TUBERCULOSIS

CLINICAL INSIGHTS INTO THE INTERACTION OF CHILDHOOD TUBERCULOSIS AND HIV IN THE WESTERN CAPE

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The mutually deleterious interaction of HIV infection and tuberculosis (TB) was recognised in adults soon after the scope of the HIV epidemic became evident. It is now clear that in populations with a high incidence of infection due to Mycobacterium tuberculosis, clinical TB disease is one of the first signs of declining immunity in HIV-infected adults.

The HIV/AIDS epidemic impacts on childhood TB, both directly and indirectly, but documentation of this interaction is less certain than for adult TB because of the difficulties in diagnosing childhood TB. Firstly, HIV-infected children probably have the same degree of susceptibility to TB infection and subsequent disease as adults, and secondly, HIV-infected and non-infected children will be exposed to increasing sources of TB because of the impact of HIV/AIDS on adult TB prevalence.

The Western Cape (WC) province of South Africa is an area with a remarkably high incidence of TB. The reported TB notification rate was 589 new cases per 100 000 population per year in 1998 (data from Department of Health: Directorate Health Systems Research and Epidemiology). It is also the province with the lowest prevalence of HIV infection at 5.2% (95% confidence interval (CI) 3.2 - 7.2%) according to the 1998 national HIV survey of women attending public antenatal clinics (data from Department of Health: Directorate Health Systems Research and Epidemiology).

TB is often an early opportunistic infection in adults in developing countries as they are already infected with the tubercle bacillus at the time that their immunity starts to decline. In children, the situation may be different because they are almost always primarily infected by mother-to-child transmission of HIV, while the TB infection usually occurs later in life. Therefore, children will more often present with other infections before they develop TB disease. This is confirmed by postmortem studies of young children dying of HIV/AIDS; TB was only found in 1/78 cases in the Ivory Coast and in 1/74 children with a median age of 2.7 years in South America.

Fig. 1. Age at time of culture-confirmed TB diagnosis in HIV-infected children (N = 29). https://www.sajhivafrica.org/article/view/46174/46206
**WESTERN CAPE EXPERIENCE**

In our own experience, working in communities with a very high incidence of TB, young HIV-infected children (less than 2 years of age) can develop culture-positive TB and have culture-confirmed TB. However, documenting childhood HIV infection has been difficult. This problem has been accentuated by HIV infection and its associated lung conditions, which on clinical and radiological evaluation can present in similar ways. Over-reporting and over-diagnosis of TB is therefore most likely, and to evaluate the role of TB in HIV-infected children, culture of the *M. tuberculosis* organism remains important. Criteria suggested by the World Health Organisation (WHO) for diagnosing TB in children, such as weight loss and cough of more than 2 weeks' duration, have become less helpful. Evaluating 29 children with culture-positive TB at TB hospital over a period of 8 years showed that some criteria may still be helpful (Table 1) and from observations done at TB hospital it is clear that determining whether the contact has drug-resistant TB is essential. Five of 13 (38%) HIV-infected children with culture-confirmed TB admitted over a 1-year period had drug-resistant TB, 4 of whom had received previous TB treatment. Three of the latter 4 children had previously had drug-sensitive TB, but thereafter had contact with adults with drug-resistant TB. In some cases this involved the same adult who also developed drug-resistant TB after previously being fully sensitive.

**DIAGNOSIS OF TB**

The diagnosis of TB in children has always been difficult. Contact history remains important, and from observations done at TB hospital it is clear that determining whether the contact has drug-resistant TB is essential. Five of 13 (38%) HIV-infected children with culture-confirmed TB admitted over a 1-year period had drug-resistant TB, 4 of whom had received previous TB treatment. Three of the latter 4 children had previously had drug-sensitive TB, but thereafter had contact with adults with drug-resistant TB. In some cases this involved the same adult who also developed drug-resistant TB after previously being fully sensitive.

