

LONDON  
SCHOOL of  
HYGIENE  
& TROPICAL  
MEDICINE



Law, GR; Smith, AG; Roman, E; Ukcccr Study Group (Peto, J., -As A Member Of), (2002) The importance of full participation: lessons from a national case-control study. *British journal of cancer*, 86 (3). pp. 350-5. ISSN 0007-0920 DOI: <https://doi.org/10.1038/sj.bjc.6600092>

Downloaded from: <http://researchonline.lshtm.ac.uk/15764/>

DOI: [10.1038/sj.bjc.6600092](https://doi.org/10.1038/sj.bjc.6600092)

#### Usage Guidelines

Please refer to usage guidelines at <http://researchonline.lshtm.ac.uk/policies.html> or alternatively contact [researchonline@lshtm.ac.uk](mailto:researchonline@lshtm.ac.uk).

Available under license: <http://creativecommons.org/licenses/by-nc-nd/2.5/>

# The importance of full participation: lessons from a national case–control study

**United Kingdom Childhood Cancer Study Investigators (writing committee: GR Law, AG Smith, E Roman)\*<sup>1,2</sup>**

<sup>1</sup>Leukaemia Research Fund, Centre for Clinical Epidemiology, 30 Hyde Terrace, University of Leeds, Leeds LS2 9LN, UK

Differential participation between cases and controls can lead to biased estimates of risk. However, the effects of participation are often ignored. We report a detailed analysis of locations of residence for participants and non-participants in a large, national case–control study of childhood cancer in Great Britain, using the 1991 census. The initial selection of 7669 controls, taken from lists of those registered with a General Practitioner, was representative of the British population in respect to an areal-based index of material deprivation. However, parents of controls agreeing to participate were living in more affluent areas than initially selected controls and their matched 3838 cases. The three components of the deprivation index, persons unemployed, households not owning a car or their home were similarly associated with participation. Other census characteristics, such as proportion of flat dwellers and centrally heated households were also associated with control participation. Population density of the local area was not different between participating controls and their matched cases. However, initially selected controls lived in more urban areas than their cases. Such differences are not unique to this study, as they are an inevitable consequence of incomplete participation. The implications of these differences are discussed, in relation to the difficulty this imposes in the interpretation of studies of disease aetiology.

*British Journal of Cancer* (2002) **86**, 350–355. DOI: 10.1038/sj/bjc/6600092 www.bjcancer.com

© 2002 The Cancer Research Campaign

**Keywords:** childhood cancer; aetiology; census; participation; case-control

Voluntary participation in health-related studies is rarely 100%, and there is evidence that in recent years it has been declining (Hartge, 1999). A particular problem for epidemiological research arises when participation is not randomly distributed across study groups. Unfortunately, whether or not an individual agrees to participate in a project is often associated with the health outcome and with the exposure(s) under investigation; invariably this leads to biased estimates of risk (Rothman and Greenland, 1998). The assessment of the impact of differential participation requires the characteristics of those who do not take part to be compared with those who do. In most studies, however, information on non-participants is often sparse or non-existent. Indeed, in some designs, such as those that employ random digit dialling, investigators are not even able to identify non-participants (Wacholder *et al*, 1992).

Previous attempts aimed at exploring this form of selection bias have relied mainly on re-approaching non-participants and asking a restricted number of questions (e.g. Holt *et al*, 1991; Madigan *et al*, 2000; Wrensch *et al*, 2000). However, even when the identities of non-participants are known, practical and ethical considerations often prohibit the use of such methods. The design of the United Kingdom Childhood Cancer Study (UKCCS) provided a valuable opportunity to investigate the potential impact of participation bias (UKCCS Investigators, 2000). The findings, which used small-area

census data to investigate differences between participating and non-participating subjects, are reported here.

## MATERIALS AND METHODS

A case–control approach was employed, based around a face-to-face interview of the parents of the index children. Study methods have been described in detail elsewhere (UKCCS Investigators, 2000). The base population was defined as all 0–14 year-olds born and resident in Great Britain (GB), registered with a General Practitioner (GP), without a prior malignancy and not in residential local authority care. Data collection was co-ordinated by 10 regional centres using a common protocol and data collection instruments.

