULTIPLE REGRESSION							
DEPENDENT VARIABLE.. SVCLMN							
MEAN RESPONSE	9.46630	STD. DEV.	1.89270				
VARIABLE(S) ENTERED ON STEP NUMBER	1..	HEIGHT					
		WEIGHT					
		AGE					
		CIGTOT					
multiple R	*.67877	ANALYSIS OF VARIANCE	DF	SUM OF SQUARES	MEAN SQUARE	F	SIGNIFICANCE
R SQUARE	*.46013	REGRESSION	4..	14.85432	3.71358	1.06194	*.460
STD DEVIATION	1.86476	RESIDUAL	5..	17.38660	3.41732		
COEFFICIENT OF VARIABILITY	19.609 PERCENT						
VARIABLES NOT IN THE EQUATION							
VARIABLE	B	STD ERROR B	F	BETA	VARIABLE	PARTIAL	TOLERANCE
HEIGHT	*.62649571	14.469230	*.18747586E-02	*.01555540			
WEIGHT	*.91210422E-01	.21841697	*.17431871	*.2170111			
AGE	*-.70529961E-01	.71114256E-01	*.98366098	*.381142			
CIGTOT	*.11152130E-04	.87665288E-05	1.705530	*.643041			
(CONSTANT)	8.3621991	30.295027	*.76190059E-01	*.13966			
			*.794				

ALL VARIABLES ARE IN THE EQUATION.

COEFFICIENTS AND CONFIDENCE INTERVALS.

VARIABLE	B	STD ERROR B	t	95.0 PCT CONFIDENCE INTERVAL
HEIGHT	*.62649571	14.469230	*.18747586E-01	*37.820270
WEIGHT	*.91210422E-01	.21841692	*.17431870	*.65766045
AGE	*-.70529961E-01	.71114256E-01	*.98366096	*.17227317
CIGTOT	*.11152130E-04	.87665288E-05	1.705530	*.31986800E-04
CONSTANT	8.3621991	30.295027	*.27602547	*.11082541E-04

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AVERAGED DATA

VALUES NINAMF CREATION DATE = 02/06/77

FILE NINAMF MULTIPLE REGRESSION

DEPENDENT VARIABLE = FEVIAN

MEAN RESPONSE 8.94727 STD. DEV. 1.44012

VARIABLE(S) ENTERED ON STEP NUMBER 1.

HEIGHT WEIGHT

AGE CIGTOT

ANALYSIS OF VARIANCE

REGRESSION DF 4

RESIDUAL DF 5

MEAN SQUARE 12.96100

5.10352

SIGNIFICANCE 2.84077 .141

R SQUARF 3.24047

1.14070

STD DEVIATION

COEFFICIENT OF VARIABILITY 11.937 PERCENT

VARIABLES IN THE EQUATION

R STD ERROR B F

SIGNIFICANCE

BETA

ELASTICITY

HEIGHT -.6.7588861 8.2872334 .66515978

WEIGHT .12093214 .12099802 .93650706

AGE *.89213152E-01 .40730601E-01 4.7975342

CIGTOT *-.10684466E-04 .50210035E-05 4.52881895

(CONSTANT) 17.451589 17.351619 1.0115770

.361

.452

.378

.080

.087

.087

.361

.2205367

.1.36153

.3781500

1.00638

.6339598

.45744

.7889241

.13786

ALL VARIABLES ARE IN THE EQUATION.

COEFFICIENTS AND CONFIDENCE INTERVALS.

VARIABLE	R	STD ERROR B	T	95.0 PCT CONFIDENCE INTERVAL
HEIGHT	-.6.7588861	8.2872334	-.8155732	.28.061534 14.543842
WEIGHT	.12093214	.12099802	.96669905	-.20063767 .44750194
AGE	-.89213152E-01	.40730601E-01	-2.1493274	-.15391110 .15886399E-01
CIGTOT	-.10684466E-04	.50210035E-05	-7.1279541	.23591170E-04 .2222238728E-05
CONSTANT	17.451589	17.351619	1.0057718	.27.151028 .02.054205

UNPACKED DATA		(CREATION DATE = 02/06/77)		MULTIPLE REGRESSION	
FILE	NAME	FILE	NAME	FILE	NAME
INDEPENDENT VARIABLE...					
MEAN RESPONSE					
11.57519	FVCMM	STD. DEV.	2.45017		
VARIABLE(S) ENTERED ON STEP NUMBER 1..					
HEIGHT		WEIGHT			
AGE		AGE			
C1G1OT					
ANALYSIS OF VARIANCE					
MULTIPLE R	.89456	ANALYSIS OF VARIANCE	DF	SUM OF SQUARES	MEAN SQUARE
N SQUARE	*.80023	REGRESSION	4,	43.73639	10.89010
STD. DEVIATION	1.46925	RESIDUAL	5,	10.79337	2.15869
COEFFICIENT OF VARIABILITY					
12.693 PERCENT					
VARIABLES IN THE EQUATION					
VARIABLE	B	STD ERROR B	F	BETA	
					SIGNIFICANCE
HEIGHT	*11.013624	11.400363	*.93669679	*.711662	
HEIGHT	*38510358	*17209154	5.0126768	*.378	*1.71805
AGE	*.59967211E-01	*.56031199E-01	1.5595986	*.075	*.7081534
C1G1OT	*.28815614E-04	*.69071617E-05	17.404298	*.267	*.41847
(CONSTANT)	9.3094474	23.869570	*.15212337	*.027733	
				*1.3505620	
				*.28139	
VARIABLES NOT IN THE EQUATION					
VARIABLE	B	STD ERROR B	F	PARTIAL	TOLERANCE
HEIGHT					
HEIGHT					
AGE					
C1G1OT					
(CONSTANT)					
ALL VARIABLES ARE IN THE EQUATION.					
COEFFICIENTS AND CONFIDENCE INTERVALS.					
VARIABLE	B	STD ERROR B	t		
HEIGHT	*11.013624	11.400363	*.93783097	*0.338744	18.211416
HEIGHT	*.69967211E-01	*.56031199E-01	*.2388955	*-.57065081E-01	*.8276724
AGE	*.46570769E-04	*.69071617E-05	*.1778188	*-2139780	*.74061392E-01
AGE	*.46570769E-04	*.69071617E-05	*.1778188	*-46570769E-04	*.11064046E-04
C1G1OT	*.28815614E-04	*.69071617E-05	*.1778188	*.047903	*.70.667598
(CONSTANT)	23.869570	*.15212337	*.113		*.28139
95.0 PCT CONFIDENCE INTERVAL					
VARIABLE	B	STD ERROR B	t		
HEIGHT					
HEIGHT					
AGE					
AGE					
C1G1OT					
(CONSTANT)					

EFFICIENTS AND CONFIDENCE INTERVALS

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AVGAGED DATA								
FILE	NAME	(CREATION DATE = 02/06/77)	MULTIPLE REGRESSION					
DEPENDENT VARIABLE...								
Y25F1NN								
MEAN RESPONSE	.28017	STD. DEV.	.10064					
VARIABLE(S) ENTERED ON STEP NUMBER	1..	HEIGHT WEIGHT AGE CIGTOT						
Coefficient of Variability	46.006 PERCENT							
VARIABLES IN THE EQUATION								
VARIABLE	B	STD. ERROR B	F	REGRESSION	R			
HEIGHT	-.15907809	1.9737071	.720804720E-02	-.0113817				
WEIGHT	-.15558059E-03	.2628113E-01	.30256387E-04	-.102119				
AGE	-.14226557E-02	.94090103E-02	.23865673E-01	-.0018745				
CIGTOT	-.37812265E-07	.11352269E-05	.11094104E-02	-.022587				
(CONSTANT)	.64870995	3.0230043	.27342794E-01	-.015555				
VARIABLES NOT IN THE EQUATION								
VARIABLE	REGRESSION	P	TOLERANCE	F	SIGNIFICANCE			
HEIGHT	-.15907809	.936	.102119					
WEIGHT	-.15558059E-03	.996	-.0018745					
AGE	-.14226557E-02	.983	-.022587					
CIGTOT	-.37812265E-07	.975	-.015555					
(CONSTANT)	.64870995	.875						

ALL VARIABLES ARE IN THE EQUATION.

COEFFICIENTS AND CONFIDENCE INTERVALS.					
VARIABLE	B	STD. ERROR B	T	95.0 PCT CONFIDENCE INTERVAL	
HEIGHT	-.15907809	1.8737071	-.84901048E-01	-4.9755224	4.6571662
WEIGHT	-.15558059E-03	.2628113E-01	-.55066351E-02	-.7246110E-01	-.7549442E-01
AGE	-.14226557E-02	.94090103E-02	-.15448119	-.2504810E-01	-.2244849E-01
CIGTOT	-.37812265E-07	.11352269E-05	-.13308113E-01	-.2055615E-05	-.2803170E-05
CONSTANT	.64870995	3.0230043	.16535713	.493357460	10.731166

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AVERAGED DATA
MALES FILE NUNAME (CREATION DATE # 02/06/77)

***** MULTIPLE REGRESSION *****

DEPENDENT VARIABLE.. VAGFRMN

MEAN RESPONSE .40321 STD. DEV. .21654

VARIABLE(S) ENTERED ON STEP NUMBER 1..

HEIGHT
WEIGHT
AGE
CIGTOTMULTIPLF R .42763
R SQUARF .18286
STD DEVIATION .26761
COEFFICIENT OF VARIABILITY 65.130 PERCENTANALYSIS OF VARIANCE DF
REGRESSION 4.
RESIDUAL 5.SUM OF SQUARS
.07717
.34462MEAN SQUARE
.01929
.06896

VARIABLES IN THE EQUATION

VARIABLE	B	STD FRMR B	F	NEFA	ELASTICITY
WEIGHT	*.45935734	2.0176768	*.50810541E+01	*.0096485	
HEIGHT	*.30534080E+01	.30759279E+01	*.95541097	*.650011	
AGE	*.13590384E+02	.10014899E+01	*.18414934E+01	*.642291	*.53852
CIGTOT	*.99845555E+06	.12345715E+05	*.65407113	*.15465	
(CONSTANT)	3.3270169	4.2663966	*.60011758	*.28587	
			.471		

ALL VARIABLES ARE IN THE EQUATION.

COEFFICIENTS AND CONFIDENCE INTERVALS.

VARIABLE	B	STD FRMR B	T	95.0 PCT CONFIDENCE INTERVAL
HEIGHT	*.45935734	2.0176768	*.22543190	*.5697296
WEIGHT	*.30534080E+01	.30759279E+01	*.99267868	*.10960217
AGE	*.13590381E+02	.10014898E+01	*.13520164	*.2484666E+01
CIGTOT	*.99845555E+06	.12345715E+05	*.80876664	*.21150013E+05
CONSTANT	3.3270169	4.2663966	*.77981894	*.14.291972

AVERAGED DATA
VALIDS 50 NAME (CREATION DATE = 02/06/77)

FILE 50 NAME ***** MULTIPLE REGRESSION

DEPENDENT VARIABLE.. CIGTOT

MEAN RESPONSE 19.27074 STD. DEV. 4.91114

VARIABLE(S) ENTERED ON STEP NUMBER 1.. HEIGHT
WEIGHT
AGE

Coefficient of Variability 21.720 PERCENT

ANALYSIS OF VARIANCE

DF SUM OF SQUARES
REGRESSION 4. 138.13382
RESIDUAL 5. 78.73953

F SIGNIFICANCE

2.19607 *205

VARIABLES IN THE EQUATION

VARIABLE	B	STD ERROR B	F	META
				ELASTICITY
HEIGHT	42.423424	30.791761	1.8982019	.4059099
WEIGHT	*.35394897	.46440991	*.57986389	*.18499
AGE	*.75456649	*.15133723	*.8295080	*.3745478
CIGTOT	*.3732448E-04	*.18655869E-04	*.0027133	*.6344213
(CONSTANT)	*47.876589	64.470413	*.55037350	*.5045522
				*.6392779
				*.8081496
				*.23583

ALL VARIABLES ARE IN THE EQUATION.

COEFFICIENTS AND CONFIDENCE INTERVALS.

VARIABLE	B	STD ERROR B	T	95.0 PCT CONFIDENCE INTERVAL
HEIGHT	42.423424	30.791761	1.3777525	*16.78113 121.57496
WEIGHT	*.35394897	.46440991	*.6149188	*.15487627 *8088478
AGE	*.75456649	*.15133723	*.6921142	*.13465232 *63358530
CIGTOT	*.3732448E-04	*.18655869E-04	*.0006832	*.0631225E-04 *85280192E-04
CONSTANT	*47.826589	64.470413	*.74193779	*117.889737 *213.55054

AVERAGED DATA

WLFSS
FILE : UNAME (CREATION DATE = 02/06/77)

MULTIPLE REGRESSION

DEPENDENT VARIABLE.. FEFVIM

MEAN RESPONSE 74.76136 STD. DEV. 9.84035

VARIABLE(S) ENTERED ON STEP NUMBER 1..

HEIGHT
WEIGHT
AGE
CIGTOT

MULTIPLE R
R SQUARE .996556
STD DEVIATION 9.94101
COEFFICIENT OF VARIABILITY 13.331 PERCENT

ANALYSIS OF VARIANCE DF SUM OF SQUARES
REGRESSION 4. 374.8091
RESIDUAL 5. 496.66132

F SIGNIFICANCE
.94338 *59

VARIABLES IN THE EQUATION

VARIABLE	B	STD ERROR B	F	BETA	ELASTICITY	SIGNIFICANCE
HEIGHT	21.811923	.77.333610	.79552047F+01	.1041572	.52585	
WEIGHT	1.0582776	1.1673716	.82182799	.4842940	.1.03398	
AGE	-.61806506	.38008395	2.6442915	-.647664	-.37931	
CIGTOT	-.33740092E-04	.46854277E-04	.51855407	-.3648992	-.02710	
(CONSTANT)	-11.095379	161.91766	.46961598E-02	.948		

ALL VARIABLES ARE IN THE EQUATION.

COEFFICIENTS AND CONFIDENCE INTERVALS.

