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Child and adult spinal tuberculosis at tertiary hospitals in the Western Cape, South Africa: 4-year burden and trend

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Abstract

The aim of this retrospective review was to assess the overall burden and trend in spinal tuberculosis (TB) at tertiary hospitals in the Western Cape Province of South Africa. All spinal TB cases seen at the province's three tertiary hospitals between 2012 and 2015 were identified and clinical records of each case assessed. Cases were subsequently classified as bacteriologically confirmed or clinically diagnosed and reported with accompanying clinical and demographic information. Odds ratios (OR) for severe spinal disease and corrective surgery in child vs. adult cases were calculated. A total of 393 cases were identified (319 adults, 74 children), of which 283 (72%) were bacteriologically confirmed. Adult cases decreased year-on-year (P=0.04), however there was no clear trend in child cases. Kyphosis was present in 60/74 (81%) children and 243/315 (77%) adults with available imaging. Corrective spinal surgery was performed in 35/74 (47%) children and 80/319 (25%) adults (OR 2.7, 95% confidence interval 1.6–4.5, P=0.0003). These findings suggest that Western Cape tertiary hospitals have experienced a substantial burden of spinal TB cases in recent years with a high proportion of severe presentation, particularly among children. Spinal TB remains a public health concern with increased vigilance required for earlier diagnosis, especially of child cases.

Introduction

Tuberculosis (TB) continues to present a considerable public health concern, particularly in developing countries where high rates of human immunodeficiency virus (HIV) infection and adverse socio-economic factors contribute to high rates of TB disease and TB-related mortality [1, 2]. In 2016, South Africa had the highest estimated incidence of TB in the world at 781 cases per 100 000 population [2]. This overall estimate includes both notified and estimated undiagnosed cases, whereas only 244 053 of the estimated 438 000 cases (56%) were notified [2]. Within South Africa, the Western Cape Province is known to carry a particularly high burden of TB, with an incidence of 681 notified cases per 100 000 population and HIV co-infection rate of 38.5% in 2015, according to the electronic TB register [3]. In this instance, the estimated number of undiagnosed/unreported cases has not been accounted for and the true incidence of TB in the province is almost certainly higher. Within such a high-burden setting, it is not surprising that sites of TB disease that constitute only a small percentage of all cases amount to a relatively large absolute number of individuals affected. For example, although spinal TB accounts for only 1–5% of all TB cases [4–6], previous reports suggest that spinal TB is a common pathology at tertiary orthopaedic units within the province [7–9].

TB of the spine typically occurs following haematogenous spread of *Mycobacterium tuberculosis* from a primary infection site and is characterised by gradual destruction of one or more spinal vertebrae [10–13]. Although progression of the disease is relatively slow and insidious, diagnosis is often delayed and advanced cases may present with kyphotic deformity, spinal instability and neurological deficits due to compression of the spinal cord [10–13]. These sequelae may be particularly prevalent in children as the disease progresses more rapidly in the high cartilage content of the immature spine and there is a greater likelihood of buckling and collapse [10, 14]. As a result, many children with spinal TB sustain significant kyphotic deformity during the active phase of the disease and all children remain at risk of secondary complications until skeletal maturity [14, 15].

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Within the Western Cape, the recommended diagnostic procedure for patients with suspected spinal TB involves a magnetic resonance imaging (MRI) scan and a biopsy of the spinal lesion or paraspinal abscess [8, 16]. While TB disease is treated with antituberculosis medication, many patients may also require corrective orthopaedic surgery [10, 17]. Anecdotal evidence suggests that in-patient diagnosis and surgical intervention for patients with spinal TB constitutes a significant burden on specialist orthopaedic services in this TB endemic setting. However, the true burden of spinal TB in the province has not been investigated. Furthermore, it is unclear whether the recent decline in the incidence of all reported TB cases within the province, from 777 per 100 000 population in 2012 to 681 per 100 000 population in 2015 [3], has had any impact on the number of new spinal TB cases over recent years.

Increased understanding of the burden of spinal TB in this setting may provide key insights for health system adjustments and the planning of future spinal TB research, a field in which Africa is currently poorly represented despite its high TB burden [18]. Therefore, the aim of the current study was to assess the burden of spinal TB at tertiary hospitals in the Western Cape, including the overall number of adult and child cases, the clinical and demographic characteristics of patients affected and the change in case numbers by year.

Methods

Study design and setting

This retrospective, descriptive study identified cases of spinal TB treated at tertiary hospitals in the Western Cape Province of South Africa between 1 January 2012 and 31 December 2015. The study was conducted in collaboration with orthopaedic services at each of the province's three tertiary hospitals (Tygerberg Hospital, Groote Schuur Hospital and Red Cross War Memorial Children's Hospital) and was approved by the Health Research Ethics Committee of Stellenbosch University and by the management of the hospitals involved. The authors assert that all procedures contributing to this work comply with the ethical standards of the relevant national and institutional committees on human experimentation and with the Helsinki Declaration of 1975, as revised in 2008.

Case identification and data collection

Cases of spinal TB in children and adults were identified using clinicians' databases, theatre log books and a search of the National Health Laboratory Service (NHLS) TB investigations. Cases identified were cross-referenced and all duplicates removed.

