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Decline in mortality, AIDS, and hospital admissions in perinatally HIV-1 infected children in the United Kingdom and Ireland

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Abstract

Objective To describe changes in demographic factors, disease progression, hospital admissions, and use of antiretroviral therapy in children with HIV.

Design Active surveillance through the national study of HIV in pregnancy and childhood (NSHPC) and additional data from a subset of children in the collaborative HIV paediatric study (CHIPS).

Setting United Kingdom and Ireland.

Participants 944 children with perinatally acquired HIV-1 under clinical care.

Main outcome measures Changes over time in progression to AIDS and death, hospital admission rates, and use of antiretroviral therapy.

Results 944 children with perinatally acquired HIV were reported in the United Kingdom and Ireland by October 2002; 628 (67%) were black African, 205 (22%) were aged ≥10 years at last follow up, 193 (20%) are known to have died. The proportion of children presenting who were born abroad increased from 20% in 1994-5 to 60% during 2000-2. Mortality was stable before 1997 at 9.3 per 100 child years at risk but fell to 2.0 in 2001-2 (trend P < 0.001). Progression to AIDS also declined (P < 0.001). From 1997 onwards the proportion of children on three or four drug antiretroviral therapy increased. Hospital admission rates declined by 80%, but with more children in follow up the absolute number of admissions fell by only 26%.

Conclusion In children with HIV infection, mortality, AIDS, and hospital admission rates have declined substantially since the introduction of three or four drug antiretroviral therapy in 1997. As infected children in the United Kingdom and Ireland are living longer, there is an increasing need to address their medical, social, and psychological needs as they enter adolescence and adult life.

Introduction

Since 1996 antiretroviral therapy with three or four drugs has been increasingly used to treat adults infected with HIV, and its substantial effect on progression of the disease has been well described. Because of difficulties in developing appropriate formulations and the lack of age specific pharmacokinetic data to guide paediatric dosing the introduction of such treatment was delayed in children. Before 1994 only one antiretroviral drug (zidovudine) was available for children. Dual antiretroviral therapy was introduced in 1995, and treatment with three or four drugs followed from mid-1997 and is now standard. Reductions in mortality and AIDS during the era of dual therapy were reported, and substantial declines attributed to therapy with three or four drugs were reported in a US paediatric cohort. However, data on changing rates in disease progression since 1997 and their effect on use of medical services remain sparse in children. We investigated changes over time in mortality, morbidity, hospital admission rates, and antiretroviral therapy received in perinatally HIV-1 infected children in the United Kingdom and Ireland.

Methods

Study design

The national study of HIV in pregnancy and childhood (NSHPC) is informed about children who present in the United Kingdom and Ireland with HIV-1 infection and infants born to HIV infected women through two active confidential reporting schemes run in collaboration with the British Paediatric Surveillance Unit and the Royal College of Obstetricians and Gynaecologists. Obstetric data include demographic information, timing of maternal diagnosis, uptake of interventions, and outcome of pregnancy; paediatric data include neonatal details and HIV status. Each year notifying paediatricians subsequently provide minimal clinical information for infected children.

In April 2000, the collaborative HIV paediatric study (CHIPS) was established between the national surveillance study at the Institute of Child Health, London, and the Medical Research Council Clinical Trials Unit, where trials in the Paediatric European Network for Treatment of AIDS (PENTA) are coordinated (www.pentatrials.org). Most perinatally infected children reported to the national surveillance...
study are followed up in 17 paediatric clinics involved in PENTA. The collaborative HIV paediatric study was established to collect more detailed clinical, laboratory, and treatment information on infected children seen at the PENTA centres since 1996. During 2000, retrospective information was collected from clinical records on to standard forms; since then, information has been collected annually by questionnaire and merged with additional data from the national surveillance study and PENTA trials. The merged dataset includes demographic information, data on clinical events, hospital admissions, outpatient visits, antiretroviral therapy, and results of T cell subset and HIV RNA viral load tests. AIDS was defined according to the classification of the Centers for Disease Control (clinical category C disease).15

Paediatric HIV surveillance started in 1986, and by October 2002, 944 perinatally infected children had been reported to the national surveillance study. All children were included in analyses of changes in progression to AIDS and death over time. In total 593 children alive in January 1996 were enrolled in the collaborative HIV paediatric study (75% of all children reported to the national surveillance study as alive on this date) and were included in analyses of antiretroviral therapy and hospital admissions.