VARIABLE	B	STD ERROR B	T	95.0 PCT CONFIDENCE INTERVAL
HEIGHT	21.811923	.77.333610	.28204971	.176.97743
WEIGHT	1.0582776	1.1673716	.90654729	-.1.9425011
AGE	-.61806506	.38008395	-.6261278	.4.0590563
CIGTOT	-.33740092E-04	.46854277E-04	-.72010698	-.3595689
CONSTANT	-11.095379	161.91766	-.68528531E-01	-.15418101E-01
				.86700831E-04
				.405.12029
				88

AVERAGED DATA
NAME=SNKERS
FILE= NO NAME (CREATION DATE = 02/06/77)

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DEPENDENT VARIABLE.. V2SAR1N

MEAN RESPONSE 2.23236 STD. DEV. 1.06927

VARIABLE(S) ENTERED ON STEP NUMBER 1.. HEIGHT

AGE

WEIGHT

MULTIPLE R .70152
R SQUARE .31573
STD DEVIATION 1.13551

ANALYSIS OF VARIANCE
REGRESSION 1.
RESIDUAL 17.

SUM OF SQUARES
1.91719
21.91941

F .24487
SIG. .864

COEFFICIENT OF VARIABILITY 50.866 PERCENT

VARIABLES IN THE EQUATION

VARIABLE	B	STD ERROR B	F	RETA	ELASTICITY	SIGNIFICANCE
WEIGHT	4.5788038	.5.5542159	*.67960412	.421	*.3734453	*.3.58181
AGE	*16440516E+01	*.41439110E+01	*.15834005	.697049	*.1897049	*.28888
*WEIGHT	*.56402431E+01	*.74069650E+01	*.57984924	*.696	*.5149522	
(CONSTANT)	-2.6561332	6.8915113	*.14854914	*.457	*1.68289	
				.705		

All VARIABLES ARE IN THE EQUATION.

COEFFICIENTS AND CONFIDENCE INTERVALS.

VARIABLE	B	STD ERROR B	T	95.0 PCT CONFIDENCE INTERVAL
WEIGHT	4.5788038	.5.5542159	*.67438348	*7.1304671
AGE	*16440516E+01	*.41439110E+01	*.19701966	*.7093978AE+01
WEIGHT	*.56402431E+01	*.74069650E+01	*.76117823	*.2126573
CONSTANT	-2.6561332	6.8915113	*.148549100	*17.195951

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AVERAGED DATA

NON-SMOKERS

FILE: NONAME (CREATION DATE = 02/06/77)

***** MULTIPLE REGRESSION *****

DEPENDENT VARIABLE.. V25HEINN

MEAN RESPONSE 2.62037 STD. DEV. 1.20154

VARIABLE(S) ENTERED ON STEP NUMBER 1.. HEIGHT
AGE
WEIGHT

MULTIPLE R	.20747	ANALYSIS OF VARIANCE	DF	SUM OF SQUARES	MEAN SQUARE	F SIGNIFICANCE
R SQUARE	.04304	REGRESSION	3.	1.24282	.41427	.25488 .857
STD DEVIATION	1.27489	RESIDUAL	17.	27.63091	1.62535	

COEFFICIENT OF VARIABILITY 48.653 PERCENT

VARIABLES IN THE EQUATION

VARIABLE	B	STD ERROR B	F	BETA	
				SIGNIFICANCE	ELASTICITY
HEIGHT	4.5318276	6.2359977	.52812275	.3289255	
			.477	3.02184	
AGE	.10010758E-01	.46525999E-01	.46295902E-01	.1025399	
			.832	.14941	
WEIGHT	-.58798092E-01	.83161723E-01	.49989582	.4777289	
			.489	-1.49459	
(CONSTANT)	-1.7730996	7.7374465	.52513496E+01		
			.821		

ALL VARIABLES ARE IN THE EQUATION.

COEFFICIENTS AND CONFIDENCE INTERVALS.

VARIABLE	B	STD ERROR B	T	95.0 PCT CONFIDENCE INTERVAL	
				L	U
HEIGHT	4.5318276	6.2359977	.72672055	-.8.6249773	.17.688633
AGE	.10010758E-01	.46525999E-01	.21516482	-.8815051RE-01	.10817203
WEIGHT	-.58798092E-01	.83161723E-01	.70703311	-.23425399	.11665781
CONSTANT	-1.7730996	7.7374465	.22915823	-.18.097685	.14.551486

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AVERAGED DATA
NAME=SMIFRS (CREATION DATE = 02/06/77)
FILE NAME

***** MULTIPLE REGRESSION *****

DEPENDENT VARIABLE= V40ARMIN

MEAN RESPONSE 3.71296 STD. DEV. 1.44135

VARIABLE(S) ENTERED ON STEP NUMBER 1..

HEIGHT
AGE
WEIGHT

MULTIPLE R .30233 ANALYSIS OF VARIANCE DF SUM OF SQUARES F SIGNIFICANCE
R SQUARE .09161 REGRESSION 3. 1.26596 .57008 .642
STD DEVIATION 1.49020 RESIDUAL 17. 37.75174 2.22069

Coefficient of Variability 39.920 PERCENT

VARIABLES IN THE EQUATION

VARIABLE	B	STD ERROR B	F	NETA	P	SIGNIFICANCE
HEIGHT	7.5825713	7.2891494	1.0821297	.4587843		
AGE	-.10445164E-01	.54383432E-01	.313	.354914		
WEIGHT	-.25163966E-01	.97206294E-01	.65014597E-01	-.0891885		
(CONSTANT)	-7.4312530	9.0441669	.67512932	-.10943		
			.423	-.44900		

ALL VARIABLES ARE IN THE EQUATION.

COEFFICIENTS AND CONFIDENCE INTERVALS.

VARIABLE	B	STD ERROR B	T	95.0 PCT CONFIDENCE INTERVAL
HEIGHT	7.5825713	7.2891494	1.0402546	-7.7961895 .22 .961337
AGE	-.10445164E-01	.54383432E-01	-.19206519	-.1251818 .10 .29385
WEIGHT	-.25163966E-01	.97206294E-01	-.25487178	-.23025132 .1749239
CONSTANT	-7.4312530	9.0441669	.82166253	+26.512777 11.650271

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AVERAGED DATA

NON-SMOKERS
FILE SONAME (CREATION DATE = 02/06/77)

***** MULTIPLE REGRESSION *****

DEPENDENT VARIABLE.. V40HEIMN

MEAN RESPONSE 4.74314 STD. DEV. 1.73937

VARIABLE(S) ENTERED ON STEP NUMBER 1.. HEIGHT
AGE
WEIGHT

		ANALYSIS OF VARIANCE	DF	SUM OF SQUARES	MEAN SQUARE	F	SIGNIFICANCE
MULTIPLE R	.39171	REGRESSION	3.	9.28398	3.09466	1.02704	.405
R SQUARED	.15343	RESIDUAL	17.	51.22406	3.01318		
STD DEVIATION	1.73585						
COEFFICIENT OF VARIABILITY	36.597 PERCENT						

----- VARIABLES IN THE EQUATION -----

VARIABLE	B	STD ERROR B	F	BETA	PARTIAL	TOLERANCE	F
				SIGNIFICANCE			
HEIGHT	12.196997	8.4907360	2.0635508	.6115360			
AGE	-.23435467E-01	.63348320E-01	.13686015	.4.49312			
WEIGHT	-.43483698E-01	.11323036	.14747799	-.1658229			
(CONSTANT)	-12.755473	10.535061	1.4659495	-.19323			
				-.2440561			
				-.61064			

ALL VARIABLES ARE IN THE EQUATION.

COEFFICIENTS AND CONFIDENCE INTERVALS.

VARIABLE	B	STD ERROR B	T	95.0 PCT CONFIDENCE INTERVAL	
HEIGHT	12.196997	8.4907360	1.4365064	-5.7168900	.30.110884
AGE	-.23435467E-01	.63348320E-01	-.36994614	-.15708874	.11021780
WEIGHT	-.43483698E-01	.11323036	-.38402863	-.28237887	.19541147
CONSTANT	-12.755473	10.535061	1.2107640	-.34.982508	.9.4715610

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AVERAGED DATA
NAME=SONERS
FILE=SONARE (CREATION DATE = 02/06/77)

***** MULTIPLE REGRESSION *****

INDEPENDENT VARIABLE.. CWINN

MEAN RESPONSE 1.4050 STD. DEV. .56942

VARIABLE(S) ENTERED ON STEP NUMBER 1.. HEIGHT

AGE
WEIGHT

MULTIPLE R .66026
R SQUARE .74007
STD DEVIATION .31488
COEFFICIENT OF VARIABILITY 22.372 PERCENT

ANALYSIS OF VARIANCE DF SUM OF SQUARES MEAN SQUARE F SIGNIFICANCE
REGRESSION 3. 4.79927 1.59976 .00000
RESIDUAL 17. 1.66559 .09915

VARIABLES IN THE EQUATION

VARIABLE	B	STD ERROR B	F	HTA	VARIABLE	PARTIAL	TOLERANCE	F	SIGNIFICANCE
HEIGHT	1.5778460	1.5407244	1.0194487	.2416519	HEIGHT				
AGE	-.11307635E-01	.11491422E-01	.96826899	.1.95874	AGE				
HEIGHT	.49047797E-01	.20540052E-01	5.7021147	.2443988	HEIGHT				
(CONSTANT)	-4.1741359	1.9110661	4.2706561	.31420	(CONSTANT)				
				.808900					
				2.22169					
				.043					

ALL VARIABLES ARE IN THE EQUATION.

COEFFICIENTS AND CONFIDENCE INTERVALS.

VARIABLE	B	STD ERROR B	T	95.0 PCT CONFIDENCE INTERVAL
HEIGHT	1.5778460	1.5407244	1.0244260	+1.611744 , 4.827454
AGE	-.11307635E-01	.11491422E-01	.98400660	-.35552015E-01 , -1.2937145E-01
HEIGHT	.49047797E-01	.20540052E-01	2.1879101	.57120165E-02 , .9288318E-01
(CONSTANT)	-4.1741359	1.9110661	-2.1841923	-.8.2061310 , -1.14213889

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AVERAGED DATA
NON-SMOKERS

FILE: NONAME (CREATION DATE = 02/06/77)

***** MULTIPLE REGRESSION *****

DEPENDENT VARIABLE.. SVC1MN

MEAN RESPONSE 8.84962 STD. DEV. 2.39814

VARIABLE(S) ENTERED ON STEP NUMBER 1.. HEIGHT
AGE
WEIGHT

	MULTIPLE R .72203	ANALYSIS OF VARIANCE	DF	SUM OF SQUARES	MEAN SQUARE	F SIGNIFICANCE 6.17160 .005
R SQUARED	.52133	REGRESSION	3.	59.96376	19.98792	
STD DEVIATION	1.79964	RESIDUAL	17.	55.05776	3.23869	
COEFFICIENT OF VARIABILITY 20.336 PERCENT						

VARIABLES IN THE EQUATION-----

VARIABLE	B	STD ERROR B	F	BETA	
				SIGNIFICANCE	ELASTICITY
HEIGHT	11.568318	8.8027346	1.7270512	.4206843	
AGE	-.12009316	.65676102E-01	3.3436575	.206 -.6163198	
WEIGHT	.18180326	.11739109	2.1984630	.085 -.53073	
(CONSTANT)	-18.776076	10.922180	2.9552282	.140 .7400855	
				.104 1.36836	

ALL VARIABLES ARE IN THE EQUATION.

COEFFICIENTS AND CONFIDENCE INTERVALS.

VARIABLE	B	STD ERROR B	T	95.0 PCT CONFIDENCE INTERVAL	
				L	U
HEIGHT	11.568318	8.8027346	1.3141712	-.0038285	.30.140464
AGE	-.12009316	.65676102E-01	-.8285671	-.25865762	.18471305E-01
WEIGHT	.18180326	.11739109	1.5486972	-.65870297E-01	.42947681
CONSTANT	-18.776076	10.922180	-1.7190777	-.41.819861	.4.2677091

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AVERAGED DATA
NON-SMOKERS

FTLK NONAME (CREATION DATE # 02/06/77)

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***** MULTIPLE REGRESSION *****

DEPENDENT VARIABLE.. FEV1NN

MEAN RESPONSE 8.15104 STD. DEV. 1.75265

VARIABLE(S) ENTERED ON STEP NUMBER 1.. HEIGHT
AGE
WEIGHT

MULTIPLE R	.85394	ANALYSIS OF VARIANCE	DF	SUM OF SQUARES	MEAN SQUARE	F	SIGNIFICANCE
R SQUARED	.72922	REGRESSION	3.	44.79996	14.93332	15.26062	.000
STD DEVIATION	.98922	RESIDUAL	17.	16.63539	.97855		

COEFFICIENT OF VARIABILITY 12.133 PERCENT

----- VARIABLES IN THE EQUATION -----

VARIABLE	B	STD ERROR B	F		BETA	PARTIAL	TOLERANCE	F	SIGNIFICANCE
			SIGNIFICANCE	ELASTICITY					
HEIGHT	6.0566135	4.8386573	2.0080273	.3411742					
			.175	1.46944					
AGE	-.13964373	.36100617E-01	14.962827	-.9805950					
			.001	-.66985					
WEIGHT	.20896499	.64527138E-01	10.487266	1.1639495					
			.005	1.70716					
(CONSTANT)	-12.284579	6.0036668	4.1868501						
			.057						

ALL VARIABLES ARE IN THE EQUATION.

COEFFICIENTS AND CONFIDENCE INTERVALS.

VARIABLE	B	STD ERROR B	T	95.0 PCT CONFIDENCE INTERVAL	
				+	-
HEIGHT	6.0566135	4.8386573	1.4170488	-3.3520609	17.065288
AGE	-.13964373	.36100617E-01	3.8681813	-.21580938	-.63478088E-01
WEIGHT	.20896499	.64527138E-01	3.2384048	.72824633E-01	.34510535
CONSTANT	-12.284579	6.0036668	-2.0461794	-24.951209	.38205044

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AVERAGED DATA
NOMINATORS (CREATION DATE = 02/06/77)
FILE: NODNAME

• • • • • MULTIPLE REGRESSION • • • • •
• • • • •

DEPENDENT VARIABLE: FVC1NN

MEAN RESPONSE 10.65141 STD. DEV. 2.98623

VARIABLE(S) ENTERED ON STEP NUMBER 1..