Clinical and demographic data including age, gender, HIV status and the results of samples submitted for bacteriological and histological testing were captured from NHLS laboratory results systems and available medical records. Additional information was captured for HIV-positive patients, including CD4 count, viral load (VL), timing of HIV diagnosis and antiretroviral therapy (ART) status. CD4 count and/or VL are typically measured only once or twice a year according to the standard ART monitoring in South Africa and results closest to hospital admission, from 12 months prior to admission to 1 month after admission, were included as available. Diagnosis with HIV at approximately the same time as presenting to the health system with spinal TB was considered a new HIV diagnosis as opposed to a prior HIV

diagnosis. HIV-positive patients initiated on ART only during or after presentation with spinal TB were categorised as not on ART within the descriptive analysis.

Radiological outcomes for all patients including the number of vertebrae affected and diagnosis were captured from MRI reports. Furthermore, the presence or absence of kyphotic deformity was assessed by a spine surgeon based on available imaging. If applicable, a description of the spine surgery performed was obtained from the afore-mentioned clinicians' databases, theatre log books or medical records. Corrective surgery included procedures such as instrumented fusion whereas needle biopsies, abscess drainage and costotransversectomy were considered minor surgical procedures. All data collected for the study was de-identified and entered into custom-designed forms using REDCap electronic data capture tools hosted at Stellenbosch University.

Case classification

Cases were classified as 'bacteriologically confirmed' or 'clinically diagnosed', according to the 2013 revised World Health Organization (WHO) case definitions [19]. In the current study, confirmed cases were those patients for whom the presence of *M. tuberculosis* in a spine biopsy was demonstrated by a positive Xpert MTB/RIF test (Cepheid, Sunnyvale, California, USA; available for routine testing from approximately 2014 onwards), positive culture for *M. tuberculosis* or positive smear microscopy for acid-fast bacilli. Clinically diagnosed cases were those patients for whom spine imaging, clinical presentation and, in some instances, confirmed TB from other sites indicated spinal TB, but there was no bacteriological confirmation from a spine biopsy. Patients 0–14 years of age and patients 15 years or older at the time of diagnosis were classified as child cases and adult cases, respectively.

Data analysis

Upon completion of the study, data were exported from REDCap to Microsoft Excel and then transferred to Graphpad Prism (GraphPad Prism version 6.00 for Windows, GraphPad Software, La Jolla, California, USA, http://www.graphpad.com) for analysis. The overall burden of spinal TB cases was presented as counts. Case counts per year and annual, age-specific Western Cape population estimates [20–23] were used to estimate the incidence of spinal TB in the Western Cape during the study period. No report of population estimates was available for 2012 and this estimate was taken as the average of 2011 [20] and 2013 [21] values. Categorical descriptive data were presented as counts and percentages, as appropriate. Odds ratios (ORs) were used to investigate the association between female gender and HIV infection among adults with known HIV status and the association between CD4 count <200 cells/µl and ≥3 vertebrae affected and CD4 count <200 cells/µl and receiving corrective surgery among HIV-positive adults. The severity of spinal TB presentation in children vs. adults was compared using the OR for ≥3 vertebrae affected, presence of kyphotic deformity and having received corrective spine surgery. All ORs were calculated with 95% confidence intervals (CIs) and significance was accepted at P < 0.05. Finally, a two-sided Jonckheere-Terpstra test was used to assess the trends in counts and TB incidence over the study period. This trend analysis was conducted using STATA version 15 (Stata Corporation, College Station, Texas, USA) and significance was accepted at P < 0.05.

Results

A total of 393 cases of spinal TB were diagnosed at tertiary hospitals in the Western Cape within the 4-year study period, including 319 adults and 74 children. The majority of cases presented from the Cape Town Metro (n = 313, 80%) with much smaller representation from the surrounding rural districts (Cape Winelands, n = 39 (10%), West Coast, n = 13 (3%), Overberg, n = 10 (3%), Eden, n = 9 (2%)) and outside of the province (n = 8, 2%). Demographic data, HIV status and the results of diagnostic tests are presented in Table 1.

Age and gender

The median age of adults was 38 years with a range of 16–82 years, while the median age of children was 7 years with a range from 1 year 7 months to 14 years 9 months. Approximately 57% of patients were female in both adult and child groups. The percentage of spinal TB cases within each 5-year age bracket, in total and by gender, is shown in Figure 1.

HIV status

Based on available HIV test results, 41% of adults with spinal TB were HIV-positive νs . only 3% of children. However, HIV status was unknown or could not be determined from the available data in 12% of adults and 28% of children. When examining HIV status by 5-year age bracket, the highest proportion of HIV-positive cases were observed among 25–54 years age groups whereas those ≤ 14 or ≥ 55 years of age were predominantly HIV-negative or of unknown HIV status (Fig. 2). When comparing HIV status between genders, 89 of 160 (56%) adult females and 43 of 121 (36%) adult males with known HIV status were HIV-positive. Based on these proportions, females were significantly more likely to be HIV-positive than males (OR 2.3, 95% CI 1.4–3.7, P = 0.001).

