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Development of a scoring system for the diagnosis of tuberculous lymphadenitis


Introduction

The HIV/AIDS epidemic has been associated with an increase in smear-negative pulmonary and extrapulmonary registered tuberculosis (TB) cases in many countries including Malawi [1]. Tuberculous lymphadenitis accounts for approximately 25% of the extrapulmonary cases [2]. Up to 50% of HIV infected individuals will develop persistent generalised lymphadenopathy (GPL) in the course of their disease [3]. It can be a challenge to distinguish between PGL and lymphadenopathy due to TB especially in resource poor settings without histopathological and cytological diagnostic facilities. The aim of this prospective study was to develop and evaluate a scoring system based on history and physical examination for the diagnosis of glandular TB that would be applicable in settings without laboratory backup in Malawi.

Methods

Study site

This study was conducted in Karonga District- northern Malawi in the context of the Karonga Prevention Study (KPS), a large epidemiological study of mycobacterial diseases and HIV. Since 1986 the KPS has been responsible for “case-finding” and “case-holding” for both TB and leprosy as part of follow-up for a large vaccine trial [4]; HIV prevalence among antenatal women and smear-positive pulmonary TB cases were approximately 10% and 60%, respectively, in the late 1990s [5].

Recruitment

From October 1997 to October 1999 individuals found with significant lymphadenopathy (>2cm) in at least one extracranial site were invited to join the study. These were identified during routine standard full body examinations (for leprosy) amongst outpatients, inpatients (both at the District Hospital and Health Centres across the district) and in households during field based studies. Each consenting individual had a “glandular TB suspect” form (figure 1) completed. The various signs carried numerical weights that were assigned on the basis of literature review and a pilot investigation. Once the form was filled in, the paramedic performing the examination calculated a total score and this determined subsequent action. If the total score was less than 10, the individual was asked to return for review after 3 months. If it was between 10 and 19 a week’s course of an antibiotic (co-trimoxazole or doxycycline) treatment trial was given, with instructions to return after completing the dosage. Individuals who scored 20 or greater were immediately referred to Karonga District Hospital (KDH) for further investigations.

Assessment of patients

At KDH, individual verbal consent for biopsy was obtained. For those less than 16 years old a parent or guardian consented. Pre-test counselling for HIV was given to all patients before a blood sample was drawn. Post-test counselling was given to those who wanted to know their status before disclosing the result. Each individual had a fine needle aspiration (FNA) performed using a 23 gauge needle and 20ml syringe mounted onto a suction gun. The material obtained was applied onto a slide, then thinly spread using a second slide, making two films in the process. Both slides were dried, air dried and then stored in a slide box. The material remaining in the syringe was applied directly onto solid Lowenstein Jensen (L-J) media. An excision biopsy was then performed under sterile conditions and local anaesthesia. The node removed was divided into two portions, and the cut surface examined macroscopically for caseation. One section was stored in a bottle containing formalin for histology. The other half was put in a plain sterile container and kept frozen. If caseation was seen at FNA and/or excision biopsy the patient was started on antituberculosis treatment according to the Malawi National TB programme treatment guidelines [6]. A blood sample for HIV testing was collected at the same time. All specimens were transported to the KPS laboratory, at the project’s headquarters in Chilumba, for further processing.

Laboratory procedures

At the laboratory one of the FNA slides was stained with auramine-phenol and examined using fluorescence microscopy. Scantly positive slides were destained and restained by Zehl-Nielsen for confirmation. The frozen section was thawed, a smear made and the rest was crushed, decontaminated and 2 cultures set up. The cultures included pyruvate-rich media to encourage growth of Mycobacterium bovis [7]. These cultures (including the one made from FNA material) were examined fortnightly for 8 weeks. Those with growth and colony morphology suggestive of M tuberculosis were sent to the

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Figure 1

GLANDULAR TB SUSPECT FORM

Name: ________________________  Sex: M/F ___________  Date of presentation: __/__/____  Reason for presentation: ________________________  with GP form

