Osteosarcoma outcomes at a South African tertiary hospital

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Background. Osteosarcoma is the most common primary malignant bone tumour. There is a high incidence of late presentation in the developing world, posing additional challenges in the treatment of this aggressive disease.

Objective. To evaluate clinical outcomes of patients with osteosarcoma at Charlotte Maxeke Johannesburg Academic Hospital (CMJAH), a tertiary hospital in South Africa, and compare these with similar studies in the developing world.

Methods. This was a retrospective study of 61 patients treated at CMJAH between 2007 and 2011, with a minimum follow-up of 1.3 years.

Results. An average of 4.5 months elapsed before patients were first seen at the CMJAH tumour unit. Fifty-eight patients (95.1%) initially sought conventional medical care. Three patients (4.9%) presented with pathological fractures. All the patients underwent biopsy, performed an average of 3 weeks after arrival at the tumour unit. In most cases the delay was due to limited access to magnetic resonance imaging. Most patients (n=41, 67.2%) were at Enneking stage 2B, 4 (6.6%) were at stage 2A and 16 (26.2%) were at stage 3. Of the patients, 13 (21.3%) underwent limb salvage procedures, 33 (54.1%) had amputations, 4 (6.6%) refused further treatment and 11 (18.0%) received palliative care only; 55 patients (90.2%) received chemotherapy. Two patients developed local recurrence, one of whom had an amputation and the other further wide excision. Two patients received palliative radiotherapy. Of the patients, 82.0% were HIV-negative, 4.9% HIV-positive and the rest of unknown status. At the time of the study, all but two patients, who came from other countries, were traceable or known to have died. Our overall 1-year and 5-year survival rates were 62.7% (95% confidence interval (CI) 49.1 - 73.9) and 38.1% (95% CI 24.6 - 51.4), respectively. Male patients and those with a higher Enneking stage had a poorer prognosis.

Conclusion. Although most patients sought conventional medical care, unacceptable delays worsened survival. However, our survival rates are better than those in other developing countries. We advocate that professional, public and political awareness of osteosarcoma be improved as a matter of urgency, to facilitate rapid tertiary referral and expedite management.

Osteosarcoma is the most common primary malignancy of bone.1 It is relatively rare, however, with an incidence of 4 per million population per annum.2-5 The peak incidence is in the second decade of life, with a smaller peak in the 7th and 8th decades, these cases usually being secondary.2-4 The tumour is more common in males than in females,5,6 and has a higher prevalence among blacks than among whites.5 The cause is unknown, but individuals with germline mutations in the retinoblastoma (Rb) and tumour protein p53 (TP53) genes have been shown to be at increased risk.7,8 Survive was as low as 5% in the 1950s, with a 5-year survival rate of only 22%, most patients undergoing amputation and adjuvant radiation only or pioneering chemotherapy.9,10 During the 1970s there was a marked improvement in 5-year survival rates, which were reportedly as high as 81.6% in one study,11,12 largely owing to significant advances in chemotherapy, with utilisation of high-dose and multiagent regimens. In addition, tumour necrosis induced by neoadjuvant chemotherapy assisted in reducing the size of the surgical margin required for wide tumour excision, facilitating successful limb salvage without negatively affecting overall survival.13-15

Unfortunately, improvements in osteosarcoma management that are routine in the developed world are still not common in developing countries, because of limitations in human resources and lack of equipment and access to new and more effective chemotherapy regimes. Furthermore, it has been reported that cultural beliefs and reliance on traditional medicine in developing countries have resulted in late presentation to conventional hospitals, resulting in poorer outcomes.16-17 Reliable data from clinical outcome and epidemiological studies in developing countries are also scarce.17 Consequently, accurate benchmarking of evidence-based management protocols is extremely difficult to achieve. Muthupei and Mariba18 reported a 5-year survival rate of 7.5% in their 2000 study of 66 osteosarcoma patients at a tertiary hospital in South Africa (SA), which is comparable to outcomes from the 1950s in developed countries.19 In 2010, Shipley and Beukes20 reported on 30 patients with osteosarcoma treated at another SA tertiary institution. Half of their patients presented with metastases.20 The majority of patients in these studies from developing countries presented with advanced disease, to which the authors attributed their poor outcomes compared with the far better survival rates in developed countries.21-22 Noor et al.22 had similar findings in a study in Cambodia, with a 5-year survival rate of only 8%, citing late presentation and cultural preferences as the main factors responsible for the poor outcomes.22