CD4 counts were available in 117/132 (89%) HIV-positive adults; the median CD4 count was 325 cells/µl (IQR 193–520 cells/µl) with a median CD4% of 21.38% (IQR 14.95–28.12%). Of 74 (56%) adults with available VL, 54 had lower than detectable VL (<20 copies/ml) and 20 had a detectable VL with a median of 3.25 log₁₀ copies/mL (IQR 2.24–5.02 log₁₀ copies/ml). The median absolute time difference between CD4 test results and admission and VL test results and admission was 2 weeks (IQR 0–10 weeks) and 10 weeks (IQR 4–18 weeks), respectively. Most adults had prior HIV diagnosis and were on ART upon presentation with spinal TB. Further HIV-specific details are shown in Table 2.

Only two children were confirmed HIV-positive; an 8-year-old boy with a prior HIV diagnosis on ART (CD4 count: 105 cells/ μ l, CD4%: 6.86% and VL: 3.13 log₁₀ copies/ml) and an 11-year-old girl newly diagnosed with HIV not yet on ART (CD4 count: 283, CD4%: 13.73 and VL: 4.04 log₁₀ copies/ml).

Diagnosis

Imaging was available for 318/319 adults (n = 314 MRI scans, n = 3 computed tomography (CT) scans and n = 1 radiograph only) and 72/74 children (n = 64 MRI scans, n = 1 CT scan, n = 7 radiograph only) with MRI or CT findings specifically suggestive of TB in 309 (97%) adults and in 62 (95%) children. Of patients who received an MRI or CT scan, 53/382 (14%) had non-contiguous

involvement of two or more vertebrae. Of 211 patients with spine biopsies submitted for Xpert MTB/RIF testing, 145 (69%) were positive. Similarly, of 354 patients with spine biopsies submitted for mycobacterial culture, 229 (65%) were positive. Further details of how MRI and spine biopsy results related to one another are included as online Supplementary Material (Fig. S1).

Among 263 (70%) patients with known drug susceptibility test (DST) results, 239 (91%) had susceptibility to isoniazid and rifampicin or rifampicin based on DST on mycobacterial culture or Xpert MTB/RIF, respectively. However, 24 patients (9%) had drug-resistant TB: eight patients (3%) had mono-resistant TB (n=4 rifampicin, n=4 isoniazid), 14 patients (5%) had multidrug-resistant (MDR)-TB and two patients (<1%) had extensively drug-resistant-TB. Although two cases of MDR were based on Xpert only, all other cases of drug resistance were identified from culture DST.

When considering all diagnostic evidence together, 283 (72%) of all spinal TB cases were bacteriologically confirmed from a spine biopsy. The proportion of confirmed cases was slightly higher in adults than in children (73% vs. 66%). However, 315 of 319 (99%) adults and only 57 of 74 (77%) children had spine biopsy specimens sent for bacteriology, affording greater opportunity for bacteriological confirmation among the adults (data not shown).

Severity of presentation

A large number of adults and children presented with severe spinal disease, based on the number of vertebrae affected, the presence of kyphotic deformity and corrective surgery received (Table 3). While children were no more likely than adults to have $\geqslant 3$ vertebrae affected or to present with kyphotic deformity, children were 2.7 (95% CI 1.6–4.5, P=0.0003) times more likely to have received corrective surgery, compared with adults (Table 3). Overall, 115 (29%) spinal TB patients received corrective surgery. Among HIV-positive adults with an available CD4 count, there was no association between CD4 count <200 cells/µl and having $\geqslant 3$ vertebrae affected or receiving corrective surgery.

Cases by year

Adult cases of spinal TB decreased year-on-year during the study period with 105 cases in 2012 and 56 cases in 2015 (Fig. 3). This decrease was accompanied by a decrease in the incidence of spinal TB among adults in the Western Cape from 2.57 cases per 1 00 000 in 2012 to 1.22 cases per 1 00 000 in 2015. Both case counts and incidence of spinal TB among adults showed a significant downward trend over the 4-year study period (z=-2.04, P=0.04 for both counts and incidence). Conversely, the number of children diagnosed with spinal TB each year showed no significant trend with 16 cases in both 2012 and 2015 (Fig. 3). The incidence of spinal TB among children also remained similar at 1.03 cases per 1 00 000 and 0.99 cases per 1 00 000 in 2012 and 2015, respectively, with no significant trend.

Discussion

The main finding of the study was that 393 new cases of spinal TB were treated at tertiary hospitals in the Western Cape between 2012 and 2015 with 72% of all cases bacteriologically confirmed

Table 1. Demographics, investigation of spine lesions and case classification of children and adults with spinal tuberculosis at tertiary hospitals in the Western Cape, 2012–2015