Statistical methods
For children whose mothers’ HIV-1 status was known before delivery, we considered the date of birth to be the date of the child’s first presentation to medical services with HIV-1. For remaining children, we estimated the date of presentation using the earliest date of: first positive results of HIV-1 antibody test, T cell subsets, or viral load tests; HIV related clinical events; clinic visit; or hospital admission. In all analyses, each child began to contribute to follow up from his or her date of presentation.

We censored records at the date children were last known to be alive in the death analysis and at the date of last clinical assessment in the AIDS/death analysis. To reduce presentation bias, we excluded children not identified at birth from analyses of AIDS or death if they first presented within one month of AIDS diagnosis or death, respectively. The effect of calendar period was further examined by using Cox proportional hazards models, adjusted for how the children were identified (at birth or later), ethnicity, sex, and place of birth (United Kingdom and Ireland or abroad). We also carried out separate adjusted analyses by place of birth. Time was measured from birth by using late entry at the age of first presentation.16 In any calendar period, children were considered at risk only from the first day

Table 1 Characteristics of perinatally HIV-1 infected children in the United Kingdom and Ireland up to October 2002 according to place of birth. Figures are numbers (percentages) of children

<table>
<thead>
<tr>
<th></th>
<th>Born in UK or Ireland (n=586)</th>
<th>Born abroad (n=353)</th>
<th>Unknown (n=5)</th>
<th>Total (n=944)</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Female</strong></td>
<td>280 (48)</td>
<td>169 (52)</td>
<td>3</td>
<td>468 (50)</td>
</tr>
<tr>
<td><strong>Ethnicity</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>White</td>
<td>148 (25)</td>
<td>13 (4)</td>
<td>0</td>
<td>161 (17)</td>
</tr>
<tr>
<td>Black</td>
<td>347 (59)</td>
<td>278 (79)</td>
<td>3</td>
<td>628 (67)</td>
</tr>
<tr>
<td>Other</td>
<td>88 (15)</td>
<td>61 (17)</td>
<td>1</td>
<td>150 (16)</td>
</tr>
<tr>
<td>Unknown</td>
<td>3 (1)</td>
<td>1 (0.3)</td>
<td>1</td>
<td>5 (1)</td>
</tr>
<tr>
<td><strong>Region of follow up</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>London</td>
<td>416 (71)</td>
<td>248 (70)</td>
<td>4</td>
<td>668 (71)</td>
</tr>
<tr>
<td>Rest of England</td>
<td>92 (16)</td>
<td>73 (20)</td>
<td>0</td>
<td>165 (17)</td>
</tr>
<tr>
<td>Scotland</td>
<td>36 (6)</td>
<td>8 (2)</td>
<td>0</td>
<td>44 (5)</td>
</tr>
<tr>
<td>Ireland</td>
<td>35 (6)</td>
<td>18 (5)</td>
<td>1</td>
<td>54 (6)</td>
</tr>
<tr>
<td>Wales and N Ireland</td>
<td>7 (1)</td>
<td>6 (2)</td>
<td>0</td>
<td>13 (2)</td>
</tr>
<tr>
<td><strong>How child was identified</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Prospectively from birth</td>
<td>121 (21)</td>
<td>11 (3)</td>
<td>0</td>
<td>132 (32)</td>
</tr>
<tr>
<td>After birth, asymptomatic</td>
<td>157 (27)</td>
<td>139 (39)</td>
<td>2</td>
<td>298 (14)</td>
</tr>
<tr>
<td>After birth, symptomatic</td>
<td>257 (47)</td>
<td>180 (51)</td>
<td>2</td>
<td>469 (50)</td>
</tr>
<tr>
<td>Unknown</td>
<td>21 (4)</td>
<td>23 (7)</td>
<td>1</td>
<td>45 (5)</td>
</tr>
<tr>