HEIGHT
AGE
WEIGHT

	ANALYSIS OF VARIANCE	DF	SUM OF SQUARES	F	SIGNIFICANCE
REGRESSION	3.	98.17659	32.72553	6.93902	.003
RESIDUAL	17.	80.17471	4.71616		

Coefficient of Variability 20.177 PERCENT

• • • • • VARIABLES IN THE EQUATION • • • • •

VARIABLE	B	STD ERROR B	F	BETA	ELASTICITY
HEIGHT	.0577257	10.622509	*.22625452	.1475582	
AGE	*.13869757	*.79251209E-01	3.0626961	.640	*.87639
WEIGHT	*.31448413	*.1415915	*.9288257	*.098	*.5716209
(CONSTANT)	-13.693677	13.180103	1.0794500	.040	*.0280881
					1.96568

All variables are in the equation.

Coefficients and Confidence Intervals.

VARIABLE	B	STD ERROR B	T	95.0 PCT CONFIDENCE INTERVAL
HEIGHT	5.0527257	10.622509	*.47566219	*17.35808
AGE	*.13869757	*.79251209E-01	*1.7500562	*.30590122
WEIGHT	*.31448413	*.1415915	2.2200058	*.15609451E-01
CONSTANT	-13.693677	13.180103	*-1.0389658	*41.501264
				14.113910

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AVERAGED DATA
NON-SMOKERS

FILE NUNAME (CREATION DATE = 02/06/77)

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***** MULTIPLE REGRESSION *****

DEPENDENT VARIABLE.. V25FR1MN

MEAN RESPONSE .28127 STD. DEV. .15669

VARIABLE(S) ENTERED ON STEP NUMBER 1.. HEIGHT
AGE
WEIGHT

MULTIPLE R	.35078	ANALYSIS OF VARIANCE	DF	SUM OF SQUARES	MEAN SQUARE	F	SIGNIFICANCE
R SQUARE	.12304	REGRESSION	3.	.06042	.02014	.79509	.513
STD DEVIATION	.15916	RESIDUAL	17.	.43062	.02533		

COEFFICIENT OF VARIABILITY 56.185 PERCENT

----- VARIABLES IN THE EQUATION -----

VARIABLE	B	STD ERROR B	F	BETA	PARTIAL	TOLERANCE	F
				SIGNIFICANCE			
HEIGHT	-1.0088783	.77849194	1.6794587	-.5615098			
AGE	-.69005764E-02	.58082310E-02	1.4115068	-.5420080			
WEIGHT	.14807359E-01	.10381776E-01	2.0342854	.9225508			
(CONSTANT)	1.3296523	.96593040	1.8948921	.187			

----- VARIABLES NOT IN THE EQUATION -----

VARIABLE	PARTIAL	TOLERANCE	F	SIGNIFICANCE

ALL VARIABLES ARE IN THE EQUATION.

COEFFICIENTS AND CONFIDENCE INTERVALS.

VARIABLE	B	STD ERROR B	T	95.0 PCT CONFIDENCE INTERVAL	
				LOWER	UPPER
HEIGHT	-1.0088783	.77849194	-1.2959193	-2.6513527	.63359611
AGE	-.69005764E-02	.58082310E-02	-1.1880685	-.19154873E-01	.51537197E-02
WEIGHT	.14807359E-01	.10381776E-01	1.4262838	-.70962740E-02	.36710991E-01
CONSTANT	1.3296523	.96593040	1.3765508	-.70828274	.3.3675872

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AVERAGED DATA

NON-SMOKERS

FILE NODNAME (CREATION DATE = 02/06/77)

***** MULTIPLE REGRESSION *****

DEPENDENT VARIABLE.. V40FRINN

MEAN RESPONSE .36373 STD. DEV. .12766

VARIABLE(S) ENTERED ON STEP NUMBER 1.. HEIGHT
AGE
WEIGHT

MULTIPLE R	.14549	ANALYSIS OF VARIANCE	DF	SUM OF SQUARES	MEAN SQUARE	F SIGNIFICANCE
R SQUARED	.02117	REGRESSION	3.	.00690	.00230	.12254 .946
STD DEVIATION	.13699	RESIDUAL	17.	.31902	.01877	

COEFFICIENT OF VARIABILITY 37.663 PERCENT

----- VARIABLES IN THE EQUATION -----

VARIABLE	B	STD ERROR B	F		ELASTICITY
			SIGNIFICANCE	P	
HEIGHT	.64901343E-01	.67006363	.93815648E-02	.924	.0443380
AGE	-.11793446E-02	.49992609E-02	.55650603E-01	.816	-.1137011
WEIGHT	.24478897E-02	.89358029E-02	.75044094E-01	.787	.1872010
(CONSTANT)	.13340099	.83139567	.25745573E-01	.874	.44827

----- VARIABLES NOT IN THE EQUATION -----

VARIABLE	PARTIAL	TOLERANCE	F
----------	---------	-----------	---

SIGNIFICANCE

ALL VARIABLES ARE IN THE EQUATION.

COEFFICIENTS AND CONFIDENCE INTERVALS.

VARIABLE	B	STD ERROR B	T	95.0 PCT CONFIDENCE INTERVAL	
				LOWER	UPPER
HEIGHT	.64901343E-01	.67006363	.96858478E-01	-1.3488093	1.4788120
AGE	-.11793446E-02	.49992609E-02	-.23590380	-.11726863E-01	.93681739E-02
WEIGHT	.24478897E-02	.89358029E-02	.27394177	-.16405006E-01	.21300786E-01
CONSTANT	.13340099	.83139567	.16045427	-1.6206905	1.8874925

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AVERAGED DATA
NON-SMOKERS
FILE NONAME (CREATION DATE = 02/06/77)

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***** MULTIPLE REGRESSION *****

DEPENDENT VARIABLE.. CSVINN

MEAN RESPONSE 15.05085 STD. DEV. 4.95143

VARIABLE(S) ENTERED ON STEP NUMBER 1., HEIGHT
AGE
WEIGHT

MULTIPLE R	.80122	ANALYSIS OF VARIANCE	DF	SUM OF SQUARES	MEAN SQUARE	F SIGNIFICANCE
R SQUARED	.64195	REGRESSION	3.	314.76959	104.92320	10.15988 .000
STD DEVIATION	3.21360	RESIDUAL	17.	175.56262	10.32721	

COEFFICIENT OF VARIABILITY 20.264 PERCENT

----- VARIABLES IN THE EQUATION -----

VARIABLE	B	STD ERROR B	F	BETA	PARTIAL	TOLERANCE	F SIGNIFICANCE
				SIGNIFICANCE			
HEIGHT	-20.214787	15.718981	1.6538251	-.3560407			
			.216	-2.22720			
AGE	-.47546817E-01	.11727735	.16436700	-.1181826			
			.690	-.11725			
WEIGHT	.59723148	.20962444	8.1171124	1.1775172			
			.011	2.50838			
(CONSTANT)	13.259144	19.503659	.46216687				
			.506				

----- VARIABLES NOT IN THE EQUATION -----

VARIABLE	PARTIAL	TOLERANCE	F
			SIGNIFICANCE

ALL VARIABLES ARE IN THE EQUATION.

COEFFICIENTS AND CONFIDENCE INTERVALS.

VARIABLE	B	STD ERROR B	T	95.0 PCT CONFIDENCE INTERVAL
HEIGHT	-20.214787	15.718981	-1.2860113	-.53.378937 , 12.949363
AGE	-.47546817E-01	.11727735	-.40542200	-.29498039 , .19988676
WEIGHT	.59723148	.20962444	2.8490547	.15496258 , 1.0395004
CONSTANT	13.259144	19.503659	.67982856	-.27.889979 , 54.408267

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AVERAGED DATA

NON-SMOKERS
FILE NONAME (CREATION DATE = 02/06/77)

***** MULTIPLE REGRESSION *****

DEPENDENT VARIABLE.. FEEVNN

MEAN RESPONSE 77.43870 STD. DEV. 10.60737

VARIABLE(S) ENTERED ON STEP NUMBER 1.. HEIGHT
AGE
WEIGHT

	MULTIPLE R	R SQUARED	STD. DEVIATION	ANALYSIS OF VARIANCE	DF	SUM OF SQUARES	MEAN SQUARE	F SIGNIFICANCE
MULTIPLE R	.70299			REGRESSION	3.	1112.11190	370.70397	5.53672 .008
R SQUARED	.49420			RESIDUAL	17.	1138.21270	66.95369	
STD. DEVIATION	8.18252							
COEFFICIENT OF VARIABILITY	10.566 PERCENT							

----- VARIABLES IN THE EQUATION -----

VARIABLE	B	STD. ERROR B	F		BETA ELASTICITY
			SIGNIFICANCE	P	
HEIGHT	11.673990	40.023960	.85074328E-01	.774	.0959782 .26340
AGE	-.56195244	.29861376	3.5414367	.077	-.6520102 .28380
WEIGHT	.13919406	.53374964	.68008971E-01	.797	-.1281055 .11972
(CONSTANT)	88.289837	49.660579	3.1608062	.093	

ALL VARIABLES ARE IN THE EQUATION.

COEFFICIENTS AND CONFIDENCE INTERVALS.

VARIABLE	B	STD. ERROR B	T	95.0 PCT CONFIDENCE INTERVAL	
				L	U
HEIGHT	11.673990	40.023960	.29167504	-.72.769184	.98.117165
AGE	-.56195244	.29861376	-.1.8818705	-.1.1919724	.68067527E-01
WEIGHT	.13919406	.53374964	-.26078530	-.1.2653074	.98691924
CONSTANT	88.289837	49.660579	1.7778656	-.16.484825	.193.06450

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AVERAGED DATA					
NON-SHIPPERS	NONNAME	CREATION DATE =	02/06/77	MULTIPLE REGRESSION	
FILE	NONNAME				
DEPENDENT VARIABLE: IVPPHN					
MEAN RESPONSE 19.32104 STD. DEV. 11.73047					
VARIABLE(S) ENTERED ON STEP NUMBER 1..					
HEIGHT WEIGHT					
ANALYSIS OF VARIANCE					
REGRESSION DF 3..					
RESIDUAL DF 17..					
SUM OF SQUARES					
115.83771 F .74900 SIGNIFICANCE .861					
2636.73431 155.017290					
COEFFICIENT OF VARIABILITY 64.452 PERCENT					
VARIABLES NOT IN THE EQUATION					
VARIABLE B STD ERROR B F SIGNIFICANCE					
HEIGHT -.39.577661 60.911706 .42217676 -.2942343					
AGE +15642631 +.45445462 +.18447822 +.1641183					
WEIGHT +15030953 +.91230346 +.14240211E+01 +.1250910					
(CONSTANT) 72.344261 75.577491 +.916266945 +.51818					
.352					
VARIABLE BETA ELASTICITY					
HEIGHT -.525 -.57916					
AGE +.735 +.31663					
WEIGHT +.855 +.1250910					
(CONSTANT) -.352 -.51818					
VARIABLE PARTIAL TOLERANCE F SIGNIFICANCE					
HEIGHT -.49475131 .60730913 .88.916005					
AGE +.4420666 +.80730913 +.1.152417					
WEIGHT +.95721965 +.1.5630009 +.1.6641200					
(CONSTANT) +.344261 +.87.116310 +.231.79883					

ALL VARIABLES ARE IN THE EQUATION.

COEFFICIENTS AND CONFIDENCE INTERVALS,					
variable	b	std error b	t	95.0 pct confidence interval	
HEIGHT	+39.577461	60.911706	.42217676	-.168.60993	.88.916005
AGE	+15642631	.45445462	.18447822	+.1641183	+.1.152417
WEIGHT	+15030953	.91230346	.14240211E+01	+.1250910	+.1.6641200
CONSTANT	72.344261	75.577493	.916266945	+.87.116310	+.231.79883

MULTIPLE REGRESSION									
DEPENDENT VARIABLE..		V22ARINN							
MEAN RESPONSE		1.88429		STD. DEV.		.88440			
VARIABLE(S) ENTERED ON STEP NUMBER		1..		WEIGHT					
MULTIPLE R		.56661		ANALYSIS OF VARIANCE		DF		SUM OF SQUARES	
R SQUARE		.32105		REGRESSION		3.		1.71187	
STD DEVIATION		.82213		RESIDUAL		11.		3.51561	
COEFFICIENT OF VARIABILITY		43.631 PERCENT						7.43479	
VARIABLES NOT IN THE EQUATION									
VARIABLE	R	STD ERROR R	F	SIGNIFICANCE	Y	VARIABLE	PARTIAL	TOLERANCE	F
WEIGHT	4.3606056	3.0764672	2.0080420	.4010128		WEIGHT			
AGE	-.454179607E-01	.22316543F=01	4.1457347	4.03607		AGE			
#EIGHT	*.20869231E=01	*.34228094E=01	.37100615	*.6694135		#EIGHT			
(CONSTANT)	-5.4814544	4.995098	1.2040635	*.9081077		(CONSTANT)			

ALL VARIABLES ARE IN THE EQUATION.