### Case selection

Potentially eligible cases were children diagnosed with a confirmed malignancy or benign tumour of the central nervous system (CNS) within the time frame of the study. All diagnostic groups were included in Scotland between 1991–1994, and 1992–1994 in England and Wales. Case accrual continued in England and Wales for non-Hodgkin's lymphomas and leukaemias during 1995 and leukaemias alone in 1996. Diagnostic confirmation was obtained via several sources: Medical Research Council (MRC) treatment trials, UK Childhood Cancer Study Group (UKCCSG), and histopathology review panels (UKCCS Investigators, 2000).

### Control selection

Two control children were matched on age and sex to each participating case, randomly selected from children registered with the

\*Correspondence: Dr GR Law; E-mail: g.r.law@leeds.ac.uk

<sup>2</sup>A complete list of investigators is given in The United Kingdom Childhood Cancer Study: Objectives, Materials and methods. *British Journal of Cancer* 2000; **82**(5): 1073–1102. See Appendix for Management Committee, Regional Investigators and Processing Group

Received 16 November 2001; revised 29 November 2001; accepted 29 November 2001

same (former) Family Health Services Authority (England and Wales) or Health Board (Scotland) as their matched case. The first two controls selected were assigned as 'first-choice', following successful eligibility checks. In order to restrict the analysis to examining the effects of subject participation and to those with a valid address that could be linked to census, GP permission was obtained for all control families classified as 'first-choice'. A proportion of the parents of first-choice controls declined to participate and the selection process was repeated until two control families were enrolled and interviewed. The two controls enrolled into the study were assigned as 'interviewed', with a proportion of these also assigned as first-choice.

### Linkage to the census of Great Britain

Cases and controls were linked, via the postcode of their residence, to the 1991 census of GB, which was conducted around the time of the commencement of the study (The 1991 census, Crown copyright, ESRC Purchase). Every household has a legal requirement to complete the census form, which contained 25 questions, the content ranging from age and sex of occupants to indicators of affluence such as adult employment and housing conditions. To maintain confidentiality, data are released at an areal level, with the smallest geographical units being indivisible and considered to be homogenous. The smallest units in England and Wales were the 108 336 Enumeration Districts (ED) and in Scotland the 38 084 Output Areas (OA). These can be aggregated into 9527 Electoral Wards (England and Wales) and 1002 Postcode Sectors (Scotland).

The address of the child at the time of diagnosis was used to link the study subjects to the census data. The address was assigned a validated postcode, using an automated system (QuickAddress™) and the Postal Address Books available from Royal Mail. All postcodes were assigned to an ED/OA using a lookup program PC2ED for England and Wales and the Postcode to OA databases for Scotland.

### Census data

The census variables of interest, detailed in Table 1, focus on material possessions, employment, household characteristics, and population density. The simple counts provided by the census are given as a proportion of the relevant population (of households or people), as the geographically delimited census regions are not the same size in terms of population or spatial area. The index of deprivation and the assessment of population density are

described in more detail below. From Table 2 onwards, all variables were divided into seven categories, with wherever possible an equal number of census units for GB in each group. An arithmetic mean for each census measure was calculated by weighting the GB data by the age, sex and regional distribution of cases from the study. The proportions of persons in each of the seven categories from the census, also weighted by the study values were calculated.

**Deprivation** An index of deprivation was generated for each ED/OA in GB (UKCCS Investigators, 2000). The proportions of unemployed economically active persons (aged 16 and over), households without a car and households not owner-occupied were calculated. Each proportion was transformed to give zero skew with a mean of zero and an overall standard deviation of one. The transformed value for each census unit was calculated and is equivalent to the number of standard deviations from the overall mean. These standard deviations, from the three variables, were summed for each unit and the resulting value is termed the index of deprivation. A high value positive value represents a region with high material deprivation and a high negative value represents an affluent area.