HISTORY

Fever: None (0) 1-14 days (1) >14 days (2)

Night sweats: None (0) 1-14 days (1) >14 days (2)

Weight loss: None (0) Moderate (2) Severe (1)

Pain in node: None (0) Moderate (2) Severe (1)

Growth (2 weeks): None (0) Moderate (1) Much (3)

TB contact: None known (0) H/H Fish <2yrs (2) H/H Fish <2 yrs(3)

CLINICAL EXAM

General conditions: Fair (1) Sick (2) Very ill (3)

Enlarged glands:

Cervical (C1) L Y/N R Y/N

Submandibular (S) L Y/N R Y/N

Subcervical (M) L Y/N R Y/N

Axillary (A) L Y/N R Y/N

Epitrochlear (E) L Y/N R Y/N

Mastoidal (T) L Y/N R Y/N

Symmetry: Site: No (3) Yes (then consider size)

Size: Equal (1) Unequal (2)

Maximum score for any node group for each characteristic:

Consistency: Soft (0) Firm (1) Hard (2) Fluctuant (3)

Mobility: Mobile (1) Doubtful (2) Fixed (3)

Fused/Matted: No (1) Doubtful (2) Yes (3)

Note: Group = one or more nodes at a particular site

TOVAI SCORE ACTION: IF <10 Review Treatment trial >20 Refer to hospital

TREATMENT TRIAL

Antibiotic: ________________________  Dates: __/__/____ to __/__/____

Result: If no change, or doubtful, MO to review; else discharge.

Date: __/__/____  Staff code: ________

Checker: ________

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PHLS Mycobacterium Reference Laboratory in Dulwich, London, UK for species identification and drug sensitivity testing. The section fixed in formalin and the remaining slide from FNA were sent to St Thomas’ Hospital, London for histopathological and cytological examination respectively (by SBL).

HIV testing was carried out following a standard KPS protocol involving dual testing by both particle agglutination (Edgware- Mast Diagnostics-modified Serodia at two dilutions) and by sandwich ELISA (Organon- Vironostika Uniform II plus 0 test-Organon Teknik). Only concordant results are reported to the patient. In case of discordant results both tests are repeated in duplicate on serum recrystallized from the original specimen.

Data management
All data were double entered and verified at the Project Headquarters in Karonga using Epi info (version 6; Center for Disease Control (CDC; U.S.A) and Foxpro (version 2.6A) software. Analyses were performed in Stata (version 6, Stata Corporation, Texas, USA).

Diagnostic criteria
For this analysis, a case was considered “confirmed” if histological findings were consistent (pathognomonic) with TB beyond reasonable doubt and/or *M. tuberculosis* was identified from culture.

Results
Overall 166 patients were enrolled into the trial as shown in figure 2. Of these 52 did not have further investigations (28 improved on treatment trial, 3 were too ill for biopsy and 21 did not return for review). Of the remaining 114 patients, 105 had excision biopsy and FNA performed (75 TB confirmed), 6 had excision biopsy only (2 TB confirmed) and 3 had FNA only (2 TB confirmed bacteriologically). The mean age of the confirmed TB cases was 28.4 years (SD 10.8, range 0-52 years). The age and sex distribution of individuals with a confirmed TB lymphadenitis diagnosis (histologically and/or bacteriologically) is shown in figure 3.

Microscopy results of FNA and frozen section slides are available on 72 confirmed TB cases. Of these 11 (15.3%) were positive on a smear made from FNA material (of these 5 were negative on frozen section smears) and 15 (20.8%) were positive from frozen section smears (of these 7 were negative on the FNA smear). Of the 75 cultures set up from confirmed TB cases, 32 (42%) yielded *M. tuberculosis* from the FNA culture (11 of these had no growth from the frozen section) whereas 36 (50%) had MTB identified from the frozen section cultures (9 of these had no growth on the FNA culture). Of note 15 (20%) of the FNA cultures were contaminated compared to 3 (4%) from the frozen section.