<td><strong>Age at first presentation (years)</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>At birth</td>
<td>121 (21)</td>
<td>11 (3)</td>
<td>0</td>
<td>132 (14)</td>
</tr>
<tr>
<td>&lt;1</td>
<td>254 (43)</td>
<td>27 (8)</td>
<td>0</td>
<td>281 (30)</td>
</tr>
<tr>
<td>1</td>
<td>72 (12)</td>
<td>38 (11)</td>
<td>0</td>
<td>110 (12)</td>
</tr>
<tr>
<td>2-4</td>
<td>104 (18)</td>
<td>113 (32)</td>
<td>2</td>
<td>221 (23)</td>
</tr>
<tr>
<td>5-9</td>
<td>25 (4)</td>
<td>123 (35)</td>
<td>2</td>
<td>154 (16)</td>
</tr>
<tr>
<td>≥10</td>
<td>6 (1)</td>
<td>41 (12)</td>
<td>1</td>
<td>48 (5)</td>
</tr>
<tr>
<td>Median (range)</td>
<td>0.46 (0.0-13.4)</td>
<td>4.6 (0.2-15.3)</td>
<td>5.0 (2.7-12.8)</td>
<td>1.6 (0.0-15.3)</td>
</tr>
<tr>
<td><strong>Age at last follow up (years)</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>&lt;1</td>
<td>104 (18)</td>
<td>11 (3)</td>
<td>0</td>
<td>115 (12)</td>
</tr>
<tr>
<td>1-4</td>
<td>203 (35)</td>
<td>99 (28)</td>
<td>2</td>
<td>304 (32)</td>
</tr>
<tr>
<td>5-9</td>
<td>158 (24)</td>
<td>121 (34)</td>
<td>1</td>
<td>300 (34)</td>
</tr>
<tr>
<td>10-14</td>
<td>67 (11)</td>
<td>97 (27)</td>
<td>2</td>
<td>166 (18)</td>
</tr>
<tr>
<td>≥15</td>
<td>14 (2)</td>
<td>25 (7)</td>
<td>0</td>
<td>39 (4)</td>
</tr>
<tr>
<td>Median (range)</td>
<td>4.8 (0.0-18.1)</td>
<td>7.3 (0.3-18.6)</td>
<td>8.0 (3.1-14.7)</td>
<td>5.7 (0.0-18.6)</td>
</tr>
<tr>
<td><strong>Stage of disease at last follow up</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>TB/HIV</td>
<td>275 (47)</td>
<td>228 (64)</td>
<td>5</td>
<td>506 (54)</td>
</tr>
<tr>
<td>C</td>
<td>159 (27)</td>
<td>86 (24)</td>
<td>0</td>
<td>245 (26)</td>
</tr>
<tr>
<td>Died</td>
<td>153 (26)</td>
<td>41 (12)</td>
<td>0</td>
<td>193 (20)</td>
</tr>
</tbody>
</table>

*According to classification of Centers for Disease Control.13
of the calendar period or the date of presentation, if later. We fitted calendar period as a time varying covariate, with adjustment for an interaction with age.

We used Kaplan-Meier curves to compare survival after initial diagnosis of AIDS by first indicator disease, grouped in hierarchical order: *Pneumocystis carinii* pneumonia, other opportunistic infection, HIV encephalopathy, failure to thrive, severe recurrent bacterial infections. We excluded six children in whom cancer was the first indicator disease. Hospital admission rates were compared with Poisson regression.

Results

Half of the 944 children reported were girls and 628 (67%) were black African (table 1). The proportion of children born abroad increased from 20% in 1994-5 to 60% in 2000-2. In total 132 (14%) children were identified at birth, 281 (30%) during the first year, and 202 (21%) at ≥5 years. Whereas the median age at presentation among children born in the United Kingdom and Ireland has remained constant at around 6 months, among children born abroad it increased from 2.5 years before 1991 to 5.2 years in 1994-5 and remained constant thereafter. At last follow up, 205 (22%) children were aged ≥10 years and 39 (4%) ≥15 years (table 1). A total of 195 children are known to have died (including 54 since 1997); and 41 left the country or were otherwise lost to follow up before 1996. Among 593 children in the collaborative HIV paediatric study, 534 (90%) were in follow up in 2001-2, 30 (5%) in 2000, and 29 (5%) were last seen in 1999 or earlier.