COEFFICIENTS AND CONFIDENCE INTERVALS.			
VARIABLE	R	STD ERROR R	T
WEIGHT	.3006056	3.0764672	1.4174068
AGE	-.454179607E-01	*.72316543E=01	-.2.0361077
#EIGHT	*.20869231E=01	*.34228094E=01	*.60917661
(CONSTANT)	-5.4814544	4.995098	*.0972981

95.0 PCT CONFIDENCE INTERVAL

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AVERAGED DATA					
SMOKEY'S	FILENAME	(CREATION DATE = 15/06/77)	MULTIPLE REGRESSION		
FILE					
DEPENDENT VARIABLE.. V75HEINN					
MEAN RESPONSE:	2.19150	STD. DEV.	1.18228		
VARIABLE(S) ENTERED ON STEP NUMBER	1...	HEIGHT			
WEIGHT		AGE			
MULTIPLE R	.54404	ANALYSIS OF VARIANCE	DF	SUM OF SQUARES	MEAN SQUARE
R SQUARE	.30014	REGRESSION	3.	5.87740	1.95913
STD DEVIATION	1.11566	RESIDUAL	11.	11.69155	1.24469
COEFFICIENT OF VARIABILITY	50.908 PERCENT				
VARIABLES NOT IN THE EQUATION					
VARIABLE	B	STD ERROR B	F	BETA	F
				SIGNIFICANCY	SIGNIFICANCE
HEIGHT	4.8798985	4.1748818	1.3662595	.3157016	.318154
AGE	-.56577339E+01	*.30311529E+01	3.4833924	-.6229661	
WEIGHT	*.70151574E+02	*.46489266E+01	.22770012E+01	-.06567	
(CONSTANT)	*4.6943919	6.7789591	.47954861	.72422	

ALL VARIABLES ARE IN THE EQUATION.

COEFFICIENTS AND CONFIDENCE INTERVALS.		
VARIABLE	B	STD ERROR B
HEIGHT	4.8798985	4.1748818
AGE	-.56577339E+01	*.30311529E+01
WEIGHT	*.70151574E+02	*.46489266E+01
CONSTANT	*4.6943919	6.7789591

95.0 PCT CONFIDENCE INTERVAL

AVERAGED DATA
SNOWPS FILE NODNAME (CREATION DATE = 15/06/77)

MULTIPLE REGRESSION
INDEPENDENT VARIABLE.. VADARINN
MEAN RESPONSE 3.53175 STD. DEV. 1.39373
VARIABLE(S) ENTERED ON STEP NUMBER 1.
MULTIPLE R .61932
R SQUARE .38555
STD. DEVIATION 1.23451
COEFFICIENT OF VARIABILITY 34.955 PERCENT

ANALYSIS OF VARIANCE
DF
REGRESSION 3.
RESIDUAL 11.

SUM OF SQUARES
10.43059
16.76415

MEAN SQUARE
3.47686
1.52401

F SIGNIFICANCE
2.28139 *136

VARIABLE	B	STD ERROR B	F	NETA	PARTIAL	TOLERANCE	F	SIGNIFICANCE
HEIGHT	9.4225878	4.6196555	4.1602449	.5408626				
AGE	*.68119421E-01	*.3540714E-01	*.1247467	*.6167594				
WEIGHT	.28850366E-01	*.51442206E-01	*.31453102	*.72145				
(CONSTANT)	-12.174512	7.5011435	2.7214640	*.57219				

ALL VARIABLES ARE IN THE EQUATION.

COEFFICIENTS AND CONFIDENCE INTERVALS.			95.0 PCT CONFIDENCE INTERVAL		
VARIABLE	B	STD ERROR B	T		
HEIGHT	9.4225878	4.6196555	2.0396777	*.70518192	19.590357
AGE	*.68119421E-01	*.3540714E-01	*2.0309472	*.14194202	*.57031920E-02
WEIGHT	.28850366E-01	*.51442206E-01	*.36081065	*.84171148E-01	*.14207348
CONSTANT	-12.174512	7.5011435	*1.6496660	*28.884435	*.11153713

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APPACED DATA

SWATCHES FILE NAME (CREATION DATE = 15/06/77)

FILE NAME MULTIPLE REGRESSION

DEPENDENT VARIABLE.. V40HEIM

MEAN RESPONSE 4.43816 STD. DEV. 1.77352

VARIABLE(S) ENTERED ON STEP NUMBER 1.

HEIGHT
AGE

WEIGHT

COEFFICIENT OF VARIABILITY 36.145 PERCENT

			ANALYSIS OF VARIANCE	DF	SUM OF SQUARES	F	SIGNIFICANCE
MULTIPLE R	.59763		REGRESSION	1,	15.72782	2.03724	*.167
R SQUARED	.35717		RESIDUAL	11,	28.30724	2.57359	
STD DEVIATION	1.60618						

VARIABLES IN THE EQUATION

VARIABLE	B	STD ERROR B	F	HETA SIGNIFICANCE	ELASTICITY
HEIGHT	12.428753	6.0029766	4.2866875	*.5609736	4.88408
AGE	+.76399192E-01	.43584323E-01	3.0726771	-.5607835	-.64389
WEIGHT	+.16034850E-01	.66846331E-01	.57540638E-01	-.0773094	.25307
(CONSTANT)	-15.503651	9.7473255	2.5296630	-.815	
					.140

ALL VARIABLES ARE IN THE EQUATION.

COEFFICIENTS AND CONFIDENCE INTERVALS.

VARIABLE	B	STD ERROR B	T	95.0 PCT CONFIDENCE INTERVAL
HEIGHT	12.428753	6.0029766	2.0704317	*.78170720 -.25.641713
AGE	+.76399192E-01	.43584323E-01	1.7559053	-.17232761 *.1612941E-01
WEIGHT	+.16034850E-01	.66846331E-01	1.3986310	-.13109291 *.5900649
CONSTANT	-15.503651	9.7473255	-.5905543	.36.957366

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AVERAGED DATA
SHEETNRS. 1000000 (CREATION DATE = 15/06/77)

FILE. NONAME MULTIPLE REGRESSION
 DEPENDENT VARIABLE. CV1MN
 MEAN RESPONSE 1.11824 STD. DEV. .48349
 VARIABLE(S) ENTERED ON STEP NUMBER 1.
 HEIGHT
 AGE
 WEIGHT
 ANALYSIS OF VARIANCE DF SUM OF SQUARES
 REGRESSION 3. 1.41599
 RESIDUAL 11. 1.88674
 MEAN SQUARE .47200
 .16879
 F SIGNIFICANCE 2.70628 .090

COEFFICIENT OF VARIABILITY 31.166 PERCENT
 VARIABLES NOT IN THE EQUATION

VARIABLE	B	STD ERROR B	F	BETA	PARTIAL	TOLERANCE	F	SIGNIFICANCE
HEIGHT	1.0232064	1.5174226	.44293495	.1721213				
AGE	.17264135E-01	*11102381E-01	2.3920804	*1.35372				
WEIGHT	.76713100E-02	*17120017E-01	*20078577	*4648313				
(CONSTANT)	-1.6491983	2.4953880	.43653654	*1.156894				
				*.40162				

ALL VARIABLES ARE IN THE EQUATION.

COEFFICIENTS AND CONFIDENCE INTERVALS.

VARIABLE	B	STD ERROR B	T	95.0 PCY CONFIDENCE INTERVAL
HEIGHT	1.0232064	1.5174226	*.66551358	*2.1606375 .4.4070502
AGE	.17264135E-01	*11102381E-01	*1.54663352	*.41812370E-01
WEIGHT	.76713100E-02	*17120017E-01	*44800125	*.30009568E-01
CONSTANT	*1.6491983	2.4953880	*.66070912	*7.1438985 .3.8451258

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AVERAGED DATA
SNAPPS FILE: NOSNAME (CREATION DATE: 15/06/77)

MULTIPLE REGRESSION

DEPENDENT VARIABLE..

MEAN RESPONSE

VARIATE(S) ENTERED ON STEP NUMBER 1..

COEFFICIENT OF VARIABILITY

20.270 PERCENT

VARIABLES IN THE EQUATION

VARIABLE	B	STD ERROR B	F	SIGNIFICANCE
HEIGHT	12.512884	6.0921565	4.212189	*.5720372
AGE	*.3989777E-01	*.44231810E-01	.81363175	*.7214724
WEIGHT	*44453948E-01	*.67839397E-01	*.42339473	*.1858142
(CONSTANT)	-15.447700	9.8921314	7.4386412	*.38769

ALL VARIABLES ARE IN THE EQUATION.

Coefficients and Confidence Intervals.

VARIABLE	B	STD ERROR B	T	95.0 PCT CONFIDENCE INTERVAL
HEIGHT	12.532884	6.0921565	2.0572161	*.87586038
AGE	*.39897773E-01	*.4423180E-01	*.90201538	*.13725131
WEIGHT	*44453948E-01	*.67839397E-01	*.65528218	*.10885934
CONSTANT	-15.447700	9.8921314	*1.5616149	*37.220131

ALL VARIABLES ARE IN THE EQUATION.

VARIABLE	B	STD ERROR B	F	SIGNIFICANCE
HEIGHT	12.532884	6.0921565	2.0572161	*.87586038
AGE	*.39897773E-01	*.4423180E-01	*.90201538	*.13725131
WEIGHT	*44453948E-01	*.67839397E-01	*.65528218	*.10885934
CONSTANT	-15.447700	9.8921314	*1.5616149	*37.220131

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AVERAGED DATA

SMOKERS

FILE NODNAME (CREATION DATE = 15/06/77)

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***** MULTIPLE REGRESSION *****

DEPENDENT VARIABLE.. FEVINN

MEAN RESPONSE 7.74222 STD. DEV. 1.84799

VARIABLE(S) ENTERED ON STEP NUMBER 1.. HEIGHT
AGE
WEIGHT

MULTIPLE R	.62036	ANALYSIS OF VARIANCE	DF	SUM OF SQUARES	MEAN SQUARE	F	SIGNIFICANCE
R SQUARED	.38484	REGRESSION	3.	18.39968	6.13323	2.29387	.135
STD DEVIATION	1.63516	RESIDUAL	11.	29.41123	2.67375		

COEFFICIENT OF VARIABILITY 21.120 PERCENT

----- VARIABLES IN THE EQUATION -----

VARIABLE	B	STD ERROR B	F	BETA		VARIABLE	PARTIAL	TOLERANCE	F
				SIGNIFICANCE	ELASTICITY				
HEIGHT	14.354027	6.1189157	5.5029884	.6317376					
			.039	3.23345					
AGE	.24944236E-01	.44426094E-01	.31525459	.1757164					
			.586	.12051					
WEIGHT	.24036828E-01	.68137375E-01	.12444673	.1112193					
			.731	.21747					
(CONSTANT)	-18.042517	9.9355816	3.2976736						
			.097						

----- VARIABLES NOT IN THE EQUATION -----

VARIABLE	PARTIAL	TOLERANCE	F	-----	
				SIGNIFICANCE	

ALL VARIABLES ARE IN THE EQUATION.

COEFFICIENTS AND CONFIDENCE INTERVALS.

VARIABLE	B	STD ERROR B	T	95.0 PCT CONFIDENCE INTERVAL			
				L	U	STDEV	NORM
HEIGHT	14.354027	6.1189157	2.3458449	.88638680	.27.821668		
AGE	.24944236E-01	.44426094E-01	.56147715	-.12272539	.72836921E-01		
WEIGHT	.24036828E-01	.68137375E-01	.35277008	-.12593250	.17400616		
CONSTANT	-18.042517	9.9355816	-1.8159498	-.39.910581	.3.8255471		

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AVERAGED DATA

SMOKERS

FILE NONAME (CREATION DATE = 15/06/77)

***** MULTIPLE REGRESSION *****

DEPENDENT VARIABLE.. FVC1MN

MEAN RESPONSE 9.61950 STD. DEV. 2.10020

VARIABLE(S) ENTERED ON STEP NUMBER 1.. HEIGHT
AGE
WEIGHT

	MULTIPLE R	.60154	ANALYSIS OF VARIANCE	DF	SUM OF SQUARES	MEAN SQUARE	F	SIGNIFICANCE
R SQUARE	.36185		REGRESSION	3.	22.3442	7.44827	2.07910	.161
STD DEVIATION	1.89273		RESIDUAL	11.	39.40689	3.58244		

COEFFICIENT OF VARIABILITY 19.676 PERCENT

----- VARIABLES IN THE EQUATION -----

VARIABLE	B	STD ERROR B	F	BETA	
				SIGNIFICANCE	ELASTICITY
HEIGHT	15.315059	7.0827833	4.6755149	.5930901	
AGE	-.37208467E-01	.51424208E-01	.52353813	.277667	
WEIGHT	.44103181E-01	.78870553E-01	.31268717	.14468	
(CONSTANT)	-18.788137	11.500660	2.6688358	.12114	
			.131		

ALL VARIABLES ARE IN THE EQUATION.

COEFFICIENTS AND CONFIDENCE INTERVALS.

VARIABLE	B	STD ERROR B	T	95.0 PCT CONFIDENCE INTERVAL	
				L	U
HEIGHT	15.315059	7.0827833	2.1622939	-.27403933	.30.904158
AGE	-.37208467E-01	.51424208E-01	-.72355935	-.15039237	.75975415E-01
WEIGHT	.44103181E-01	.78870553E-01	.55918438	-.12948971	.21769607
CONSTANT	-18.788137	11.500660	-1.6336572	-.44.100916	.6.5246423

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AVERAGED DATA
SMOKERS

FILE NAME (CREATION DATE = 15/06/77)

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***** MULTIPLE REGRESSION *****
DEPENDENT VARIABLE.. V25FRINN

MEAN RESPONSE .31622 STD. DEV. .18581

VARIABLE(S) ENTERED ON STEP NUMBER 1.. HEIGHT
AGE
WEIGHT

MULTIPLE R	.27469	ANALYSIS OF VARIANCE	DF	SUM OF SQUARES	MEAN SQUARE	F	SIGNIFICANCE
R SQUARE	.07545	REGRESSION	3.	.03647	.01216	.29924	.825
STD DEVIATION	.20156	RESIDUAL	11.	.44687	.04062		

Coefficient of Variability 63.740 PERCENT

VARIABLES IN THE EQUATION				
VARIABLE	B	STD ERROR B	F	BETA
			----- SIGNIFICANCE	----- ELASTICITY
HEIGHT	.39547983	.75423718	.27493676 .610	.1731112 2.18122
AGE	-.18470099E-02	.54761029E-02	.11376148 .742	-.1294045 -.21848
WEIGHT	-.46208006E-02	.83988315E-02	.30268905 .593	-.2126644 -1.02356
(CONSTANT)	.19232596E-01	1.2246917	.24661660E-03 .988	

VARIABLES NOT IN THE EQUATION				
VARIABLE	PARTIAL	TOLERANCE	F	SIGNIFICANCE

ALL VARIABLES ARE IN THE EQUATION.