**Population density** The measures of population density are reported at the electoral ward/postcode sector level. The traditional estimate of population density  $d_j$  for each geographical region  $j$  is given as

$$d_j = \frac{p_j}{a_j} \quad (1)$$

where  $p_j$  is the number of persons and  $a_j$  the spatial area. This will be referred to as the 'area-based' population density and is expressed as persons per hectare. However, this does not necessarily reflect the density at which the 'average' person lives, but more closely reflects the land use and type of region. A more useful measure for the density at which an average person lives is termed the 'person-weighted' population density (Dorling and Atkins, 1995). This is defined as

$$w_j = \sum_{i=1}^n d_i \left( \frac{p_i}{p_j} \right) \quad (2)$$

where region  $j$  is divided into  $n$  smaller regions  $i$ ,  $d_i$  is the area-based population density and  $p_i$  the number of persons for the smaller region. For the purposes of this analysis, this may be regarded as a population-weighted sum of the area-based population densities for the ED/OA in each electoral ward/postcode sector. In order to provide a more intuitive assessment of urban-

**Table 1** Description and summary of the census variables used in the analysis

Variable		Areal level <sup>a</sup>	GB weighted mean <sup>b</sup>	Min	Max
Index of deprivation	Composite index of unemployment, car and home ownership (see text)	ED/OA	0.17	-6.15	7.75
<i>Components of the deprivation index</i>					
Unemployed persons	Proportion economically-active persons unemployed	ED/OA	0.11	0.00	1.00
Houses with no car	Proportion of households without a car	ED/OA	0.33	0.00	1.00
Non-owner occupied houses	Proportion of households not owner-occupied	ED/OA	0.35	0.00	1.00
<i>Other census variables</i>					
Household overcrowding	Proportion households with more than one person per room	ED/OA	0.03	0.00	0.55
Households with child dependants	Proportion households with at least one dependant child	ED/OA	0.29	0.00	0.95
Flat dwellings	Proportion dwellings which are flats	ED/OA	0.18	0.00	1.00
Households with central heating	Proportion households with central heating	ED/OA	0.83	0.00	1.00
Persons working over 40 hours/week	Proportion employed persons working over 40 h per week	ED/OA	0.16	0.00	1.00
<i>Population density<sup>c</sup></i>					
Area-based	Area based population density (see text)	W/PC	28.09	0	712.26
Person-weighted	Person-weighted population density (see text)	W/PC	40.37	0	998.68

<sup>a</sup>ED/OA-enumeration district/output area; W/PC-ward/postcode area; <sup>b</sup>Weighted for case distribution for sex, 5-year age group (0-14 year-olds) and study region; <sup>c</sup>Based on persons per unit area.

**Table 2** Frequency (%) of interviewed cases and control groups in the UK Childhood Cancer Study by deprivation category<sup>a</sup>

Deprivation category <sup>a</sup>	Interviewed cases	Controls			GB <sup>b</sup> % distribution
		First-choice	First-choice non-interviewed	Total interviewed	
Least	532 (14)	1041 (14)	183 (9)	1154 (15)	13
	559 (16)	1090 (14)	204 (10)	1215 (16)	14
	591 (15)	1057 (14)	221 (10)	1160 (15)	14
	558 (15)	1109 (14)	260 (12)	1154 (15)	14
	504 (13)	1059 (14)	307 (14)	1038 (14)	14
	510 (13)	1097 (14)	384 (18)	990 (13)	14
Most	544 (14)	1162 (15)	526 (25)	918 (12)	17
Not linked <sup>c</sup>	0	54	54	0	
Total	3838 (100)	7669 (100)	2139 (100)	7629 (100)	100

<sup>a</sup>Deprivation for the enumeration district/output Area; <sup>b</sup>Weighted by case distribution for a 5-year age group, study region and sex; <sup>c</sup>Address not available to link census data.

rural status both measures of population density for all persons were categorized as urban (more than 25 persons per hectare), rural (less than 1.5 persons per hectare) or suburban (in between).

### Statistical methods

The risk associated with each explanatory exposure measure is presented as odds ratios from a logistic regression model. Adjustment was made for single year of age, sex and study region as a representation of the matching variables (UKCCS Investigators, 1999). Two comparison groups, not mutually exclusive, were used: the first-choice controls and the interviewed controls.