Mortality and progression to AIDS

While the number of perinatally infected children in follow up has increased steadily over time, the annual number of deaths declined markedly after 1996 (table 2). Among 734 children identified at birth, or who survived at least a month after presentation, the crude mortality was stable before 1997 at 9.3 per 100 child years but declined thereafter to 3.0 per 100 child years at the 95% confidence interval 0.12 to 0.30 than in those aged >1 year (0.72, 0.35 to 1.48; P = 0.03, heterogeneity for age) (table 3). Adjusted hazard ratios were similar when we carried out separate analyses for children aged >1 year born in the United Kingdom and Ireland (0.18, 0.11 to 0.31) and born abroad (0.20, 0.08 to 0.46).

Fifty four children are known to have died from 1997 onwards, 48 (89%) of whom were born to mothers whose HIV status was not known during pregnancy. Eighteen of 22 children who died aged <1 year had *P carinii* pneumonia or cytomegalovirus disease, or both. Information on antiretroviral therapy was available for 36 children who died; 14 (including six infants) had not received three or four drug therapy, six died within two months of starting it, and 16 died a median of 10 (range 2-53) months after starting it.

A total of 438 children developed an AIDS indicator disease, but 13 of these died before AIDS was diagnosed. Two hundred and seventeen children (50%) developed AIDS within one month of presentation (table 2). Of the children with AIDS, 119 (28%) had more than one indicator disease for AIDS at diagnosis. The proportion of children with opportunistic infections was higher in 1997-2002 than previously (57% versus 29% for *P carinii* pneumonia; 57% versus 22% for other opportunistic infections), but the distribution of other indicator diseases was similar in the two periods. Prognosis from initial AIDS diagnosis varied: mortality was higher in children with *P carinii* pneumonia or HIV encephalopathy compared with mortality in children with other opportunistic infections, failure to thrive, or severe recurrent bacterial infection (fig 2).

Among 587 children who had not progressed to AIDS within one month of presentation after birth, crude progression rates declined from 15.4 per 100 child years before 1997 to 3.0 per 100 child years at
risk in 2001-2 (test for trend P < 0.001). Figure 1 shows the adjusted risks relative to 1996. As with mortality, this decline was more marked in children aged > 1 year (hazard ratio 0.37, 95% confidence interval 0.25 to 0.53, for before 1997 compared with 1997 onwards, table 3), while progression rates were similar in both periods for children aged < 1 (1.03, 0.60 to 1.77, P = 0.002, heterogeneity for age).

Antiretroviral therapy
Of 593 children followed since 1996 in the collaborative HIV paediatric study, 137 (23%) had not started antiretroviral therapy at last report (121) or by the time of death (16). The percentage of child time spent on three or four drug antiretroviral therapy increased from 1% in 1996 to 56% in 1999 and 69% in 2001-2 (fig 3). There was no evidence that this differed in children aged < 1. There was a shift over time from initial drug regimens containing a protease inhibitor to those containing a non-nucleoside reverse transcriptase inhibitor (table 4). The proportion of children who had previously been on therapy but had currently stopped remained constant after 1996 at around 5% of child time (fig 3).

Among 371 children on antiretroviral therapy when they were last seen in 2000 or later, 48 (13%) were taking four drugs, 301 (81%) three, and 22 (6%) two. Overall 91 different drug combinations were used.

Discussion
Mortality
From 1997 onwards we have seen reductions of around 80% in mortality and 50% in progression to AIDS among children perinatally infected with HIV-1 in the United Kingdom and Ireland. In the collaborative HIV paediatric study, hospital admission rates fell substantially. The introduction and increased use of antiretroviral therapy may explain these reductions.