COEFFICIENTS AND CONFIDENCE INTERVALS.

VARIABLE	B	STD ERROR B	T	95.0 PCT CONFIDENCE INTERVAL
HEIGHT	.39547983	.75423718	.52434412	-.12645848 , 2.0555444
AGE	-.18470099E-02	.54761029E-02	.33728546	-.13899829E-01, .10205809E-01
WEIGHT	-.46208006E-02	.83988315E-02	-.55017184	-.23106501E-01, .13864900E-01
CONSTANT	.19232596E-01	1.2246917	.15704031E-01	-.2.6762952 , 2.7147604

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AVERAGED DATA SMOKERS FILE: NODNAME (CREATION DATE = 15/06/77)						
***** MULTIPLE REGRESSION *****						
DEPENDENT VARIABLE: V40FRMN						
MEAN RESPONSE	*.37500	STD. DEV.	.24196			
VARIABLE(S) ENTERED ON STEP NUMBER	1..	HEIGHT AGE WEIGHT				
MULTIPLE R	*.24381	ANALYSIS OF VARIANCE	DF	SUM OF SQUARES	MEAN SQUARE	F SIGNIFICANCE
R SQUARE	*.05944	REGRESSION	3,	*.04972	*.01624	*.23174 *.872
STD DEVIATION	*.26473	RESIDUAL	11,	*.77090	*.07008	
COEFFICIENT OF VARIABILITY	70.595 PERCENT					
***** VARIABLES NOT IN THE EQUATION *****						
VARIABLE	B	STD ERROR B	F	BETA	PARTIAL	TOLERANCE
				SIGNIFICANCE		
HEIGHT	*.54018610	*.99064410	*.29755918	*.1816050		
AGE	*.3008795E=02	*.71925239E=02	*.17499393	*.594	*.515121	
WEIGHT	*.2550486E=02	*.11031348E=01	*.51455144E=01	*.684	*.1618792	
(CONSTANT)	*.50135136	1.6085571	*.91143757E=01	*.821	*.0011236	
					*.47660	

ALL VARIABLES ARE IN THE EQUATION.

COEFFICIENTS AND CONFIDENCE INTERVALS.

VARIABLE	B	STD ERROR B	T	95.0 PCT CONFIDENCE INTERVAL
H HEIGHT	*.54018630	*.99064410	*.54548985	*-1.6400063 2.7207789
AGE	*.3008795E=02	*.71925239E=02	*.41932276	*.17821839E=01 *1.8839432E=01
WEIGHT	*.2550486E=02	*.11031348E=01	*.23120148	*.26630155E=01 *2.1129441E=01
CONSTANT	*.50135136	1.6085571	*.31167893	*-4.0417631 *3.0390564

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AVERAGED DATA

SMOKERS
FILE NODNAME (CREATION DATE = 15/06/77)

***** MULTIPLE REGRESSION *****

DEPENDENT VARIABLE.. CSVIMN

MEAN RESPONSE 16.49667 STD. DEV. 6.01552

VARIABLE(S) ENTERED ON STEP NUMBER 1.. HEIGHT
AGE
WEIGHT

		ANALYSIS OF VARIANCE	DF	SUM OF SQUARES	MEAN SQUARE	F	SIGNIFICANCE
MULTIPLE R	.63922	REGRESSION	3.	207.00313	69.00104	2.53335	.111
R SQUARE	.40860	RESIDUAL	11.	299.60762	27.23708		
STD DEVIATION	5.21891						

COEFFICIENT OF VARIABILITY 30.887 PERCENT

----- VARIABLES IN THE EQUATION -----

VARIABLE	B	STD ERROR B	F		BETA ELASTICITY
			SIGNIFICANCE	P	
HEIGHT	6.1550719	19.529642	.99329374E-01	.0832190	.63532
AGE	.31621902	.14179403	4.9734769	.6843157	.70003
WEIGHT	-.10157205	.21747293	.21814137	-.1443788	-.42107
(CONSTANT)	1.4484697	31.711231	.20863762E-02	.964	

----- VARIABLES NOT IN THE EQUATION -----

VARIABLE PARTIAL TOLERANCE F SIGNIFICANCE

ALL VARIABLES ARE IN THE EQUATION.

COEFFICIENTS AND CONFIDENCE INTERVALS.

VARIABLE	B	STD ERROR B	T	95.0 PCT CONFIDENCE INTERVAL	
				L	H
HEIGHT	6.1550719	19.529642	.31516563	-.16.829173	.49.139517
AGE	.31621902	.14179403	2.2301294	.41125211E-02	.42830552
WEIGHT	-.10157205	.21747293	-.46705607	-.58022667	.37708257
CONSTANT	1.4484697	31.711231	.45676867E+01	-.68.347468	.71.244408

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AVERAGED DATA
S'DFERS

FILE NNAME (CREATION DATE = 15/06/77)

***** MULTIPLE REGRESSION *****

DEPENDENT VARIABLE.. FEEVIMN

MEAN RESPONSE 79.59778 STD. DEV. 7.69802

VARIABLE(S) ENTERED ON STEP NUMBER 1.. HEIGHT
AGE
WEIGHT

MULTIPLE R .42475
R SQUARE .18042
STD DEVIATION 7.86218

COEFFICIENT OF VARIABILITY 9.877 PERCENT

ANALYSIS OF VARIANCE DF SUM OF SQUARES
REGRESSION 3. 149.67940
RESIDUAL 11. 679.95295

MEAN SQUARE 49.89313
61.81390
F SIGNIFICANCE .80715 .516

VARIABLES IN THE EQUATION -----

VARIABLE	B	STD ERROR B	F		BETA ELASTICITY
			SIGNIFICANCE	P	
HEIGHT	.26.128064	.29.420990	.78867833 .394		.2760520 .57249
AGE	-.12344103	.21360968	.33394673 .575		-.2087480 .05801
WEIGHT	-.28822179	.32761834	.77395767 .398		-.3201477 .25363
(CONSTANT)	58.835036	47.772295	1.5167704 .244		

VARIABLES NOT IN THE EQUATION -----

VARIABLE PARTIAL TOLERANCE F
SIGNIFICANCE

ALL VARIABLES ARE IN THE EQUATION.

COEFFICIENTS AND CONFIDENCE INTERVALS.

VARIABLE	B	STD ERROR B	T	95.0 PCT CONFIDENCE INTERVAL	
				L	U
HEIGHT	.26.128064	.29.420990	.88807564	-.38.627088	.90.881217
AGE	-.12344103	.21360968	-.57788124	-.59359270	.34671064
WEIGHT	-.28822179	.32761834	-.87974864	-.1.0093048	.41288121
CONSTANT	58.835036	47.772295	1.2315723	-.46.311060	.161.98113

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MULTIPLE REGRESSION						15/06/77	PAGE: 31
DEPENDENT VARIABLE.. IVYPMN		25.86263	STD. DEV.	19.30242			
INDEPENDENT VARIABLE(S) ENTERED ON STEP NUMBER		1..	HEIGHT AGE WEIGHT				
MULTIPLE R		.43375	ANALYSIS OF VARIANCE	DF	SUM OF SQUARES	F SIGNIFICANCE	
R SQUAPE		.18814	REGRESSION	1,	981.36405	*.64970 .495	
STD DEVIATION		19.62097	RESIDUAL	11.	4234.80617	327.12135	
COEFFICIENT OF VARIABILITY		75.866 PERCENT			384.98238		
VARIABLES IN THE EQUATION						VARIABLES NOT IN THE EQUATION	
VARIABLE	B	STD ERROR B	F	SIGNIFICANCE	BETA	VARIABLE	PARTIAL TOLERANCE F SIGNIFICANCE
HEIGHT	*109.36951	73.473422	7.2188263	*.4608365			
AGE	*533343904E-01	*53108723	*1001170235E-01	*7.37534			
WEIGHT	*79631253E-01	*81760878	*94958444E-02	*.037715			
(CONSTANT)	209.03516	119.22119	3.0741978	*.21567			
			*.107				

All variables are in the equation.

COEFFICIENTS AND CONFIDENCE INTERVALS.						95.0 PCT CONFIDENCE INTERVAL
VARIABLE	B	STD ERROR B	T			
HEIGHT	*109.36951	73.473422	*1.4805725	*270.97315		.52.218325
AGE	*533343904E-01	*53108723	*10006600	*-1.1199710		1.2266608
WEIGHT	*79631253E-01	*81760878	*97195300E-01	*-1.7199133		1.1791758
CONSTANT	209.03516	119.22119	1.7533390	*-53.368862		.471.45918

AVERAGED DATA
DRUG ONE FILE NODNAME (CREATION DATE = 02/06/77)

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GROUP COUNTS		GROUP 1	GROUP 2	TOTAL
NUMBER		12.	7.	19.

MFANS	GROUP 1	GROUP 2	TOTAL
V25AB1MN	2.208667	2.060000	2.203116
V25HE1MN	2.688667	2.550000	2.636132
V40AB1MN	4.01813	3.10143	3.301556
V40HE1MN	4.91667	4.72714	4.84684
V40HE1WJ	1.45750	1.15000	1.34421
COLUMN			30895
V25FP1AU	3.26667	2.71457	3.0832
V40FP1AU	3.20883	2.81143	3.15158
IVFP1MN	21.39570	26.70571	21.40474
V25B13MX	**.41667	**.38429	**.62168
V25B13MX	**.61500	**.61857	**.61847
V40B13MX	**.60417	**.61286	**.53621
V40B13MX	**.41750	**.70000	**.02268
CV3WJ	**.05583	**.01343	**.25189
V25F13MX	**.41133	**.00857	**.13895
V40F13MX	**.30250	**.14143	**.14286
IVFP1MX	**.73833	**.524286	1.69263

STANDARD DEVIATIONS

STANDARD DEVIATIONS	GROUP 1	GROUP 2	TOTAL
V25AB1MN	1.02671	.69414	.90411
V25HE1MN	.87636	1.02018	.91601
V40AB1MN	1.64245	1.07984	1.43594
V40HE1MN	1.61517	1.36751	1.49213
CV1MN	.60849	*.02190	.55572
V25C91MN	*.21216	*.18702	*.18797
V25D91MN	*.15900	*.09754	*.13785
V25E91MN	21.45575	27.52008	23.26222
IVFP1AU	.60262	.56515	.60473
V25A83MX	.60262	.56515	.60473
V25HE3MX	.62260	.70761	.63555
V60AB3MX	.87767	1.04607	.9125
V40HE3MX	2.00581	1.12259	1.70166
CV9WJ	.48135	.52694	.48634
V25FP3MX	1.28649	.46806	1.06245
V40FP3MX	1.31611	.52149	1.09434
IVFP3MX	29.73983	20.62607	26.68191

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AVERAGED DATA
DRUG ONE
FILE NAME (CREATION DATE = 02/06/77)

----- DISCRIMINANT ANALYSIS -----

ANALYSIS NUMBER	1	MAXIMUM STEPS
TOLERANCE LEVEL	.00010	32
F FOR INCLUSION	.01000	F FOR DELETION .00500
SOLUTION METHOD = STEPWISE. SELECT VARIABLE WHICH WILL MINIMIZE DISTANCE FUNCTION.		
PRIOR PROBABILITIES = EQUAL		
GROUP 1	GROUP 2	
.50000	.50000	
ALL ELIGIBLE VARIABLES INCLUDED		

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ANALYSIS NUMBER	V25AP1MN	V25AR1MN	V40HFMN	CV1MN	V25FR1MN	V40FR1MN	IVFP1MN	V25AR3MN
V25AP1MN	.45215	.86427	2.15709	2.34807	.30240	.02911	.01972	
V25AR1MN	.77216	1.00716	2.14419	2.0492	.01054	.01681	.01972	
V40AP1MN	1.18613	1.12172	1.12172	1.04680	.01480	.02300	.0245	.565.67770
V40HFMN	1.20720	1.15744	1.15744	1.04681	.01480	1.04685	.0245	
CV1MN	.09129	-.09129	-.09129	-.09129	-.09129	.93229	.93229	
V25FR1MN	-.09194	-.04661	-.04661	-.04661	-.04661	.476192	.476192	
IVFP1MN	-.06317	-.03975	-.03975	-.03975	-.03975	.48380	.48380	
V25AR3MN	.26465	8.67111	8.67111	8.67111	8.67111	.38088	.38088	
V25MF3MN	-.28250	-.26120	-.26120	-.26120	-.26120	.91870	.91870	
V40AR1MN	-.21588	-.20425	-.20425	-.20425	-.20425	.91870	.91870	
V40HFMN	-.37210	-.32356	-.32356	-.32356	-.32356	.1869	.1869	
V40AP1MN	-.95574	-.24431	-.24431	-.24431	-.24431	.1869	.1869	
V40HFMN	-.04553	-.01207	-.01207	-.01207	-.01207	.01432	.01432	
CV1MN	-.44510	-.34454	-.34454	-.34454	-.34454	.15057	.15057	
V25FR1MN	-.50058	-.36351	-.36351	-.36351	-.36351	.75559	.75559	
V40HFMN	-.50058	-.59330	-.59330	-.59330	-.59330	.05342	.05342	
IVFP1MN	-1.45546	-6.90449	10.01107	7.75678	5.0282	.014934	.014934	
V25FR3MN		V40FR1MN	V25FR1MN	V40FR1MN	V25FR3MN	V40FR3MN	V40FR3MN	IVFP1MN
V25MF3MN		V40HFMN	V40HFMN	V40HFMN	V25MF3MN	V25MF3MN	V25MF3MN	V25MF3MN
V25AP1MN		V25AR1MN						
V40AP1MN		V40HFMN						
CV1MN		V25FR1MN						
V25FR1MN		V25AR3MN						
V40HFMN		V40FR1MN						
IVFP1MN		V25MF3MN						

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AVERAGED DATA
DRUG ONE
FILE NAME (CREATION DATE = 02/06/77)