	Children (0–14 years) n=74 (%)	Adults (>14 years) n = 319 (%)	All n = 393 (%)
Demographics			
Age (median [IQR])	7 [4-11]	38 [30–50]	34 [23–47]
Gender (males)	31 (42)	137 (43)	168 (43)
HIV status			
Positive	2 (3)	132 (41)	134 (34)
Negative	51 (69)	149 (47)	200 (51)
Unknown	21 (28)	38 (12)	59 (15)
Spine imaging			
MRI/CT findings ^a	n = 65 (88)	n = 317 (99)	382 (97)
Suggestive of TB	62 (95)	309 (97)	371 (97)
Not suggestive of TB ^b	3 (5)	8 (3)	11 (3)
Spine biopsy pathology			
Histology	n = 41 (55)	n = 272 (85)	n = 313 (80)
Necrotizing granulomatous inflammation	31 (76)	126 (46)	157 (50)
Inconclusive	10 (24)	146 (54)	156 (50)
Ziehl Neelsen stain	n = 40 (54)	n = 251 (79)	n = 291 (74)
Positive	13 (33)	36 (14)	49 (17)
Negative	27 (67)	215 (86)	242 (83)
Spine biopsy microbiology			
Auramine stain	n = 51 (69)	n = 243 (76)	294 (75)
Positive	11 (22)	55 (23)	66 (22)
Negative	40 (78)	188 (77)	228 (78)
Xpert MTB/RIF	n = 31 (42)	n = 180 (56)	n = 211 (54)
Positive	24 (77)	121 (67)	145 (69)
Negative	7 (23)	59 (33)	66 (31)
M. tuberculosis culture	n = 56 (76)	n = 298 (93)	n = 354 (90)
Positive	35 (63)	194 (65)	229 (65)
Negative	21 (37)	104 (35)	125 (35)
Case classification			
Bacteriologically confirmed	49 (66)	234 (73)	283 (72)
Clinically diagnosed ^c	25 (34)	85 (27)	110 (28)

Data are presented as n (%) for non-missing data, unless otherwise indicated, with % values rounded to the nearest whole number.

from a spine biopsy. The majority of patients presented with kyphotic deformity and 115 (29%) of all patients received corrective spinal surgery. However, indications for surgical intervention were particularly prevalent among children with approximately one in two children vs. one in four adults receiving corrective surgery (Table 3). These findings highlight the considerable burden of spinal TB in the current setting, including the impact on affected individuals and on the public health system.

Spinal TB among adults in the Western Cape

In the current study, 50% of all spinal TB cases were between 23 and 47 years old with a median age of 38 years among adult cases (Table 1). While previous studies from the USA [4] and Eastern Mediterranean [11] reported a somewhat older median age of approximately 50 years among spinal TB cases, the current age distribution is similar to findings from Kenya [24] and China

^aMRI, magnetic resonance imaging (n = 378); CT, computed tomography (n = 4).

bAlthough MRI findings did not suggest TB, spinal TB was diagnosed following bacteriological (n = 10) or histological (n = 1) spine biopsy findings.

^cAmong the clinically diagnosed cases (no bacteriological confirmation from a spine biopsy), seven adults and seven children had bacteriological confirmation from another site (lungs, *n* = 8; lymph node, *n* = 2; pleura, *n* = 1; knee, *n* = 1; groin, *n* = 1; liver, *n* = 1).

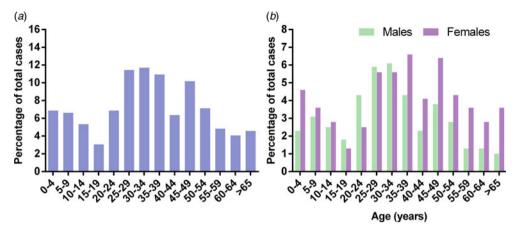


Fig. 1. Cases of spinal tuberculosis presented as (A) total cases and (B) male and female cases, per 5-year age bracket.

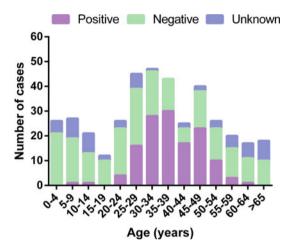


Fig. 2. HIV status of spinal tuberculosis cases by 5-year age bracket.

[12] and suggests that spinal TB primarily affects adults from a relatively young, economically active population in regions with a high TB burden.

Fifty-seven per cent of adults in the current study were female, a finding in keeping with previous spinal TB studies from South Africa [7, 8, 25, 26] but in contrast to other large studies in which 51–61% of spinal TB cases were male [4, 11, 12, 24]. One possible explanation for this difference is HIV infection and its oft-cited association with extra-pulmonary TB disease [27, 28]. For example, whereas HIV co-infection was $\leqslant\!4\%$ [4, 24] or not reported [11, 12] in the afore-mentioned studies from other parts of the world, 56% of adult females and 36% of adult males with known HIV status in the current study were HIV-positive, indicating a significantly higher HIV-positive prevalence among adult females in this study.

Overall, 41% of adults in the current study were HIV-positive, an HIV prevalence notably higher than the 20% HIV prevalence reported among patients with spinal TB treated at Groote Schuur Hospital in 2013–2014 [9]. This difference is most likely explained by the inclusion of children \leq 14 years (none of whom were confirmed HIV-positive) in the HIV prevalence from this previous study along with the inclusion of high HIV prevalence areas in the Tygerberg Hospital catchment area for the current study [29]. Unknown HIV status among 15% of all cases in the current

Table 2. Clinical characteristics at spinal tuberculosis diagnosis among HIV-infected cases