Hospital admissions
The number of children seen in collaborative HIV paediatric study centres increased from 299 during 1996 to 493 in 2001-2, while the number of hospital admissions fell by 26% from 350 to 258. Admission rates declined by 80% from 4.4 per 100 child years of follow up in 1996 to 0.9 in 2001-2 (test for trend P < 0.001). As expected, the rates were lower in children born to mothers whose HIV status was known in pregnancy compared with those in whom it was not (relative risk 0.49, 0.42 to 0.57, P < 0.001). Among those presenting after birth, rates of admission were higher in the first six months after presentation than later (4.21, 3.79 to 4.69, P < 0.001).

Table 3
Adjusted hazard ratios (95% confidence intervals) for factors associated with progression to death and AIDS/death

<table>
<thead>
<tr>
<th>Calendar year:</th>
<th>Death</th>
<th>P value</th>
<th>AIDS/death</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Before 1997</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
</tr>
<tr>
<td>1997-2002</td>
<td>0.26 (0.17 to 0.38)</td>
<td>P=0.001</td>
<td>0.49 (0.36 to 0.67)</td>
<td>P=0.001</td>
</tr>
<tr>
<td>1997-2002, age &lt;1 year</td>
<td>0.72 (0.56 to 1.14)</td>
<td>P=0.005</td>
<td>1.03 (0.60 to 1.77)</td>
<td>P=0.002</td>
</tr>
<tr>
<td>1997-2002, age ≥1 year</td>
<td>0.19 (0.13 to 0.30)</td>
<td>P=0.002*</td>
<td>0.37 (0.25 to 0.53)</td>
<td>P=0.002*</td>
</tr>
<tr>
<td>Sex:</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
</tr>
<tr>
<td>Female</td>
<td>0.84 (0.61 to 1.15)</td>
<td>P=0.3</td>
<td>1.06 (0.81 to 1.40)</td>
<td>P=0.7</td>
</tr>
<tr>
<td>Ethnicity:</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>White</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
</tr>
<tr>
<td>Black</td>
<td>0.67 (0.48 to 1.27)</td>
<td>P=0.5</td>
<td>0.77 (0.55 to 1.08)</td>
<td>P=0.5</td>
</tr>
<tr>
<td>Other or not known</td>
<td>1.42 (0.65 to 2.37)</td>
<td>P=0.1</td>
<td>1.09 (0.69 to 1.72)</td>
<td>P=0.1</td>
</tr>
<tr>
<td>How child was identified:</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Prospectively from birth</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
</tr>
<tr>
<td>After birth</td>
<td>0.49 (0.31 to 0.78)</td>
<td>P=0.001</td>
<td>0.44 (0.30 to 0.64)</td>
<td>P=0.001</td>
</tr>
<tr>
<td>Place of birth:</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>UK or Ireland</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
</tr>
<tr>
<td>Abroad</td>
<td>0.96 (0.62 to 1.48)</td>
<td>P=0.9</td>
<td>1.06 (0.73 to 1.53)</td>
<td>P=0.8</td>
</tr>
</tbody>
</table>

*Test of heterogeneity for age.
uptake of three or four drug antiretroviral therapy accompanied these changes and is likely to be the major contributing factor. Similar findings, concurrent with the widespread uptake of antiretroviral therapy, have been reported from cohorts of children in the United States and Italy. Follow up in these studies was only to 1999, but we observed that survival stabilised during 2000-2 after dramatic reductions during 1997-9. A similar pattern has been observed in adult cohorts.

There are clearly limitations in using cohort studies to ascribe changes in patterns of disease to particular interventions. As in the Italian study we allowed for late entry at first presentation to reduce bias due to children with a good prognosis surviving longer and being more likely to be enrolled. In addition we repeated analyses separately for children born in the United Kingdom and Ireland or born abroad and observed similar results; this is particularly important in our cohort with the increasing proportion of children born abroad in recent years. As we included all children ever reported in the United Kingdom and Ireland, biases related to referral patterns to specialist centres are not an issue.