Objective
To evaluate clinical outcomes of patients with osteosarcoma at Charlotte Maxeke Johannesburg Academic Hospital (CMJAH), a tertiary hospital in SA, and compare these with similar studies
from the developing world to evaluate factors contributing to any differences encountered.

**Methods**

This was a retrospective study of 61 patients treated for osteosarcoma at the CMJAH tumour unit over the 5-year period January 2007 - December 2011. The minimum follow-up was 1.3 years (mean 3.1, range 1.3 - 6.3). CMJAH is a level 1 trauma centre and one of the two main teaching hospitals of the University of the Witwatersrand, catering not only for the city of Johannesburg but also for Gauteng Province and neighbouring provinces and countries. Only patients with biopsy-confirmed osteosarcoma were included in the study.

Ethics clearance to conduct the study was obtained from the University of the Witwatersrand (ref. no. M150783). The data were obtained from theatre registry and patient records. All patients admitted to the tumour unit had a comprehensive form (available from the first author at lisendal@hotmail.com) completed on admission, throughout the course of admission and during follow-up. The tumour form had three main sections: (i) history and clinical findings; (ii) investigations (blood tests and radiography) and procedures (biopsy/surgical operations) done; and (iii) follow-up. All patients received a standard osteosarcoma work-up consisting of blood investigations and imaging before biopsy and then definitive treatment. Most patients were followed up at orthopaedic and oncology outpatient clinics, and those who had defaulted were contacted telephonically.

The main outcome measures were demographics, tumour stage according to the Enneking staging system, and treatment options offered (surgery (amputation and limb salvage), chemotherapy, radiotherapy) and survival. Survival was calculated in years from the date of presentation to the time of the study. Kaplan-Meier analysis was used to portray survival functions. Equality of survivor functions was investigated using the log-rank test.

**Results**

A total of 61 patients were included in the study (41 males and 20 females, male/female ratio 2:1). The mean age was 19.4 years (range 7 - 48). Most of the patients (n=36, 59.0%) were in their second decade, followed by the third decade (n=13, 21.3%) and then the first (n=5, 8.2%). The fourth and fifth decades accounted for 4 (6.6%) and 3 (4.9%) patients, respectively.

Of the patients, 43 (70.5%) and 15 (25.6%) initially sought care from their local clinic and general practitioner, respectively, before being referred to the tumour unit. Only 3 (4.9%) went to a traditional healer before being referred. On average, it took 4.5 months for patients to be seen in the tumour unit for the first time. Most patients (n=42, 68.9%) were from the CMJAH catchment area of Gauteng Province, the other SA provinces and other countries accounting for 16 (26.2%) and 3 (4.9%), respectively.

The majority of the patients (n=58, 95.1%) presented with a painful swollen limb and the rest (n=3, 4.9%) with pathological fractures. The tibia was the most common site of involvement, accounting for 34 (55.7%) cases, the femur and humerus accounting for 19 (31.1%) and 12 (19.7%), respectively. Only 3 patients (4.9%) were HIV-positive; 8 (13.1%) were of unknown status, and the rest (n=50, 82.0%) were HIV-negative.

Biopsy was done at a mean of 3 weeks after presentation. In most cases, the delay was because magnetic resonance imaging (MRI) was not readily available. The majority of the patients (n=41, 67.2%) were at Enneking stage 2B, 4 (6.6%) were at stage 2A, and 16 (26.2%) presented with metastases (stage 3). Conventional osteosarcoma was the most common histological type, accounting for 93.4% of cases (Table 1).