### RESULTS

Of the 4433 cases identified as eligible, permission to approach 4306 (97%) was obtained from the treating consultant, and 3838 (87%) agreed to participate (Table 2). Case interview rates varied by diagnostic group, from 93% for acute lymphoblastic leukaemia to 82% for the CNS tumours. Of the 7669 matched controls eligible for the study, 5530 (72%) agreed to participate; a lower proportion than for the cases. The proportion for control participation was broadly similar across the different diagnostic groups.

The distribution of interviewed cases and interviewed and first-choice controls by deprivation category is shown in Table 2. The frequency distribution of persons recorded on the census within each of the seven deprivation categories in GB, weighted for the case distribution, is shown on the far right-hand side. The first-choice control group was similarly distributed to that for the whole of GB, with a slightly lower proportion of controls from the most deprived group. The non-interviewed first-choice controls show a skew towards the more deprived groups, with 25% of those refusing to be interviewed coming from the most deprived seventh (where 15% would be expected). This resulted in the interviewed control group having fewer controls from the more deprived areas, with corresponding excesses in the more affluent areas. The distribution for interviewed cases was more similar to the first-choice controls, and to GB as a whole, than to the interviewed controls.

Table 3 shows the distribution of cases and controls for all chosen census characteristics. The results from a logistic model, adjusting for age, sex and study region are presented, comparing cases to interviewed controls (right hand columns) and to the first-choice controls (central columns). Overall, the trends for the two comparison groups – which are not mutually exclusive – appear to convey conflicting messages. Comparing cases and first-choice controls with respect to deprivation and

its component indices, there is little evidence of any consistent associations, with the possible exception of home ownership. However, comparison of interviewed controls with cases suggests elevated risks for childhood cancer associated with increasing material deprivation, unemployment, decreased car ownership and home ownership.

There were no discernable associations between cancer in children and the proportion of houses with child dependants. A non-significant protective effect of proportion of flat dwellers in an area was shown for the first-choice controls. The patterns of association are less clear for the measure of proportion of households with central heating and persons working more than 40 h per week, but the trends are contradictory between the two comparison groups.

Comparison with the interviewed control group, suggested no association between population density and childhood cancer. When making comparison with first-choice controls, no risk estimates for population density were observed to be significantly different from one. However, the point estimates suggested association between cancer and increasing rurality, as measured by both population density methods.

### DISCUSSION

Issues of participation are significant for all studies of disease aetiology that rely on individual compliance. The issue of non-participation, and consequential bias that may be introduced, is critically important in case-control studies that rely on personal contact to assess environmental experiences and exposures. Our findings indicate that in such studies, the profiling of non-participants may be as important as that of participants. In the study described here, the involvement of parents required active participation, as information was primarily collected through a face-to-face interview (UKCCS Investigators, 2000). As with all interview-based case-control studies, however, whilst the motivation of affected families to participate was strong, the motivation of those who were unaffected is less clear.

Comparisons of census data for cases and first-choice controls provided results that were free from participation bias: the findings suggesting that, on average, case families tended to live in areas that were more affluent than those of control families. In contrast, comparison with participating control families suggested that, on average, case families tended to live in areas that were less affluent than those of control families. This observation is consistent with other reports suggesting that participants often belong to a higher socio-economic groups than non-participants as measured by housing tenure, income, level of education and occupation (Holt

*et al*, 1991; Hatch *et al*, 1998; Madigan *et al*, 2000; Wrensch *et al*, 2000).