ANALYSIS NUMBER	1	MAXIMUM STEPS
TOLERANCE LEVEL	.00010	32
F FOR INCLUSION	.01000	F FOR DELETION .00500
SOLUTION METHOD = STEPWISE. SELECT VARIABLE WHICH WILL MINIMIZE DISTANCE FUNCTION.		
PRIOR PROBABILITIES = EQUAL		
GROUP 1	GROUP 2	
.50000	.50000	
ALL ELIGIBLE VARIABLES INCLUDED		

AVERAGED DATA
DIFC ONE:

***** VARIABLES IN THE ANALYSIS *****

VARIABLE	ENTER CRITERION	F TO REMOVE
V75AR1MN	*.90164	*.00029
V75AF1MN	*.64074	1.02741
V40AR1MN	-.65153	1.11107
V40HF1MN	-.78907	3.77631
C1MN	-.91750	1.79463
V75F1MN	-.71421	.04603
V10F1MN	-.61944	.86716
IVPP1MN	-.61290	.81644

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***** VARIABLES NOT IN THE ANALYSIS *****

VARIABLE	TOLERANCE	F TO ENTER	F TO ENTRY CRITERION
NINHFR REMOVED	.58074	7.60831	8
0	*.72194	*.64750	100.0
			.473

1 FUNCTIONS WILL BE USED IN REMAINING ANALYSES

STANDARDIZED DISCRIMINANT FUNCTION COEFFICIENTS

V75AR1MN	-.04680
V75AF1MN	-.2-.67796
V40AF1MN	-.90193
V40HF1MN	5.90678
C1MN	-.1.27735
V75F1MN	*.30338
V10F1MN	-.1.17596
IVPP1MN	*.67209

UNSTANDARDIZED DISCRIMINANT FUNCTION COEFFICIENTS

V75AR1MN	*.511250E-01
V75HF1MN	-.2-.95556
V40HF1MN	-.2-.84193
V40F1MN	3.95863
C1MN	-2.29854
V75F1MN	1.61934
V10F1MN	-.8.53081
IVPP1MN	*.288918E-01
CONSTANT	4.10867

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AVERAGED DATA
DRUG ONE

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CENTROIDS OF GROUPS IN REDUCED SPACE

GROUP 1 = .61384
GROUP 2 = 1.05230

PREDICTION RESULTS -

ACTUAL GROUP NAME	GROUP CODE	N OF CASES	PREDICTED GROUP MEMBERSHIP	
			GROU P 1	GROU P 2
GROUP 1	2	14	10. 52.6 PCT	4. 21.1 PCT
GROUP 2	3	11	2. 10.5 PCT	9. 47.4 PCT

100.0 PERCENT OF KNOWN CASES CORRECTLY CLASSIFIED

CHI-SQUARE = 19.000 SIGNIFICANCE = .000

AVERAGED DATA
DRUG ONE
FILE: NONAME (CREATION DATE = 02/06/77)

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- - - - - DISCRIMINANT ANALYSIS - - - - -

ANALYSIS NUMBER 2

TOLERANCE LEVEL .00010 MAXIMUM STEPS 32
F FOR INCLUSION .01000 F FOR DELETION .00500

SOLUTION METHOD = STEPWISE. SELECT VARIABLE WHICH WILL MINIMIZE DISTANCE FUNCTION.

PRIOR PROBABILITIES = EQUAL

GROUP 1 GROUP 2
.50000 .50000

ALL ELIGIBLE VARIABLES INCLUDED

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AVERAGED DATA
DRUG ONE

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VARIABLES IN THE ANALYSIS

VARIABLE	ENTRY CRITERION	F TO REMOVE
V25AP3W	" .75479	.11189
V25HF3W	" .72830	.08495
V40AP3W	" .75523	.09686
V40HF3W	" .80326	1.00664
V40SF3W	" .73217	.07584
CV3W	" .73211	.01095
V25P3W	" .90115	.55008
V40P3W	" .95494	1.06608
IVPP3W		

NUMBER REMOVED	EIGENVALUE	CANONICAL CORRELATION	PERCENT OF TRACE	WILKS LAMBDA	CHI-SQUARE	D.F.	SIGNIFICANCE
0	.39238	.51085	100.0	.71820	4.63417	8	.796

1 FUNCTIONS WILL BE USED IN REMAINING ANALYSES

STANDARDIZED DISCRIMINANT FUNCTION COEFFICIENTS

1

V25AP3W	" .73652
V25HF3W	" .11294
V40AP3W	" .60507
V40HF3W	1.46289
CV3W	" .11743
V25P3W	.36468
V40P3W	" 1.70405
IVPP3W	1.04375

UNSTANDARDIZED DISCRIMINANT FUNCTION COEFFICIENTS

V25AP3W	" 1.21795
V25HF3W	" .449460
V40AP3W	" .6161821
V40HF3W	" .959465
CV3W	" .262453
V25P3W	" .343245
V40P3W	" 1.55715
IVPP3W	" .392610E-01
CONSTANT	" .217281

1220

AVERAGED DATA
DRUG ONE

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CENTROIDS OF GROUPS IN REDUCED SPACE

GROUP 1 .45254
GROUP 2 -.77578

PREDICTION RESULTS -

ACTUAL GROUP NAME	CODE	N OF CASES	PREDICTED GROUP MEMBERSHIP	
			GROUP 1	GROUP 2
GROUP 1	2	14	11. 57.9 PCT	3. 15.8 PCT
GROUP 2	3	11	3. 15.8 PCT	8. 42.1 PCT

100.0 PERCENT OF KNOWN CASES CORRECTLY CLASSIFIED

CHI-SQUARE = 19.000 SIGNIFICANCE = .000

AVERAGED DATA
DRUG ONE

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TASK NAME DRUG TWO
(*DRUG ED 2)
DISCRIMINANT GROUPS=SMGCAT(0 1)/VARIABLES=V25AH1MN V25HE1MN V40AR1MN V40HE1MN
CV1MN V25FR1MN V40FR1MN IVFP1MN V25AR3MX V25HE3MX V40AR3MX
V40HE3MX CV3MX V25FR3MX V40FR3MX IVFP3MX/
ANALYSIS=V25AH1MN TO CV1MN V25FR1MN V40FR1MN IVFP1MN/
METHOD=MNRESID/
OPTIONS 2 3 4 5 11 12 13 19
STATISTICS 1 2 3

DISCRIMINANT OPTIONS 13 THRU 19 NOT YET IMPLEMENTED AND WILL BE IGNORED

20096(DECIMAL) CM NEEDED FOR DISCRIM

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AVGPAGED DATA

DRUG TWO FILE: RONNAME (CREATION DATE: 02/06/77)

GROUP COUNTS

NUMBER	GROUP 1	GROUP 2	TOTAL
9.	6.	15.	

MEANS	GROUP 1	GROUP 2	TOTAL
V25AP1MN	2.09222	1.90667	1.97800
V25MF1MN	2.54333	2.28667	2.44067
V40MF1MN	3.27667	3.46500	3.41200
V40MF1MK	4.73111	4.75500	4.74067
V40MF1MX	1.18000	1.19133	1.18533
CV1MN	2.91111	2.92000	2.90667
V75AP1MN	4.47778	4.29333	4.40000
V40MF1MN	17.12333	35.61111	24.51933
V1P1MN	*.41000	*.18333	*.31933
V25MF1UX	*.36222	*.37167	*.36600
V75MF1UX	*.47889	*.67333	*.54667
V40MF3UX	*.84444	*.04000	*.92267
V40MF3MK	.02778	*.20500	*.06533
C1MN	*.19333	*.26833	*.27133
V25FR1MK	*.02667	*.03111	*.02933
V50FR3MK	17.62111	7.50667	13.57533

STANDARD DEVIATIONS

GROUP 1	GROUP 2	TOTAL	
V25AP1MN	*.80986	*.54149	*.70799
V25MF1MN	*.85025	*.68104	*.77171
V40MF1MN	1.01583	1.66258	1.01501
V40MF1MK	1.31547	1.21118	1.24337
V40MF1MX	*.51242	*.51168	*.50623
C1MN	*.10971	*.09798	*.10342
V25FR1MN	*.01090	*.17151	*.11916
V10Y1LUX	3.93850	31.71863	21.15577
V1P1MN	*.56232	*.61834	*.57485
V25MF3UX	*.85621	1.04695	*.91311
V25MF3MX	*.83912	*.77209	*.79123
V10MF3MK	*.83137	*.12529	*.92576
V10MF3MX	*.37662	*.42151	*.39415
C1MN	*.21560	*.23017	*.23665
V25FR3MK	*.40853	*.42107	*.39913
V40FP3MK	34.11469	19.52359	28.76585

AVERAGED DATA
DRCG TWO

WITHIN GROUPS COVARIANCE MATRIX

	V25A1MN	V25E1MN	V40A1MN	V40E1MN	CV1MN	V25F1MN	V40F1MN	IVFP1MN	V25AR3MN
V25A1MN	*51722	*62613	1.10733	1.66471					
V25E1MN	*53422	*59768	1.28248	*27553					
V40A1MN	*62271	*65897	*11084	*01632	*0075	*01110	*01534		
V40E1MN	*67041	*24007	*07194	*07755	*0383	*0414	*09596	396.47683	
CV1MN	*21307	*03459	*01856	*05766	*2.1253	*45743	*1.3729	*2.32929	34164
V25F1MN	*05028	*05110	*3.48934	*7.61364	*04112	*07339	*07497	*3509	
V40F1MN	*05486	*03364	*1.48505	*2.60018	*01625	*01715			
IVFP1MN			*26215	*11429	*0939	*01715	*05065		
V25A2MX	*23521	*41620	*41620	*21258	*16607	*01715	*47454		
V25E2MX	*32370	*35851	*36677	*49855	*00121	*02043	*04098	4.20796	
V40A2MX	*28455	*18437	*21789	*45188	*00268	*01206	*08492		
V40E2MX	*10590	*15211	*05468	*05081	*1.11340	*02708	*00019	*06400	
CV3MX	*32235	*35187	*22250	*27914	*17142	*01221	*02917	*04146	
V25F3MX	*02023	*00536	*00536	*07092	*04013	*052276	*09835	*04166	
V40F3MX	*01794	*8.68397	*12.84615	*12.44259	*1.52276		*1.80199	*144.57884	
IVFP3MX	*9.0K918								
V25E3MX									
V40A3MX									
V40E3MX									
V40F3MX									
IVFP3MX									

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AVERAGED DATA
DRCG TWO
FILE NNAME (CREATION DATE = 02/06/77)
DISCRIMINANT ANALYSIS

ANALYSIS NUMBER 1

MAXIMUM STEPS 12
TOLERANCE LEVEL .00010
F FOR DELETION .00500

F FOR INCLUSION .01000
SOLUTION METHOD = STEPWISE. SELECT VARIABLE WHICH WILL MINIMIZE DISTANCE FUNCTION.
PRIOR PROBABILITIES = EQUAL

GROUP 1 GROUP 2

.50000 .50000

123

AVERAGED DATA
DRUG TAU

VARIABLES IN THE ANALYSIS		F TO REMOVE
VARIABLE	ENTRY CRITERION	
V2AP1MN	*.79710	2.85567
V2HE1MN	*.35017	.86490
V4AR1MN	*.65011	2.08697
V4HE1MN	*.38638	*.20247
CV1MN	*.60039	3.43595
V2FP1MN	*.38638	1.13231
V4CP1MN	*.41634	*.23936
IVFP1MN	*.82266	4.93354

NUMBER REMOVED 0 EIGENVALUE .215690

VARIABLE	CANONICAL CORRELATION	PERCENT OF TRACE	WILKS LAMBDA	CHI-SQUARE	D.F.	SIGNIFICANCE
	.82658	100.0	*.31677	11.49389	8	.175

1 FUNCTIONS WILL BE USED IN REMAINING ANALYSES

STANDARDIZED DISCRIMINANT FUNCTION COEFFICIENTS

V25AR1MN	*7.15185
V25FP1MN	2.47233
V4AP1MN	5.40940
V4HE1MN	*1.42724
CV1MN	*1.38254
V25FA1MN	*1.47916
V4F1MN	2.21495
IVFP1MN	1.59824

UNSTANDARDIZED DISCRIMINANT FUNCTION COEFFICIENTS

V25AP1MN	*10.1017
V25FP1MN	3.1244
V4AP1MN	5.37499
V4HE1MN	*1.14788
CV1MN	*1.91277
V25FA1MN	*16.10728
V4F1MN	18.4946
IVFP1MN	*76.3988E-01
CONSTANT	*2.91130

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AVERAGED DATA
DRUG TWO

CENTROIDS OF GROUPS IN REDUCED SPACE

GROUP 1 -1.11634
GROUP 2 1.67450

PREDICTION RESULTS *

ACTUAL GROUP NAME	GROUP CODE	N OF CASES	PREDICTED GROUP MEMBERSHIP	
			GROUP 1	GROUP 2
GROUP 1	2	13	12. 80.0 PCT	1. 6.7 PCT
GROUP 2	3	9	4. 26.7 PCT	5. 33.3 PCT

113.3 PERCENT OF KNOWN CASES CORRECTLY CLASSIFIED

CHI-SQUARE = 24.067 SIGNIFICANCE = .000

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AVERAGED DATA
DRUG TWO
FILE NONAME (CREATION DATE = 02/06/77)

DISCRIMINANT ANALYSIS

ANALYSIS NUMBER 2

TOLERANCE LEVEL

.00010

MAXIMUM STEPS

12

F FOR INCLUSION

.01000

F FOR DELETION

.00500

SOLUTION METHOD = STEPWISE. SELECT VARIABLE WHICH WILL MINIMIZE DISTANCE FUNCTION.