AIDS and hospital admissions

Our study is among the first to report reductions in hospital admissions over time in children with HIV, paralleling reductions in mortality and morbidity. The pattern of AIDS diagnoses before and since 1997 were similar, except for a trend towards a higher proportion of opportunistic infections in the later period. Improvements in the uptake of antenatal HIV testing and interventions to prevent transmission from mother to child have substantially increased the proportion of women diagnosed before delivery (from 32% in 1997 to 77% in 2001 in the United Kingdom) and led to a corresponding decrease in the proportion of infected infants. As expected, mortality, progression to AIDS, and hospital admission rates were lower in infected children born to mothers whose diagnosis was known before the child was born. Caution must be exercised in interpreting these results to suggest that antenatal testing has benefits beyond prevention of transmission by improving the outcome of infected children; presentation bias could play a part as asymptomatic infected children born to undiagnosed women will not be identified until they present with symptoms. Despite the marked reduction in hospital admission rates, total admissions have declined by only one quarter since 1996, underlying the need for continued inpatient as well as outpatient services for HIV infected children.

Infants compared with older children

Whereas in children aged over 1 year under clinical care mortality decreased by 81% and progression to AIDS or death decreased by 63%, we saw only modest improvements among infants. This may reflect a decrease in the proportion of infants in the cohort followed from birth, due to the reduction in mother to child transmission rates, although we attempted to control for this. Other explanations include the possibility that three or four drug antiretroviral therapy is not started early enough in infants. Pharmacokinetic issues may also hamper the effectiveness of antiretroviral therapy in this age group. In recent years, most infants presented with symptoms as they had been born to mothers undiagnosed at delivery. As reported here and previously, some infants die from P. carinii pneumonia or cytomegalovirus disease, or both, before antiretroviral therapy can be started. The outlook for infants infected despite maternal diagnosis in pregnancy cannot easily be deduced from this study as numbers are small. However, we can speculate that transmission of maternal viruses that are drug resistant or more virulent may be also important. Ethnicity, sex, and place of birth had no significant effect on outcome.

Antiretroviral therapy

We observed a change over time from first use of antiretroviral therapy regimens based on protease inhibitors to those containing a non-nucleoside reverse transcriptase inhibitor. Although an increasing proportion of children receive four drug antiretroviral therapy, nearly all start with triple therapy. Further analyses of changes in prescribing practices for antiretroviral therapy and associated changes in HIV RNA and CD4 cell count responses are planned.

Conclusions

Rates of death, progression to AIDS, and hospital admission in children with HIV in the United Kingdom and Ireland, biases related to referral patterns to specialist centres are not an issue.
Kingdom and Ireland have significantly fallen. As ante-
natal detection rates improve and fewer children born to
infected women are themselves infected, children present-
ing to paediatric services with HIV are likely to be
older and to have been born abroad. This, combined with
improved life expectancy, means that the demand for special-
ist paediatric HIV services will continue to increase.21
Transitional links with adult services are required to deal
with the medical, social, and psychological needs of children entering adoles-
cence and adult life.

National surveillance of paediatric HIV is undertaken by
the National Study of HIV in Pregnancy and Childhood (NSHPC) at
the Institute of Child Health, London, in collaboration with
the Health Protection Agency Communicable Disease Surveil-
 lance Centre and the Scottish Centre for Infection and Environ-
 mental Health. NSHPC relies on active reporting from
paediatricians through the British Paediatric Surveillance Unit
of the Royal College of Paediatrics and Child Health, and from
obstetric respondents reporting through an active reporting
scheme run under the auspices of the Royal College of
Obstetricians and Gynaecologists. We thank all those reporting
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Royal Hospital for Sick Children, Bristol), W Tarnow-Mordi, J
Petrie (Ninewells Hospital and Medical School, Dundee), K
Slopper, V Shah (Ealing Hospital, London), J mok (Royal
Edinburgh Hospital for Sick Children), DMG, VN, N Klein, LF,
M clapson, B Òhene-Keva (Great Ormond St Hospital, London),
C Ball, D Navagum, D graham, A waters (King’s College Hospital, London), DMG, E cooper, T
Fisher, R Barrir, S wong (Newham General Hospital, London),
V Van someren, K moshal, S McKenna (Royal Free Hospital, London), M sharland, S donaghy, W Faulknall (St George’s
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