Measures of material deprivation are often closely associated with possible aetiological factors such as smoking, occupation, and previous illness history. Indeed, in many epidemiological studies it

is virtually impossible to identify potentially harmful exposures that are not – either directly or indirectly – related to measures of social class, deprivation or affluence. Further, somewhat surprisingly, the treatment of bias as a real confounder (risk factor) rarely impacts on the magnitude of risk estimates (e.g. UKCCS Investigators,

**Table 3** Odds ratios (95% CI) of the census characteristics or residential area, from logistic regression<sup>a</sup> using the first-choice and interviewed controls

	GB <sup>b</sup>	Cases	First-choice controls		Interviewed controls	
		n (%)	n (%)	OR (95%CI) <sup>a</sup>	n (%)	OR (95%CI) <sup>a</sup>
<i>Deprivation</i>						
Least	13	532 (14)	1041 (14)	1 (–)	1154 (15)	1 (–)
	14	599 (16)	1090 (14)	1.08 (0.94–1.25)	1215 (16)	1.07 (0.93–1.23)
	14	591 (15)	1057 (14)	1.12 (0.97–1.29)	1160 (15)	1.11 (0.96–1.28)
	14	558 (15)	1109 (14)	1.00 (0.86–1.16)	1154 (15)	1.05 (0.91–1.21)
	14	504 (13)	1059 (14)	0.94 (0.81–1.10)	1038 (14)	1.06 (0.91–1.22)
	14	510 (13)	1097 (14)	0.94 (0.81–1.09)	990 (13)	1.12 (0.97–1.30)
Most	17	544 (14)	1162 (15)	0.95 (0.82–1.10)	918 (12)	1.30 (1.12–1.51)
<i>Components of the deprivation index</i>						
<i>Unemployment</i>						
Least	12	503 (13)	910 (12)	1 (–)	1041 (14)	1 (–)
	15	593 (15)	1192 (16)	0.91 (0.78–1.05)	1297 (17)	0.94 (0.82–1.09)
	14	580 (15)	1158 (15)	0.92 (0.79–1.07)	1247 (16)	0.96 (0.83–1.11)
	14	564 (15)	1041 (14)	1.00 (0.86–1.16)	1094 (14)	1.07 (0.92–1.24)
	14	531 (14)	1056 (14)	0.93 (0.80–1.08)	1043 (14)	1.05 (0.91–1.22)
	15	518 (14)	1080 (14)	0.90 (0.77–1.04)	956 (13)	1.12 (0.97–1.31)
Most	17	549 (14)	1178 (15)	0.87 (0.75–1.02)	951 (12)	1.20 (1.04–1.40)
<i>Car ownership</i>						
Most	15	616 (16)	1154 (15)	1 (–)	1281 (17)	1 (–)
	15	610 (16)	1120 (15)	1.02 (0.89–1.18)	1223 (16)	1.04 (0.90–1.19)
	15	595 (16)	1177 (15)	0.96 (0.83–1.10)	1274 (17)	0.97 (0.85–1.11)
	15	570 (15)	1115 (15)	0.97 (0.84–1.12)	1157 (15)	1.03 (0.89–1.18)
	14	551 (14)	1145 (15)	0.91 (0.79–1.05)	1057 (14)	1.09 (0.94–1.25)
	14	472 (12)	1041 (14)	0.87 (0.75–1.01)	923 (12)	1.07 (0.93–1.25)
Least	12	424 (11)	863 (11)	0.95 (0.81–1.11)	714 (9)	1.26 (1.07–1.47)
<i>Home ownership</i>						
Most	16	661 (17)	1230 (16)	1 (–)	1413 (19)	1 (–)
	15	659 (17)	1223 (16)	1.01 (0.89–1.16)	1327 (17)	1.06 (0.93–1.21)
	14	546 (14)	1111 (14)	0.93 (0.81–1.08)	1135 (15)	1.03 (0.90–1.18)
	14	535 (14)	1053 (14)	0.96 (0.83–1.11)	1052 (14)	1.09 (0.95–1.25)
	14	511 (13)	1007 (13)	0.96 (0.83–1.11)	995 (13)	1.10 (0.95–1.27)
	14	513 (13)	1009 (13)	0.97 (0.84–1.11)	898 (12)	1.22 (1.06–1.41)
Least	13	413 (11)	982 (14)	0.80 (0.69–0.93)	809 (11)	1.09 (0.94–1.27)
<i>Other measures</i>						
<i>Household overcrowding</i>						
Least	18	788 (21)	1463 (19)	1 (–)	1587 (21)	1 (–)
	15	629 (16)	1114 (15)	1.06 (0.93–1.21)	1242 (16)	1.02 (0.89–1.16)
	14	552 (14)	1094 (14)	0.95 (0.83–1.09)	1118 (15)	0.99 (0.87–1.14)
	13	485 (13)	991 (13)	0.92 (0.80–1.06)	989 (13)	0.99 (0.86–1.13)
	13	491 (13)	1003 (13)	0.93 (0.81–1.06)	980 (13)	1.01 (0.88–1.16)
	13	454 (12)	993 (13)	0.86 (0.75–1.00)	956 (13)	0.96 (0.83–1.10)
Most	13	439 (11)	957 (12)	0.87 (0.75–1.00)	757 (10)	1.17 (1.01–1.36)
<i>Households with child dependants</i>						
Least	5	198 (5)	416 (5)	1 (–)	424 (6)	1 (–)
	10	443 (12)	800 (10)	1.16 (0.95–1.43)	806 (11)	1.17 (0.96–1.45)
	12	485 (13)	908 (12)	1.12 (0.92–1.38)	912 (12)	1.14 (0.93–1.39)
	15	541 (14)	1163 (15)	0.98 (0.80–1.19)	1214 (16)	0.95 (0.78–1.16)
	17	621 (16)	1263 (16)	1.03 (0.85–1.26)	1261 (17)	1.05 (0.87–1.28)
	19	719 (19)	1380 (18)	1.10 (0.90–1.33)	1399 (18)	1.10 (0.91–1.33)
Most	23	831 (22)	1685 (22)	1.04 (0.86–1.25)	1613 (21)	1.03 (0.91–1.33)
<i>Flat dwellings</i>						
Least	21	851 (22)	1595 (21)	1 (–)	1667 (22)	1 (–)
	17	620 (16)	1264 (16)	0.92 (0.81–1.05)	1313 (17)	0.92 (0.81–1.05)
	15	588 (15)	1179 (15)	0.93 (0.82–1.06)	1213 (16)	0.95 (0.83–1.08)
	16	575 (15)	1139 (15)	0.94 (0.83–1.08)	1154 (15)	0.97 (0.85–1.11)
	14	550 (14)	1093 (14)	0.94 (0.82–1.07)	1062 (14)	1.01 (0.89–1.16)
	11	418 (11)	838 (11)	0.93 (0.81–1.08)	793 (10)	1.04 (0.89–1.20)
Most	7	236 (6)	507 (7)	0.87 (0.72–1.04)	427 (6)	1.10 (0.91–1.32)