HIV-INTECTED Cases					
	Children N = 2 (%)	Adults N = 132 (%)			
CD4 count					
<200 cells/μl	1 (50)	30 (23)			
200–499 cells/μl	1 (50)	56 (42)			
≽500 ells/μl	-	31 (23)			
Unknown	-	15 (11)			
Viral load					
Lower than detectable ^a	-	54 (41)			
Detectable viral load	2 (100)	20 (15)			
Unknown	-	58 (44)			
HIV diagnosis					
New diagnosis	1 (50)	17 (13)			
Prior diagnosis	1 (50)	104 (79)			
Unknown	-	11 (8)			
Antiretroviral therapy					
On treatment	1 (50)	95 (72)			
Not on treatment	1 (50)	15 (11)			
Unknown	-	22 (17)			

a<20 viral RNA copies/ml

study is in keeping with the 16% unknown HIV status in the afore-mentioned study at Groote Schuur [9]. The apparent absence of HIV testing among these cases is a notable concern. However, it is acknowledged that some patients may have received HIV testing without this being reflected in the records available for the current study, e.g. a rapid test conducted at a primary care clinic.

The overall HIV prevalence in South Africa and in the Western Cape has been estimated at 12% and 5% [30], respectively, suggesting that HIV prevalence may be considerably higher among individuals with spinal TB than in the general population. Most adults with confirmed HIV were diagnosed prior to presentation with spinal TB, were on ART and had lower than

spine deformity and spine surgery:	among children and adults with spinal tuberculosis
	e, spine deformity and spine surgery a

	Children N=74 (%)	Adults N = 319 (%)	Children vs. adults OR (95% CI)	<i>P</i> -value
Vertebrae affected ^a				
1–2 vertebrae	33 (46)	172 (54)	Reference	
≽3 vertebrae affected ^b	39 (54)	146 (46)	1.4 (0.8–2.3)	P = 0.24
Spine deformity ^c				
No kyphosis	14 (19)	72 (23)	Reference	
Kyphosis	60 (81)	243 (77)	1.3 (0.7-2.4)	P = 0.54
Spine surgery				
Minor procedure/none ^d	39 (53)	239 (75)	Reference	
Corrective surgery	35 (47)	80 (25)	2.7 (1.6–4.5)	P = 0.0003

an = 72 children and 318 adults due to missing data.

dNo spine surgery (n = 13 (18%) children, n = 3 (1%) adults), biopsy or other minor procedure (n = 26 (35%) children, n = 236 (74%) adults).

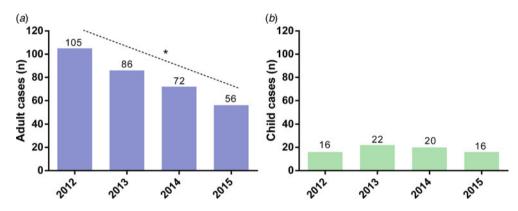


Fig. 3. Number of (A) adults and (B) children diagnosed with spinal tuberculosis at tertiary hospitals in the Western Cape between January 2012 and December 2015 *Significant trend for decrease in cases over the study period (*P* = 0.04).

detectable VLs (Table 2). Although the majority had CD4 counts <500 cells/μl, only 23% had severe immunosuppression. Those with CD4 count <200 cells/μl did not show increased odds of having ≥3 vertebrae affected or receiving corrective surgery. However, variation in the timing of CD4 tests and little sensitivity in the measure of disease severity limit the interpretation of these findings.

In addition to the findings of a large burden of cases, relatively young median age and high HIV prevalence, the current study suggests that many adults present with relatively advanced spinal TB disease, given that 46% of adult cases had three or more vertebrae affected and 77% had kyphotic deformity (Table 3). Furthermore, 25% of adults required corrective surgery, on the basis of indications such as spinal instability or progressive neurological deficit [17]. Notably, those who received corrective surgery comprised only 30% of all adults with kyphosis and it is assumed that surgery could not be justified in the remainder of the patients with spinal deformity. While treatment outcomes fell beyond the scope of the current study, it has previously been reported that at least 23% of patients managed non-surgically have kyphosis of ≥10 degrees at the affected level post-treatment [31]. This permanent deformity may or may not impact on the individual's physical function in the long term. Nevertheless, it is likely to

have significant cosmetic and psychological implications for the relatively young, economically active population concerned.

Spinal TB among children in the Western Cape

Approximately 19% of spinal TB cases in the current study were in children (≤14 years old) with 14% of all cases involving children younger than 10 years old. In contrast to previous findings of a large peak in all TB disease among 0-4 years old in our setting [32, 33], the current study found an equal distribution of spinal TB cases in 0-4 and 5-9 years old. This is likely due to a time delay in the onset of spinal TB after initial TB infection; Perez-Velez and Marais suggested that osteoarticular TB manifests 1-3 years after primary infection in children [34]. However, it is possible that delayed presentation to the health system and delayed diagnosis may also contribute to the observed distribution of cases. In comparison to young children, there were relatively fewer cases of spinal TB among adolescents (defined as 10-19 years), in keeping with previous findings of an increase in pulmonary rather than extra-pulmonary TB in this age group [33, 35].