Macroscopic examination records were available for 74 confirmed TB cases. Out of these, 64 (85.6%) were correctly identified as having TB based on caseation. Of ten that were not recognised on this basis, one was recorded as having no caseation and 9 were classified as “doubtful”. Of the 34 non-TB cases, 2 (5.9%) were identified as having caseation on naked eye (macroscopic) examination (1 Kaposi Sarcoma and 1 Non-Hodgkin’s Lymphoma). Of the 63 TB confirmed cases with an HIV result, 54 (85.7%) were seropositive. All eight cases of Kaposi sarcoma were HIV seropositive.

The distribution of total points scored by individuals with a confirmed diagnosis is shown in figure 4. The distribution of clinical features amongst individuals with a confirmed diagnosis is shown in the table. Significant *p* values are highlighted.

Discussion
This is a preliminary analysis. A detailed multivariate analysis of diagnostic scores and results is underway.
Overall there was no difference in the proportion of confirmed TB cases by sex ($p=0.16$) or by age ($p=0.13$). However we note that there is marked female excess in young adults (15-29 years) with glandular TB. This distribution mirrors that of HIV in Malawi[8]. The high HIV seroprevalence amongst glandular TB patients has implications for care. The relatively low proportion of smear positive from FNA and biopsy sections are similar to those reported previously [9] [10]. One possible explanation is that we included individuals from outpatient clinics and from the community, hence in early stages of their disease with low bacillary load. It is worth adding that nodes which yielded much material at FNA were more likely to be smear positive ($p=0.02$) and culture positive ($p=0.04$) than were the nodes that yielded little material. The high contamination rate (20%) of the FNA cultures is not unexpected as these cultures were set up in the operating room without any decontamination process. Notably there was no significant difference in the total number of positive cultures between FNA and frozen section cultures ($p=0.8$). This study confirms the rarity of human M bovis disease in Karonga, as all the positive cultures were identified as M. tuberculosis. This is despite setting up cultures on pyruvate-rich media to encourage growth of M. bovis.

Sensitivity and specificity of macroscopic examination of the sectioned nodes were 97.0% and 76.3% respectively. The low specificity achieved here is of concern. This is due to the 9 cases that were classified as “doubtful” for cation. Of these, 7 had an HIV result and 6 were seropositive. These atypical macroscopic and histological features of tuberculous lymphadenitis in HIV infected individuals have been described elsewhere[11]. The potential for error in macroscopic examination for cation in malignancy (e.g. Kaposi Sarcoma, Hodgkin’s) is known [12]. This calls for particular attention in examining for typical Kaposi sarcoma mucocutaneous lesions in individuals suspected of tuberculosis lymphadenitis.

With regard to the performance of the scoring system, confirmed tuberculous lymphadenitis patients tended to achieve higher total scores than those patients with a non-TB diagnosis (12 test for heterogeneity $p<0.01$). Among the clinical features fever, night sweats, weight loss, growth, anatomical distribution and consistency achieved statistical significant level ($p<0.05$). Logistic regression analysis will determine the best combination of these features for optimum performance. A possible limitation of this study could be that since experienced pathomacros examined carried out enrolment, they recruited clients in whom a TB diagnosis was most likely. Thus this study could be measuring their clinical acumen. It is imperative therefore that this tool be evaluated in other settings of equally high HIV prevalence.

Acknowledgement

We thank the staff at KPH and Karonga District Hospital for their dedication to the field and laboratory work for this study. We also thank all the patients who accepted to take part in this trial. The KPH is funded largely by the Wellcome Trust. BN received support from the British High Commission.

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Reference

8. UNAIDS. Epidemiological fact sheets on HIV/AIDS and sexually transmitted infections; Malawi. WWW. UNAIDS. 2000.

Table: Distribution of confirmed TB lymphadenitis cases (n=79) and of total number of suspects (n=104) by level (weight point) of each clinical feature

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<th>Weighted points</th>
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<th>Weight loss</th>
<th>Pain</th>
<th>Growth</th>
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* For description of each weighted point for each clinical characterisitc - refer to figure 1

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