*Continued*

Table 3 (Continued)

	GB% <sup>b</sup>	Cases		First-choice controls		Interviewed controls	
		n (%)	n (%)	OR (95%CI) <sup>a</sup>	n (%)	OR (95%CI) <sup>a</sup>	
Households with central heating							
Most	15	624 (16)	1182 (15)	1 (-)	1209 (16)	1 (-)	
	16	600 (16)	1183 (15)	0.96 (0.83-1.10)	1190 (16)	0.98 (0.85-1.12)	
	15	581 (15)	1106 (15)	0.99 (0.86-1.14)	1194 (16)	0.94 (0.82-1.08)	
	14	564 (15)	1057 (14)	1.10 (0.87-1.16)	1087 (14)	1.00 (0.87-1.16)	
	13	513 (13)	1027 (13)	0.94 (0.81-1.09)	1045 (14)	0.95 (0.82-1.10)	
	13	455 (12)	1038 (14)	0.83 (0.71-0.96)	1000 (13)	0.88 (0.76-1.02)	
Least	14	501 (13)	1022 (13)	0.92 (0.80-1.09)	904 (12)	1.07 (0.93-1.25)	
Persons working over 40 h per week							
Most	9	391 (10)	730 (10)	1 (-)	752 (10)	1 (-)	
	15	536 (14)	1067 (14)	0.93 (0.79-1.09)	1103 (14)	0.94 (0.80-1.10)	
	13	500 (13)	1015 (13)	0.93 (0.79-1.10)	1025 (13)	0.92 (0.78-1.08)	
	17	683 (18)	1326 (17)	0.99 (0.85-1.16)	1316 (17)	0.96 (0.82-1.12)	
	16	648 (17)	1228 (16)	0.97 (0.83-1.14)	1275 (17)	0.99 (0.84-1.15)	
	18	629 (16)	1324 (17)	0.91 (0.78-1.06)	1322 (17)	0.89 (0.76-1.04)	
Least	11	440 (11)	894 (12)	1.04 (0.87-1.24)	821 (11)	0.92 (0.77-1.09)	
Data n/a		11 (0)	31 (0)		15 (0)		
Not linked <sup>c</sup>		0	54		0		
Population density							
Area-based							
Urban	47	1662 (43)	3437 (45)	1 (-)	3311 (43)	1 (-)	
Surburban	42	1693 (44)	3268 (43)	1.07 (0.94-1.22)	3345 (44)	1.01 (0.93-1.10)	
Rural	11	483 (13)	919 (12)	1.07 (0.94-1.22)	973 (13)	0.99 (0.87-1.12)	
Person weighted							
Urban	69	2542 (66)	5173 (68)	1 (-)	5048 (66)	1 (-)	
Surburban	26	1069 (28)	2046 (27)	1.05 (0.96-1.15)	2142 (28)	0.99 (0.91-1.08)	
Rural	5	227 (6)	405 (5)	1.12 (0.95-1.33)	439 (6)	1.03 (0.87-1.22)	