Gender distribution among children was almost identical to that of adults with 58% of spinal TB cases in children involving

 $^{^{}b}$ Three to four vertebrae affected (n = 26 (36%) children, n = 94 (30%) adults), \geqslant 5 vertebrae affected (n = 13 (18%) children, n = 52 (16%) adults).

cn = 315 adults due to missing data.

females (Table 1). However, the smaller overall number of child cases translated into a smaller absolute difference in male vs. female cases and limits the extent to which this gender distribution can be interpreted. In a similar way, only two of the children in the current study were known to be HIV-positive and the relationship between HIV status and gender in children could not be assessed. The low prevalence of HIV among children in the current study is in keeping with the previous reports of a 2% HIV prevalence among South African children ≤14 years of age [30]. However, it is unclear whether this is a true reflection of HIV prevalence within this group given that HIV status was unknown in 28% of children. The relatively large proportion of children with unknown HIV status is of major concern, as young children remain at risk for late diagnosis of perinatal HIV infection [36]. While it is possible that evidence of HIV testing may have existed beyond the records available for the current study, previous studies in our setting support a comparatively high proportion of unknown HIV status among children under 5 years old [9, 33].

The majority of children with spinal TB could be described as having severe disease with 54% having ≥3 vertebrae affected, 81% presenting with kyphosis and 47% receiving corrective surgery (Table 3).

While children were no more likely than adults to have ≥3 vertebrae affected or to have kyphotic deformity, they were almost three times more likely to have received corrective surgery (Table 3). The most probable explanation for this is that active spinal TB disease progresses more rapidly in children than in adults with a particularly high risk of deformity among children younger than 10 years old [15]. The current study recorded kyphotic deformity as a binary outcome rather than quantifying it and one would expect that deformity would have been more severe among children, increasing the risk of spinal instability or progressive neurological deficit. Furthermore, in contrast to adults, children remain at risk of progressive deformity through disproportional growth of vertebral remnants after the active stage of the spinal TB disease [15]. It follows that increased corrective surgery among children may also partly be explained by surgery to prevent severe deformity in the future. Risk factors for severe deformity may be identified very early in the course of the disease and include age below 10 years, cervical thoracic or thoracolumbar junctional lesions and the well-established 'spine at risk' signs described by Rajasekaran [15].

Trends in the burden of spinal TB

The current study found that adult cases of spinal TB showed a significant decline between 2012 and 2015, whereas cases of spinal TB among children showed no clear trend over the study period (Fig. 3). These patterns were similarly reflected in incidence measures with the incidence of spinal TB among adults decreasing from 2.57 per 100 000 in 2012 to 1.22 per 100 000 in 2015. While these values are broadly similar to the 1.32 spinal cases per 100 000 reported among adults in KwaZulu-Natal in 2005–2006 [25], factors such as changes in TB control programmes and the provision of ART in the intervening years limit the scope for further incidence comparison.

A decline in spinal TB cases is consistent with a decrease in the incidence of all TB cases at both national level [37, 38] and in the Western Cape [39] during the period of study – progress that may be largely attributed to the National Strategic Plan for HIV, STIs and TB (2012–2016) [40]. This plan was developed as the next phase of response to the HIV and TB epidemics within the

country and some key applications have included the introduction of the Xpert test for TB diagnosis from 2011, increased TB screening and prophylaxis in high-risk populations and increased ART coverage [41]. While these factors are likely to contribute to a decrease in the incidence of all TB cases, it could be speculated that increased ART coverage may be particularly relevant to decrease the incidence of extra-pulmonary TB; earlier initiation of ART would be expected to protect against severely impaired immunity and increased risk of disseminated TB disease among HIV-positive individuals [42].

Unfortunately there is little data available on other types of extra-pulmonary TB in South Africa with which to compare the current trends in child and adult cases of spinal TB. A retrospective review of TB registers at hospitals in the rural Mopani District reported an increase in bone and joint TB from five to 13 cases between 2009 and 2013 [43]. However, the study had a number of limitations and it is unclear whether the change in bone and joint TB should be regarded as meaningful. Data from the Electronic Treatment Register (ETR) in the Western Cape suggest a decrease in all new TB cases among children ≤14 years (6 350 cases in 2012 vs. 5 095 cases in 2015) (unpublished data, Western Cape Department of Health). However, in principle, it is likely to be more difficult to observe trends among child cases of spinal TB given the small absolute numbers involved. A further explanation could be that although the incidence of TB has decreased in recent years, it remains exceptionally high and much larger decreases may be necessary before the benefit can be seen among the vulnerable paediatric population.

The current study focused on trends in absolute case numbers and it fell beyond the scope of the study to estimate trends in the proportion of spinal TB among all TB cases. Some studies from developed countries have reported an increase in extra-pulmonary TB as a proportion of all TB cases [44–46]. However, this trend has been attributed to an increase in immigrant populations from Africa and elsewhere and is therefore not necessarily applicable in the developing world. WHO estimates suggest that the proportion of exclusively extra-pulmonary TB cases in South Africa decreased from 14% in 2012 to 12.7% in 2013 and 10% in 2015 [37, 38, 47].

In the case of spinal TB, previous studies from developed countries are inconclusive with some suggesting an increase [6, 44] and others a decrease [4] in spinal TB cases. At present, trends in the proportion of spinal TB among all TB cases in South Africa and other developing countries remain unclear, partly due to reporting methods that tend to group all types of extra-pulmonary TB together and to record only pulmonary TB when patients have concomitant pulmonary and extra-pulmonary infection. In the future, increased differentiation for reporting severe forms of TB would provide greater insight into the true impact of TB and allow for trends over time to be established.