Tuberculin skin tests have often been reported to be of little value in dually infected children mainly from developed countries. However, in our series, 50% of 24 culture-proven TB cases had a Mantoux test of 15 mm (H S Schaaf, M F Cotton - unpublished data). That no child had a Mantoux size between 5 mm and 14 mm leads one to consider an 'all or nothing' type of response to the tuberculin test. More important, this shows that the Mantoux test remains a useful tool in the diagnosis of TB in children. It is even truer than in the past that a negative tuberculin skin test does not exclude the diagnosis of TB. These children probably had significant tubercul reactions because they presented quite early in the course of their HIV disease and probably still had relatively good immunity. Clinical signs are not of much help in terms of diagnosing TB in children.

Extrapulmonary TB such as lymph node TB, tuberculous meningitis and abdominal TB are reported to be present in about 25% of children with TB. Extrapulmonary TB has been reported to be more common in adults with HIV infection. Forty-one per cent of the children in our series had confirmed extrapulmonary TB.

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**HIV-infected children**

<table>
<thead>
<tr>
<th>Feature</th>
<th>N</th>
<th>%</th>
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<tbody>
<tr>
<td>Adult TB contact</td>
<td>19</td>
<td>66</td>
</tr>
<tr>
<td>Cough &gt; 2 weeks</td>
<td>14</td>
<td>48</td>
</tr>
<tr>
<td>Confirmed weight loss</td>
<td>22</td>
<td>76</td>
</tr>
<tr>
<td>Hepatomegaly</td>
<td>26</td>
<td>90</td>
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<tr>
<td>Respiratory signs</td>
<td>20</td>
<td>69</td>
</tr>
<tr>
<td>Splenomegaly</td>
<td>16</td>
<td>55</td>
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<tr>
<td>Otorrhea acute/chronic</td>
<td>10</td>
<td>34</td>
</tr>
<tr>
<td>Extrapulmonary TB</td>
<td>12</td>
<td>41</td>
</tr>
<tr>
<td>Anaemia (Hb &lt; 10 g/dL)</td>
<td>26</td>
<td>90</td>
</tr>
<tr>
<td>Mantoux ≥ 15 mm (N = 24)</td>
<td>12</td>
<td>50</td>
</tr>
</tbody>
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*Table 1. Symptoms, signs and side-room investigations at culture-confirmed diagnosis of TB in 29 HIV-infected children.*

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**Chest radiograph demonstrating paratracheal and hilar lymph nodes with the compression of the left main bronchus.**
Interpreting chest radiographs in HIV-infected children has become increasingly difficult. The reticulonodular pattern typical of miliary TB can be identical to lymphoid interstitial pneumonia (LIP), and both conditions could have enlarged hilar and/or mediastinal lymphadenopathy. Hepatosplenomegaly occurs in both and is also of no value with regard to diagnosis. Furthermore, patients with LIP may also develop TB. Recurrent pneumonias, more nonspecific presentations of LIP, bronchiectasis, Pneumocystis carinii pneumonia, and other lung conditions may add to the diagnostic dilemma. We have observed a very high prevalence of significant paratracheal adenopathy (52%), hilar adenopathy (67%), and bronchial compression (30%), the latter even in older children (2-3 years of age), in 27 culture-confirmed cases (H S Schaaf, M F Cotton - unpublished data). (See radiographs) This incidence is significantly higher than in HIV-negative patients.12

Fig. 2. Final confirmation of HIV diagnosis in children with confirmed TB (N = 29) (ELISA = enzyme-linked immunosorbent assay, PCR = polymerase chain reaction; p24ag = p24 antigen test)

RELAPSE OF TB

Relapse (re-activation or re-infection) occurred in 10 of our patients (34%). Two children had been on treatment for between 4 and 5 months when M. tuberculosis was again cultured. In the remainder, TB was reconfirmed by culture after a median of 23 months (range 10-83 months) of initial diagnosis. We recently confirmed by DNA fingerprinting that one of these patients had true reinfection. Drug susceptibility testing was available for 6 patients, with only 1 child drug-resistant. His mother initially had drug-susceptible TB and developed multidrug-resistant TB over a period of 28 months.

REFERENCES