PRIOR PROBABILITIES = EQUAL

GROUP 1 GROUP 2

.50000 .50000

ALL ELIGIBLE VARIABLES INCLUDED

125

AVERAGED DATA
DRUG TWO

----- VARIABLES IN THE ANALYSIS -----

VARIABLE	ENTRY CRITERION	F TO REMOVE
V25AR3MX	-.80319	5.67790
V25HE3MX	-.47897	1.46140
V40AR3MX	-.51871	2.00705
V40HE3MX	-.59495	.05878
CV3MX	-.91997	1.19273
V25FR3MX	-.41457	.98593
V40FR3MX	-.47782	.02246
IVFP3MX	-.85520	1.35114

----- VARIABLES NOT IN THE ANALYSIS -----

VARIABLE	TOLERANCE	F TO ENTER	ENTRY CRITERION
----------	-----------	------------	-----------------

NUMBER REMOVED	EIGENVALUE	CANONICAL CORRELATION	PERCENT OF TRACE	WILKS LAMBDA	CHI-SQUARE	D.F.	SIGNIFICANCE
0	1.57379	.78196	100.0	.38853	9.45379	8	.305

1 FUNCTIONS WILL BE USED IN REMAINING ANALYSES

STANDARDIZED DISCRIMINANT FUNCTION COEFFICIENTS

	1
V25AR3MX	2.95530
V25HE3MX	-.1.75918
V40AR3MX	-.1.56592
V40HE3MX	-.29985
CV3MX	-.67593
V25FR3MX	.77122
V40FR3MX	-.12728
IVFP3MX	-.83326

UNSTANDARDIZED DISCRIMINANT FUNCTION COEFFICIENTS

	1
V25AR3MX	.5.14102
V25HE3MX	-.1.95181
V40AR3MX	-.1.97908
V40HE3MX	-.323895
CV3MX	-.69765
V25FR3MX	.823349
V40FR3MX	-.318896
IVFP3MX	-.289670E-01
CONSTANT	-.384109

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AVERAGED DATA
DRUG TWO

CENTROIDS OF GROUPS IN REDUCED SPACE

GROUP 1	.95357
GROUP 2	1.43036

PREDICTION RESULTS -

ACTUAL GROUP NAME	CODE	N OF CASES	PREDICTED GROUP MEMBERSHIP	
			GROUP 1	GROUP 2
GROUP 1	2	13	10. 66.7 PCT	3. 20.0 PCT
GROUP 2	3	9	4. 26.7 PCT	5. 33.3 PCT

100.0 PERCENT OF KNOWN CASES CORRECTLY CLASSIFIED

CHI-SQUARE = 15.000 SIGNIFICANCE = .000

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AVERAGED DATA
DRUG TWO

TASK NAME DRUG THREE
*SELECT IF (DRUG EQ 3)
DISCRIMINANT GROUPS=SMSGCAT(0 1)/VARIABLES=V25AH1MN V25HE1MN V40AR1MN V40HE1MN
CV1MN SVC1MN FVC1MN V25FR1MN V40FR1MN CSV1MN FEFV1MN
IVFP1MN V25AR3MX V25HE3MX V40AR3MX V40HE3MX CV3MX SVC3MX FEV3MX
FVC3MX V25FR3MX V40FR3MX CSVV3MX FEFV3MX IVFP3MX/
ANALYSIS=V25AR1MN TO IVFP1MN/METHOD=MINRESID/
ANALYSIS=V25AR3MX TO IVFP3MX/METHOD=MINRESID/

OPTIONS 2 3 4 5 11 12 13 19
STATISTICS 1 2 3

DISCRIMINANT OPTIONS 13 THRU 19 NOT YET IMPLEMENTED AND WILL BE IGNORED

21056(DECIMAL) CM NEEDED FOR DISCRIM

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AVERAGED DATA
DRUG THREE
FILE: NQHME (CREATION DATE = 02/06/77)

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02/06/77

GROUP COUNTS

NUMBER	GROUP 1	GROUP 2	TOTAL
12.	8.	20.	

MEANS	GROUP 1	GROUP 2	TOTAL
V75AP1MH	1.97111	1.71875	1.87950
V75HF1MH	2.23467	2.19275	2.22160
V10AP1MH	3.38750	3.14625	3.37100
V10HF1MH	4.51167	4.15600	4.44760
V40HF1MH	1.52250	1.39010	1.43150
CV1%	9.10750	8.46500	8.80550
SVC1%	8.10500	7.61875	7.91050
FVC1%	10.91813	9.27875	10.32500
FVC1%	*24917	*13875	*25500
V25HF1MH	*36750	*16000	*36450
V40HF1MH	*16.5083	15.28510	16.01750
CV5V1%	76.18117	80.51750	77.91550
FFV1%	20.12811	17.18625	19.31150
V25AH1%	*13583	*51175	*31100
V25HF1%	*57167	*25125	*41450
V10AP1%	*45917	*91250	*64850
V10HF1%	1.04843	*42125	*79750
V40HF1%	*113750	*111000	*12650
CV3%	*19331	*310000	*316000
SVC1%	*04817	*12750	*09550
FVC1%	*81111	*00250	*88760
V25HF1%	*13500	*18750	*151600
V25HF1%	*03750	*09000	*01150
V40HF1%	*1.35250	*34125	*37500
CV5V1%	22.42167	1.72500	14.16300
FFV1%	*60333	33.30825	13.68450

STANDARD DEVIATIONS

STANDARD DEVIATIONS	GROUP 1	GROUP 2	TOTAL
V75AP1MH	*82882	*56146	*72653
V75HF1MH	*87785	*41724	*87190
V40AP1MH	1.11421	*75165	*66298
V40HF1MH	1.57480	1.11483	1.38150
V40HF1%	*51106	*39609	*50903
CV1%	2.45808	1.94291	2.26647
SVC1%	1.82400	1.41450	1.66447
FVC1%	3.11247	2.30462	2.83292
V25HF1%	*10.94	*18117	*16343
V40HF1%	*12835	*12284	*12479
CV5V1%	4.12297	3.26170	3.76382

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AVERAGED DATA

DRUG THREE

	10.11342	7.54792	9.21129
FFFV1MN	6.88457	9.86914	7.24501
V25AP1MN	.60042	.40454	.56343
V25H13MX	.45449	.68917	.56632
V30AR3MX	.68593	.47346	.64155
V40HF3MX	.59116	1.46737	1.01640
CV3MX	.41971	.40366	.42108
SVC1MX	1.14680	.57456	.97581
FEV3MX	.2401	.55851	.71567
FVC1MX	2.51189	.62064	2.00514
V25FP3MX	.35682	.80932	.60235
V40FR3MX	.33556	.28571	.31802
CVSV3MX	4.76705	4.31354	4.55371
FFFV3MX	77.13175	6.17619	59.71668
V25P3MX	20.82039	32.80810	30.29372

WITHIN GROUPS COVARIANCE MATRIX

	V25AR1MN	V25HE1MN	V40AR1MN	V40HF1MN	CV1MN	SVC1MN	FEV1MN	FVC1MN	V25FR1MN
V25AR1MN	.54250								
V25HE1MN	.60941	.84236							
V40AP1MN	.61846	.72131	.97839						
V40HE1MN	.90909	1.11770	1.31586	2.01618					
CV1MN	.03598	-.01154	.11544	.21322	.26030				
SVC1MN	.43208	.46757	.69085	1.15501	.92311	.51220			
FEV1MN	.30177	.36719	.68215	1.24708	.69414	3.61483	2.87875		
FVC1MN	-.39922	-.36079	-.01776	.34024	1.04571	.544859	4.14353	7.99278	
V25FP1MN	.01272	.05203	.02410	.05714	.00531	.07993	.08149	.05255	.02605
V40FR1MN	.02183	.04532	.03053	.08050	.03200	.13687	.10915	.15902	.01116
CVSV1MN	-.48018	.95088	.08842	-.01932	1.41996	1.61348	1.82258	2.96124	-.02113
FFFV1MN	5.29251	5.92154	5.81996	8.24111	-.1.47048	-.6.19527	-.3.77791	-.17.61105	.00781
V25P1MN	-.41492	-.2.28130	.34984	-.97263	1.46817	4.70849	1.82482	3.86748	-.09747
V25AP3MX	-.18988	-.14159	-.17083	-.19121	-.04857	-.33669	-.14832	-.07452	.00941
V25H13MX	-.02090	-.10320	.06454	-.01020	-.00519	.06164	-.00764	-.10936	.00117
V30AR3MX	-.05533	.01988	-.03440	-.04978	-.01245	-.06072	-.01362	-.02677	.03197
V40HF3MX	-.08347	-.27791	.02433	-.04276	-.00196	.25835	.14743	.26489	-.03770
CV3MX	.00108	.01501	.01030	-.0324	-.14892	-.44453	-.28954	-.44164	-.00153
SVC1MX	-.07849	-.19977	-.13478	-.32973	-.10961	-.47607	-.31791	-.38887	-.07020
FEV1MX	-.01638	-.05124	-.07584	-.08024	-.07647	-.12962	-.05970	-.05669	-.00949
FVC1MX	-.01409	-.13745	-.75740	-.70289	-.04291	.11604	-.30908	-.25899	-.03179
V25FP3MX	.07647	.00121	.20670	.18783	.01720	.03612	.03777	.11655	.04508
V40FR3MX	.01875	-.04084	.01179	-.03806	-.01195	-.07164	-.05220	-.12997	-.02920
CVSV3MX	.31283	.67694	.72295	.55489	-.1.51127	-.3.41894	-.1.78908	-.3.36326	.08049
FFFV3MX	4.59567	4.84435	25.87007	22.79230	1.98000	-.5.14390	7.16876	-.2.41665	.58120
V25P3MX	7.12247	11.78846	5.36435	8.18286	-.1.37491	-.4.63356	-.0.50300	-.22.61373	2.12370
	V40FR1MN	CVSV1MN	FFFV1MN	V25P1MN	V25AR3MX	V25HE3MX	V40HF3MX	V40AR3MX	CV3MX
V25FR1MN	.01612								
CVSV1MN	.13136	14.55595							
FFFV1MN	-.06110	-.7.75513	84.95511						
V25P1MN	.19265	8.57479	-.18.48246	52.48256					
V25AP3MX	.00156	.10532	-.1.06646	-.8.80077	-.28395				
V25H13MX	-.03336	-.02466	-.18889	2.79110	-.01531	-.31116			
V30AR3MX	.02777	-.01563	-.06578	-.37256	-.26198	-.04916	-.37470		

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DISCRIMINANT ANALYSIS

AVERAGED DATA
DRUG THREE
FILE "NAME" (CREATION DATE # 02/06/77)

PAGE 26

	02/06/77	02/06/77	02/06/77
ANALYSIS NUMBER	1		
TOLERANCE LEVEL	*.00010	MAXIMUM STEPS	52
F FOR INCLUSION	*.0100	F FOR REJECTION	*.00500
SOLUTION METHOD = STEPWISE. SELECT VARIABLE WHICH WILL MINIMIZE DISTANCE FUNCTION.			
PRIOR PROBABILITIES = EQUAL			
GROUP 1	GROUP 2		
*.50000	*.50000		

AVERAGED DATA
DRUG THREE
FILE "NAME" (CREATION DATE # 02/06/77)

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AVERAGED DATA
DRUG THREE
FILE "NAME" (CREATION DATE # 02/06/77)

PAGE 26

	02/06/77	02/06/77	02/06/77
ANALYSIS NUMBER	1		
TOLERANCE LEVEL	*.00010	MAXIMUM STEPS	52
F FOR INCLUSION	*.0100	F FOR REJECTION	*.00500
SOLUTION METHOD = STEPWISE. SELECT VARIABLE WHICH WILL MINIMIZE DISTANCE FUNCTION.			
PRIOR PROBABILITIES = EQUAL			
GROUP 1	GROUP 2		
*.50000	*.50000		

CENTROIDS OF GROUPS IN REDUCED SPACE

GROUP 1	*9539
GROUP 2	*1.47809

UNSTANDARDIZED DISCRIMINANT FUNCTION COEFFICIENTS

	1
V25A1MN	5.09770
V25H1MN	*2.05704
V25H1AN	*2.61406
V25A1AN	2.65781
V40H1MN	*5.86634
C1MN	*64095
SIC1MN	1.40014
FFV1MN	*16154
V40F1MN	3.59664
CUS1MN	*3.21281
FFP1MN	*16565
IVP1MN	

STANDARDIZED DISCRIMINANT FUNCTION COEFFICIENTS

	1
V25A1MN	7.01650
V25H1MN	*2.30119
V25H1AN	*2.1512
V40H1MN	1.94135
V40F1MN	-1.15836
C1MN	*26.766
SIC1MN	*38694
FFV1MN	*1.79449
V40F1MN	0.956111
CUS1MN	*1.44852
FFP1MN	*2227391E-01
IVP1MN	11.8258
CONSTANT	

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AVERAGED DATA
DURING THREE
PREDICTION RESULTS =

ACTUAL GROUP NAME	CASES	N OF CASES	PREDICTED GROUP MEMBERSHIP
GROUP 1	2	14	GROUP 1 GROUP 2
CAMP 1			11*. 55.0 PCT 15.0 PCT
GROUP 2	3	11	3*. 15.0 PCT 40.0 PCT

95.0 PERCENT OF KNOWN CASES CORRECTLY CLASSIFIED

CHI-SQUARE = 16.200 SIGNIFICANCE = .000

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AVERAGED DATA
DURING THREE
FILE NUMBER (CREATION DATE = 02/06/77)
***** DISCRIMINANT ANALYSIS *****

ANALYSIS NUMBER 2 MAXIMUM STEPS 52
TOLERANCE LEVEL .00010
F FOR INCLUSION .01000 F FOR DELETION .00500
SOLUTION METHOD = STEPWISE. SELECT VARIABLE WHICH WILL MINIMIZE DISTANCE FUNCTION.
PRIOR PROBABILITIES = EQUAL
GROUP 1 GROUP 2
.50000 .50000

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AVERAGED DATA
DRUG THREE
STANDARDIZED DISCRIMINANT FUNCTION COEFFICIENTS

	1
V25AP3W	1.65436
V25HF3W	-1.95677
V40LW3W	-1.14441
V40HF3W	2.54939
CV3W	-1.48938
SVC3W	.32766
FV3W	-2.47390
FVC3W	-2.24344
V25HF3W	.16198
V40HF3W	.41010
CVS3W	1.79485
IVFP3W	2.68247