Limitations

Limitations of the current study include the retrospective nature of the study and the fact that it relied on possible spinal TB patients being referred to tertiary orthopaedic units for diagnostic confirmation. It is possible that some patients may have been diagnosed presumptively without tertiary referral, although these numbers should be low. The current study also does not account for any cases of spinal TB diagnosed in the private health sector. However, these numbers would be expected to be very low, given that only 16% of South Africans belong to private medical

schemes [48] and that these individuals are likely to be better resourced with less of the socio-economic adversity associated with TB. Nevertheless, the current findings should be taken to represent the minimum burden for spinal TB in the province rather than the total burden.

Other limitations of the study include the fact that it did not include information about previous episodes of TB or symptoms at presentation, may have overestimated the number of cases with unknown HIV status and had incomplete information regarding the state of HIV illness among HIV-positive individuals. Finally, it is noted that the number of bacteriologically confirmed cases may have been higher had the Xpert test been in use for the whole study period rather than primarily 2014 and 2015.

Conclusion

The current study demonstrated a high absolute burden of spinal TB at tertiary hospitals in the Western Cape in recent years with 19% of cases involving children. Adult cases appear to have declined between 2012 and 2015, although the trend among child cases is less obvious. While this apparent progress is encouraging, the number of cases remains considerable and severe presentation requiring corrective surgery was common, particularly in children. It follows that spinal TB remains a public health concern in the Western Cape, with increased vigilance required for earlier diagnosis, especially among children.

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Conflict of interest. None.

References

- Abdool Karim SS et al. (2009) HIV infection and tuberculosis in South Africa: an urgent need to escalate the public health response. The Lancet 374, 921–933.
- World Health Organization (2017) Global Tuberculosis Report 2017.
 Geneva: World Health Organization. http://www.who.int/tb/publications/global_report/archive/en/ (Accessed 12 January 2018).
- Massyn N et al. (ed.) (2016) District Health Barometer 2015/16. Health Systems Trust. Durban: Health Systems Trust. http://www.hst.org.za/publications/Pages/-District-Health-Barometer-201516.aspx (Accessed 12 January 2018).
- De la Garza Ramos R et al. (2017) The epidemiology of spinal tuberculosis in the United States: an analysis of 2002-2011 data. Journal of Neurosurgery Spine 26, 507-512.
- Johansen IS et al. (2015) Characteristics and clinical outcome of bone and joint tuberculosis from 1994 to 2011: a retrospective register-based study in Denmark. Clinical Infectious Diseases 61, 554–562.
- Jutte PC et al. (2004) Increase of bone and joint tuberculosis in The Netherlands. Journal of Bone and Joint Surgery British Volume 86, 901–904.

 Held M et al. (2014) Genexpert polymerase chain reaction for spinal tuberculosis: an accurate and rapid diagnostic test. The Bone & Joint Journal 96–B, 1366–1369.

- Watt JP and Davis JH (2014) Percutaneous core needle biopsies: the yield in spinal tuberculosis. South African Medical Journal 104, 29–32.
- Held MFG et al. (2017) Epidemiology of musculoskeletal tuberculosis in an area with high disease prevalence. Asian Spine Journal 11, 405–411.
- Rajasekaran S, Kanna RM and Shetty AP (2014) Pathophysiology and treatment of spinal tuberculosis. JBJS Reviews 2, e4.
- Batirel A et al. (2015) The course of spinal tuberculosis (Pott disease): results of the multinational, multicentre backbone-2 study. Clinical Microbiology and Infection 21, 1008e9–1008e18.
- Shi T et al. (2016) Retrospective study of 967 patients with spinal tuberculosis. Orthopedics 39, e838–e843.
- Turgut M (2001) Spinal tuberculosis (Pott's disease): its clinical presentation, surgical management, and outcome. A survey study on 694 patients. Neurosurgical Review 24, 8–13.
- Jain AK et al. (2014) Tubercular spondylitis in children. Indian Journal of Orthopaedics 48, 136–144.
- Rajasekaran S (2013) Natural history of Pott's kyphosis. European Spine Journal 22, 634–640.
- Dunn R and Zondagh I (2008) Spinal tuberculosis: diagnostic biopsy is mandatory. South African Medical Journal 98, 360–362.
- Mak KC and Cheung KMC (2013) Surgical treatment of acute TB spondylitis: indications and outcomes. European Spine Journal 22, 603-611
- Held M et al. (2018) Most influential literature in spinal tuberculosis: a global disease without global evidence. Global Spine Journal 8, 84–94.
- World Health Organization (2013) Definitions and Reporting Framework for tuberculosis-2013 Revision. Geneva. http://www.who.int/tb/publications/definitions/en/ (Accessed 12 January 2018).
- Statistics South Africa (2011) Mid-year Population Estimates 2011.
 Statistical Release P0302. Pretoria. http://www.statssa.gov.za/publications/ P0302/P03022011.pdf (Accessed 6 August 2018).
- Statistics South Africa (2013) Mid-year Population Estimates 2013.
 Statistical Release P0302. Pretoria. https://www.statssa.gov.za/publications/P0302/P03022013.pdf (Accessed 6 August 2018).
- Statistics South Africa (2014) Mid-year Population Estimates 2014.
 Statistical Release P0302. Pretoria. https://www.statssa.gov.za/publications/P0302/P03022014.pdf (Accessed 6 August 2018).
- Statistics South Africa (2015) Mid-year Population Estimates 2015.
 Statistical Release P0302. Pretoria. https://www.statssa.gov.za/publications/P0302/P03022015.pdf (Accessed 6 August 2018).
- Mwachaka PM et al. (2011) Spinal tuberculosis among human immunodeficiency virus-negative patients in a Kenyan tertiary hospital: a 5-year synopsis. Spine Journal 11, 265–269.
- Godlwana L et al. (2008) Incidence and profile of spinal tuberculosis in patients at the only public hospital admitting such patients in KwaZulu-Natal. Spinal Cord 46, 372–374.
- Daniel K and Dunn R (2013) Comparison of platelet count in tuberculosis spine to other spine pathology. European Spine Journal 22, 2810–2814.
- Naing C et al. (2013) Meta-analysis: the association between HIV infection and extrapulmonary tuberculosis. Lung 191, 27–34.
- 28. **Shivakoti R** *et al.* (2017) Association of HIV infection with extrapulmonary tuberculosis: a systematic review. *Infection* **45**, 11–21.
- Western Cape Provincial Aids Council (2016) Annual Progress Report 2014/15, Provincial Strategic Plan 2012-2016. http://sanac.org.za/wp-content/uploads/2016/04/WC_PSP-ANNUAL-PROGRESS-REPORT-Final.pdf (Accessed 6 August 2018).
- Shisana O et al. (2014) South African National HIV Prevalence, Incidence and Behaviour Survey, 2012. Cape Town: HSRC Press. http://www.hsrc.ac. za/en/research-data/view/6871 (Accessed 12 January 2018).
- Nene A and Bhojraj S (2005) Results of nonsurgical treatment of thoracic spinal tuberculosis in adults. The Spine Journal 5, 79–84.
- Snow K et al. (2017) Tuberculosis in adolescents and young adults: epidemiology and treatment outcomes in the Western Cape. The International Journal of Tuberculosis and Lung Disease 21, 651–657.