<sup>a</sup>Odds ratio and 95% confidence interval adjusted for a single year of age, sex and study region; <sup>b</sup>Weighted for case distribution for sex, 5-year age group and study region;

<sup>c</sup>Address not available to link to census data.

1999; 2001). The challenge is to disentangle the artifactual consequences of participation bias from genuine aetiological factors. Some researchers have suggested using the variable most closely related to participation as a confounder to 'adjust' for participation. Such adjustments are, however, only appropriate when the antecedents of both exposure and disease, or their joint distributions, are available for the entire study population (Greenland, 1998). As in the majority of studies, neither of these were available for the UKCCS, where individual reasons for participation could only be indirectly related with the deprivation index.

As expected, our findings show that the primary care sampling frame provided control families that were broadly representative of the general GB population, where the majority of people register with a GP in the area in which they live – access to NHS medical care requiring registration which covers approximately 98% of the population (RCGP, 1987). The UKCCS sampling frame compares favourably with other methods of control selection. For example, random digit dialling, a method commonly employed in the USA and Canada (Robison and Daigle, 1984), prohibits collection from homes without a telephone, those not at home when telephoned, and those who answer but who either refuse to answer any questions or deliberately lie about their families eligibility.

In conclusion, our findings confirm that differential participation is a potentially major source of bias in case-control studies that estimate risks on the basis of information reported from respondents alone. Studies that ignore this source of bias may produce misleading results. There is a clear need to address this

issue in terms of study design, and in the application of appropriate statistical methods to try to overcome this bias.

## ACKNOWLEDGEMENTS

The UKCCS was sponsored and administered by the United Kingdom Co-ordinating Committee on Cancer Research and was supported by the UKCCS paediatric oncologists and by the National Radiological Protection Board. Financial support was provided by: Cancer Research Campaign, Imperial Cancer Research Fund, Leukaemia Research Fund, and Medical Research Council through grants to their units; Leukaemia Research Fund, Department of Health, member companies of Electricity Association, Irish Electricity Supply Board, National Grid Company plc, and Westlakes Research (Trading) Ltd for general expenses of the study; Kay Kendall Leukaemia Fund for associated laboratories studies; and Foundation of Children with Leukaemia for study of electrical fields. The investigation in Scotland is funded by The Scottish Office, Scottish Power plc, Scottish Hydro-Electric plc, and Scottish Nuclear Ltd. We thank the members of the UKCCSG for their support, the staff of the local hospitals, the family physicians and their practice staff. We especially thank the families of the children included in the study for their help.