UNSTANDARDIZED DISCRIMINANT FUNCTION COEFFICIENTS

	1
V25AP3W	2.93624
V25HF3W	-1.45965
V40AP3W	-1.79163
V40HF3W	2.41635
CVS3W	-3.53148
SVC3W	.155890
FV3W	-1.45541
FVC3W	-1.121196
V25HF3W	.268417
V40HF3W	1.35241
CVS3W	.284390
IVFP3W	.485448E-01
CONSTANT	-1.68510

CENTROIDS OF GROUPS IN REDUCED SPACE

GROUP 1	*1.11499
GROUP 2	1.67249

AVERAGED DATA
THRU THREE
PREDICTION RESULTS =

ACTUAL GROUP CODE	N OF CASES	PREDICTED GROUP MEMBERSHIP	GROUP 1	GROUP 2
GROUP 1	2	14	12.	7.
			60.0 PCT	11.0 PCT
GROUP 2	3	11	4.	7.
			20.0 PCT	35.0 PCT

95.0 PERCENT OF KNOWN CASES CORRECTLY CLASSIFIED

CHI-SQUARE = 16.200 SIGNIFICANCE = .000

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AVERAGED DATA
THRU THREE
DISCRIMINANT
STATISTICS
FINISH

DBUG 4(OR 4)
(DRUG EQ 4)
GRN01\$55C05C010 1) /VARIABLES=V25AR1NN V25H1NN V40H1NN V40H1NN
CVMM SVCMM FETMM FV1MM V25FR1NN V40FR3NN V40HE3NN CVMM SVCMM FETMM
IVFP1NN V25FR3NN V40FR3NN CVMM FETV3NN IVFP3NN
FVC3NN V25FR3NN V40FR3NN CVMM FETV3NN IVFP3NN
ANALYSIS=DISCRIMINANT TU IVFP1NN/METHOD=INSTEP10/
ANALYSIS=DISCRIMINANT TO IVFP3NN/METHOD=MINRES10/
OPTIONS
1 2 3
STATISTICS
1 2 3
FINISH

DISCRIMINANT OPTIONS 13 THRU 19 NOT YET IMPLEMENTED AND WILL BE IGNORED
21056 DECIMAL CM NEEDED FOR DISCRIM

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AVERAGED DATA

(CREATION DATE = 02/06/77)

FILE NUMBER

GROUP COUNTS

NUMBER	GROUP 1	GROUP 2	TOTAL
6.	6.	14.	

MEANS

	GROUP 1	GROUP 2	TOTAL
V25AP14N	1.87500	1.58000	1.74857
V25AF14N	2.25175	1.89667	2.10071
V40AF14N	3.31500	3.31813	3.35031
V40AF14N	4.49125	4.29167	4.40571
V40AF14N	1.29175	1.18833	1.24857
FV14N	8.35175	8.04000	8.21929
SV14N	7.86175	7.38667	7.62641
FEV14N	10.00000	9.65500	9.87214
V25AF14N	2.51250	2.28433	2.41443
V40AF14N	3.66225	2.96000	3.27366
CUS14N	10.85250	15.02500	14.72643
FFV14N	75.24600	76.49333	78.06857
FUP14N	15.98250	19.41500	17.66766
V25AF14N	1.88750	0.73333	1.39299
V25AF14N	4.20000	0.61667	1.56443
V40AF14N	4.12900	0.01000	0.07571
V40AF14N	4.23750	0.41667	4.20211
CV34X	15.00000	0.01000	0.08143
SU14X	29.125	1.1167	21.429
FFV34X	21.250	0.7333	0.04000
FUC34X	0.88475	0.6833	1.2286
V25AF14N	0.02125	0.04167	0.00000
V40AF14N	0.02500	0.12500	0.05214
V40AF14N	1.61125	0.69333	1.33214
CSV34X	1.30000	1.25667	2.0429
FFV34X	0.01375	3.489500	2.91000

STANDARD DEVIATIONS

	GROUP 1	GROUP 2	TOTAL
V25AF14N	.53364	.31843	.46399
V25AF14N	.64314	.51750	.60619
V40AF14N	.69320	.68473	.78151
V40AF14N	1.26206	1.16457	1.19045
CV34X	.70177	.52625	.61207
SU14X	2.47956	2.11130	2.21858
FV14N	1.68883	1.49714	1.56442
FV14N	2.76156	1.80765	2.32113
V25AF14N	2.11482	1.0944	1.0483
V40AF14N	1.0028	1.4408	1.2065
CSV34X	5.77080	6.30538	5.76366

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AVERAGE DATA	
FREQUENCY	1000
PERIOD	6.36150
PERIOD	6.25177
V250F100K	5.98470
V250F100K	4.26816
V250F100K	5.8870
V250F100K	4.91436
V250F100K	4.5509
V250F100K	6.01756
V250F100K	6.76084
CUSUM	3.18775
SUM10X	6.86840
FF100K	6.60433
FF100K	6.3100
FF100K	6.0988
FF100K	9.295894
V250F100K	9.0509184
CUSUM	3.19052
FF100K	7.11442
V250F100K	21.18129
	6.71145
	5.98470
	4.26816
	5.8870
	4.91436
	4.5509
	6.01756
	6.76084
	3.18775
	6.86840
	6.60433
	6.3100
	6.0988
	9.295894
	9.0509184
	3.19052
	7.11442
	21.18129
	7.54920
	6.21697
	3.6788
	5.6385
	5.5044
	7.94721
	4.47020
	6.82979
	5.7162
	5.91141
	1.76930
	1.41950
	3.09440
	4.89521
	5.81173
	17.27450

WITHIN GROUPS COVARIANCE MATRIX

AVERAGED DATA		PAGF		3H	
	DRIVING FORUM				
V49HE3MX	*.06287	*.05554	*.2.99714	1.23799	.09820
V49HE3MX	*.00242	*2.19556	*.89817	*.60397	.07946
SIC3MX	*1.47448	*.20816	*.03291	.13516	.17400
FF3MX	*.00090	*.03041	1.41187	*.21947	.10201
FVC3MX	*.01331	1.03740	2.28551	*.27076	*.11143
V49FB3MX	*.00035	*2.08843	*.51256	.75119	*.01076
U49FB3MX	*.01675	*.52565	*.50142	.26183	*.00784
C15V3MX	*.00184	*21.49620	*.96086	*4.76585	*.82818
F15V3MX	*.09448	*9.49603	*16.85631	12.77413	1.66064
115FP3MX	*15265	*7.91669	44.40122	*74.52817	*2.17932
SVC3MX	FEV3MX	FVC3MX	V25FB3MX	V40FB3MX	CVSV3MX
SIC3MX	*73671	33064	25980	25980	CVSV3MX
FF3MX	*12527	*11585	*.02018	*.01433	CVSV3MX
V49FB3MX	*.02819	*.06998	*.02554	*.02026	V49FB3MX
V49HE3MX	*.04684	*.04855	*.02554	*.01915	*.02554
CVSV3MX	1.96825	9.20455	*.05790	.02571	*.16810
FF3V3MX	2.10886	1.50517	*2.33617	*.02571	*.05040
115FP3MX	*.58137	*.60322	3.93655	*1.85651	*1.56706

VERAGED DATA
WING, FOUR
FILE NUMBER (CRAFT)

(CERTIFICATION DATE = 02/06/77)

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TRUSTANCE LEVEL	MAXIMUM STEPS	52
F FOR INCLUSION	F FOR DELETION	*00500
SOLUTION METHOD = STEPWISE.	SELECT VARIABLE WHICH WILL MINIMIZE DISTANCE FUNCTION.	
PHIPIP PROBABILITIES = EQUAL		

GROUP 1 GROUP 2
• \$50000 • \$50000

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DISCREMINANT ANALYSIS

AVERAGEN DATA
NOTIC P-OUT

ALL ELIGIBLE VARIABLES INCLUDED

VARIABLES IN THE ANALYSIS

VARIABLE	ENTRY CRITERION	F TO REMOVE
V25F1MN	*.40446	1.31142
V25H1MN	*.05119	1.13863
V40H1MN	*.79229	3.91419
V40M1MN	*.49441	5.92441
V40W1MN	*.57776	5.49357
CV1MN	*.15320	3.77264
SVC1MN	*.07111	5.70068
FFV1MN	*.02787	*.89260
FFV1M1	*.01747	*.55723
V25F1MN	*.26448	4.60744
V40W1MN	*.20394	4.04080
CVS1MN	*.71632	7.14783
IVF1MN		

SUMMARY OF APPROVED

0 62.63311 .9211 100.0

FUNCTIONS WILL BE USED IN REMAINING ANALYSES

VARIABLE	TOLERANCE	F TO ENTER	F TO REMOVE	P-OUT
FFV1MN	.00000	0	*.07269	

1 FUNCTIONS WILL BE USED IN REMAINING ANALYSES

STANDARDIZED DISCRIMINANT FUNCTION COEFFICIENTS

V25H1MN	*25.56442
V25W1MN	30.61669
V40H1MN	*31.61044
V40M1MN	46.62501
CV1MN	*38.66759
SVC1MN	29.05224
FFV1MN	*19.65752
IVF1MN	3.38605
V25F1MN	*8.61709
V40F1MN	*15.48510
CVS1MN	30.13121
IVF1MN	5.51837

AVERAGED DATA
DRUG FOUR

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UNSTANDARDIZED DISCRIMINANT FUNCTION COEFFICIENTS

	1
V25H1MN	-55.0985
V25H2MN	50.5099
V40H1MN	+41.0069
V10H1MN	19.1657
CV1MN	+61.1752
SVC1MN	11.0950
FV1MN	-12.5714
FVC1MN	1.45479
V25L4MN	-79.1632
ZAPF1MN	+121.519
CVS1MN	5.20989
TVF1MN	.884785
CONSTANT	-17.8793

CENTROIDS OF GROUPS IN REDUCED SPACE

GROUP 1 -6.34540
GROUP 2 8.46054

PREDICTION RESULTS *

ACTUAL GROUP NAME	CODE	N OF CASES	PREDICTED GROUP MEMBERSHIP	
			GROUP 1	GROUP 2
GROUP 1	2	14	11. 78.6 PCT	3. 21.4 PCT
GROUP 2	3	11	4. 28.6 PCT	7. 50.0 PCT

128.6 PERCENT OF KNOWN CASES CORRECTLY CLASSIFIED

CHI-SQUARE = 34.571 SIGNIFICANCE = .000

AVERAGED DATA
DRUG FOUR
FILE: VNAME (CREATION DATE = 02/06/77)

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----- DISCRIMINANT ANALYSIS -----
ANALYSIS NUMBER 2
TOLERANCE LEVEL .00010 MAXIMUM STEPS 52
F FOR INCLUSION .01000 F FOR DELETION .00500
SOLUTION METHOD = STEPWISE. SELECT VARIABLE WHICH WILL MINIMIZE DISTANCE FUNCTION.
PRIOR PROBABILITIES = EQUAL
GROUP 1 GROUP 2
.50000 .50000

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AVERAGED DATA
DRUG FOUR

ALL ELIGIBLE VARIABLES INCLUDED

----- VARIABLES IN THE ANALYSIS -----

VARIABLE	ENTRY CRITERION	F TO REMOVE
V25AP3MX	-.03657	.81011
V25HE3MX	-.62601	2.04194
V30AP3MX	-.19458	5.08812
V40HE3MX	-.06684	1.42535
CV3MX	-.55036	3.53707
SVC3MX	-.01331	.09825
FEV3MX	-.35072	6.44251
FVC3MX	-.25406	2.21823
V25FR3MX	-.22212	1.74299
V40FR3MX	-.12399	4.10587
FEVV3MX	-.41641	5.47958
IVFP3MX	-.88858	.19906

----- VARIABLES NOT IN THE ANALYSIS -----

VARIABLE	TOLERANCE	F TO ENTER	ENTRY CRITERION
CVSV3MX	.00000	0	-.03553

NUMBER REMOVED	EIGENVALUE	CANONICAL CORRELATION	PERCENT OF TRACE	WILKS LAMBDA	CHI-SQUARE	D.F.	SIGNIFICANCE
0	33.16183	.98526	100.0	.02927	24.71776	12	.016

1 FUNCTIONS WILL BE USED IN REMAINING ANALYSES

STANDARDIZED DISCRIMINANT FUNCTION COEFFICIENTS

1

V25AP3MX	-2.06687
V25HE3MX	-2.59913
V30AP3MX	5.82905
V40HE3MX	-6.01691
CV3MX	3.31936
SVC3MX	-.64386
FEV3MX	8.24332
FVC3MX	-3.02748
V25FR3MX	6.04226
V40FR3MX	5.77933
FEVV3MX	-6.05906
IVFP3MX	1.60586

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AVERAGED DATA
MEAN FOUR
UNSTANDARDIZED DISCRIMINANT FUNCTION COEFFICIENTS

	1
V25AB3W	+5.69569
V75MF3W	+4.616457
V10AB3W	10.5821
V10HE3W	+5.6571
C1JW	7.05869
S1C3W	+.775936
FV3W	14.4209
FUC4W	+6.16446
V2KVP3W	14.1571
V10FP3W	18.6794
FFFU3W	+1.04256
TFP1W	.9296115+01
CONSTANT	3.58067

CENTROIDS OF GROUPS IN REDUCED SPACE

GROUP 1	+4.61718
GROUP 2	6.15621

PREDICTION RESULTS *

ACTUAL GROUP NAME	N OF CASES	PREDICTED GROUP NAME	GROUP 1	GROUP 2
GROUP 1	2	14	11*	3*
GROUP 2	3	11	74.6 PCT	21.4 PCT
			2*	9*
			14.3 PCT	64.3 PCT

147.9 PERCENT OF RANDOM CASES CORRECTLY CLASSIFIED

CHI-SQUARE = 48.286 SIGNIFICANCE = .000

AVERAGED DATA
DRUG + MUR

PUS COMPLETED
NUMBER OF CONTROL CARDS READ 75
NUMBER OF ERRORS DETECTED 0

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