33. Wood R et al. (2011) Burden of new and recurrent tuberculosis in a major South African city stratified by age and HIV-status. PLoS ONE 6, 1-9.

- Perez-Velez CM and Marais BJ (2012) Tuberculosis in children. New England Journal of Medicine 367, 348–361.
- 35. Marais BJ et al. (2004) The natural history of childhood intra-thoracic tuberculosis: a critical review of literature from the pre-chemotherapy era. The International Journal of Tuberculosis and Lung Disease 8, 392–402.
- 36. Byamungu LN et al. (2018) Timing of HIV diagnosis in children with tuberculosis managed at a referral hospital in Cape Town, South Africa. The International Journal of Tuberculosis and Lung Disease 22, 488–495.
- 37. World Health Organization (2013) Global Tuberculosis Report 2013. Geneva. http://www.who.int/tb/publications/global_report/archive/en/ (Accessed 12 January 2018).
- 38. World Health Organization. (2016) Global Tuberculosis Report 2016. Geneva. http://www.who.int/tb/publications/global_report/en/ (Accessed 12 January 2018).
- Massyn N et al. (ed.) (2015) District Health Barometer 2014/2015. Health Systems Trust. Durban: Health Systems Trust. http://www.hst.org.za/publications/Pages/HSTDistrictHealthBarometer.aspx (Accessed 12 January 2018).
- South African National AIDS Council. National Strategic Plan on HIV, STIs and TB (2012-2016). http://www.gov.za/sites/www.gov.za/files/natio-nalstrategicplanonhivstisandtb_0.pdf (Accessed 12 January 2018).

- Churchyard GJ et al. (2014) Tuberculosis control in South Africa: successes, challenges and recommendations. South African Medical Journal 104, 244–248.
- Jones BE et al. (1993) Relationship of the manifestations of tuberculosis to CD4 cell counts in patients with human immunodeficiency virus infection. American Review of Respiratory Disease 148, 1292–1297.
- Hoogendoorn JC et al. (2017) Reduction in extrapulmonary tuberculosis in context of antiretroviral therapy scale-up in rural South Africa. Epidemiology and Infection 145, 2500–2509.
- Kruijshaar ME and Abubakar I (2009) Increase in extrapulmonary tuberculosis in England and Wales 1999-2006. Thorax 64, 1090-1095.
- Peto HM et al. (2009) Epidemiology of extrapulmonary tuberculosis in the United States, 1993–2006. Clinical Infectious Diseases 49, 1350– 1357.
- Sandgren A, Hollo V and van der Werf MJ (2013) Extrapulmonary tuberculosis in the European union and European economic area, 2002 to 2011. Eurosurveillance 18, 1–9.
- World Health Organization. (2014) Global Tuberculosis Report 2014.
 Geneva. http://www.who.int/tb/publications/global_report/archive/en/ (Accessed 12 January 2018).
- Council for Medical Schemes. (2016) Annual Report 2015/16. Pretoria. http://pmg-assets.s3-website-eu-west-1.amazonaws.com/CMS_Annual_ Report_2015-2016.pdf (Accessed 6 August 2018).