## REFERENCES

- Dorling D, Atkins D (1995) *Population density, change and concentration in Great Britain 1971, 1981 and 1991*. OPCS Series SMPS no 58. London: HMSO
- Greenland S (1998) Basic methods for sensitivity analysis and external adjustment. In *Modern Epidemiology*, 2nd edn, Rothman KJ, Greenland S (eds) pp 115–134. Philadelphia: Lippincott-Raven
- Hartge P (1999) Raising response rates: Getting to yes. *Epidemiology* **10**: 105–107
- Hatch EE, Linet MS, Kleinerman RA, Tarone RE, Severson RK, Hartsock CT, Haines C, Kaune WT, Friedman D, Robison LL, Wacholder S (1998) Association between childhood acute lymphoblastic leukaemia and use of electrical appliances during pregnancy and childhood. *Epidemiology* **9**: 234–245
- Holt VL, Daling JR, Stergachis A, Voigt LF, Weiss NS (1991) Results and effect of refusal recontact in a case-control study of ectopic pregnancy. *Epidemiology* **2**: 375–379
- Madigan MP, Troisi R, Potischman N, Brogan D, Gammon MD, Malone KE, Brinton LA (2000) Characteristics of respondents and non-respondents from a case-control study of breast cancer in younger women. *International J Epidemiol* **29**: 793–798
- RCGP (1987) *Profile of UK practices*. RCGP Information Sheet No 2. London: Royal College of General Practitioners
- Robison LL, Daigle A (1984) Control selection using random digit dialling for cases of childhood cancer. *Am J Epidemiol* **120**: 164–166
- Rothman KJ, Greenland S (1998) Precision and validity in epidemiologic studies. In *Modern Epidemiology*, 2nd edn, Rothman KJ, Greenland S (eds) pp 115–134. Philadelphia: Lippincott-Raven
- UKCCS Investigators (1999) Exposure to power-frequency magnetic fields and the risk of childhood cancer. *Lancet* **354**: 1925–1931
- UKCCS Investigators (2000) The United Kingdom Childhood Cancer Study: objectives, materials and methods. *Br J Cancer* **82**: 1073–1102
- UKCCS Investigators (2001) Breast feeding and childhood cancer. *Br J Cancer* **11**: 1685–1694
- Wacholder S, Silverman DT, McLaughlin JK, Mandel S (1992) Selection of controls in case-control studies: II Types of controls. *Am J Epidemiol* **135**: 1029–1041
- Wrensch M, Miike R, Lee M, Neuhaus J (2000) Are prior head injuries or diagnostic X-rays associated with glioma in adults? The effects of control selection bias. *Neuroepidemiology* **19**: 234–244

## APPENDIX 1

*Writing Committee* – GR Law, AG Smith, E Roman

*Management Committee* – KK Cheng, Central Region; N Day, East Anglia Region; R Cartwright, A Craft, North East Region; JM Birch, OB Eden, North West Region; PA McKinney, Scotland; J Peto, South East Region; V Beral, E Roman, South Midlands Region; P Elwood, South Wales Region; FE Alexander, South West Region; CED Chilvers, Trent Region; R Doll, Epidemiological Studies Unit, University of Oxford; GM Taylor, Immunogenetics Laboratory, University of Manchester, Manchester; M Greaves, Leukaemia Research Fund Centre, Institute of Cancer Research; D Goodhead, Radiation and Genome Stability Unit, Medical Research Council, Harwell; FA Fry, National Radiological Protection Board; G Adams, UK Co-ordinating Committee for Cancer Research.

*Regional Investigators* – KK Cheng, E Gilman, Central Region; N Day, J Skinner, D Williams, East Anglia Region; R Cartwright, A Craft, North

East Region; JM Birch, OB Eden, North West Region; PA McKinney, Scotland; J Deacon, J Peto, South East Region; V Beral, E Roman, South Midlands Region; P Elwood, South Wales Region; FE Alexander, M Mott, South West Region; CED Chilvers, K Muir, Trent Region.

*Data Processing Group* – GR Law, J Simpson, E Roman.

A complete list of investigators is given in: The United Kingdom Childhood Cancer Study: objectives, materials, and methods. *Br J Cancer* 2000; **82**: 1073–1102.