In 46 cases (75.4%) surgery was undertaken with the aim of achieving wide local resection margins (Table 2). Of these 46 patients, 13 (28.3%) had limb salvage procedures and 33 (71.7%) had amputations. Four patients (6.6%) refused any form of treatment despite multidisciplinary counselling and were discharged from the hospital for home-based care, and the rest (n=11, 18.0%) received palliative care (palliative care only without amputation) because they presented late (these included two patients who received radiotherapy).

Fifty-five patients (90.2%) received chemotherapy. Of these, 52 (94.5%) had both neoadjuvant and post-adjuvant chemotherapy and 3 (5.5%) post-adjuvant chemotherapy only. Palliative radiotherapy was administered to 2/61 patients (3.3%) who were not fit for surgery. Two patients developed local recurrence, of whom one was treated with amputation and the other with further excision.

At the time of the study, 59 patients (96.7%) were traceable or known to have died. The two untraceable patients were from other countries and had been referred home for further treatment after limb salvage. Of the patients who were alive (25/59, 42.4%), 10/13 (76.9%) had been treated with limb salvage and 14/33 (42.4%) with amputation, and the rest (1/4, 25.0%) had refused surgery. Of the 34/59 patients who were dead, 1/11 (9.1%) had been treated with limb salvage and 19/33 (57.6%) with amputation, and 3/4 had refused surgery (Table 2). Of the 13/61 patients (21.3%, including the two untraceable patients) who defaulted from follow-up or had refused treatment, 3/13 (23.1%) had been treated with limb salvage and 9/33 with amputation (27.3%).

Our 1-year survival rate was 62.7% (95% confidence interval (CI) 49.1 - 73.9) and our 5-year survival rate 38.1% (95% CI 24.6 - 51.4) (Fig. 1). Fig. 2 shows that there was a statistically significant

| Table 1. Histological typing (World Health Organization classification) (N=61) |
|----------------|-----------------|
| Type            | Patients, n (%) |
| Conventional    | 57 (93.4)       |
| Chondroblastic  | 9               |
| Fibroblastic    | 5               |
| Osteoblastic    | 22              |
| Not specified   | 19              |
| Periosteal      | 1 (1.6)         |
| Parosteal       | 2 (3.3)         |
| Telangiectatic  | 1 (1.6)         |
| Small cell, low-grade surface, secondary | 0 |

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<th>Table 2. Survival v. treatment options</th>
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<td>Limb salvage</td>
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1- and 5-year survival rates of 25.8% and 7.5%, while Noor et al.\[17\] in countries:\[15-17\] Muthuphei and Mariba’s\[15\] SA study in 2000 had proximal tibia and was treated with above-knee amputation (AKA). She was a 12-year-old girl who had a pathological fracture of the left pathological fracture was alive, without recurrence at 54 months. diagnosed with conventional osteosarcoma of the proximal tibia, a 9-month history of a painful swollen knee and was subsequently Enneking stage 2B. delays may be attributable to poor accessibility of health facilities, especially for individuals living in rural areas. The higher the Enneking stage on presentation the poorer the survival according to time (in months) before arrival at CMJAH did not differ significantly (p=0.587) (Fig. 3). All the patients who were staged as Enneking 2A were alive at 5 years (range 5.08 - 6) without recurrences. Patients with Enneking stage 2B disease had 1-year and 5-year survival rates of 69.2% (95% CI 52.2 - 81.2) and 47.9% (95% CI 30.4 - 63.4), respectively. For patients with Enneking stage 3 disease, the 1- and 5-year survival rates were 37.5% (95% CI 15.4 - 59.7) and 3.75% (95% CI 0.06 - 23.4), respectively (Fig. 4). This difference is statistically significant (p=0.0008).

Only one patient who refused treatment was alive, 24 months after discharging herself from the hospital. She had presented with a 9-month history of a painful swollen knee and was subsequently diagnosed with conventional osteosarcoma of the proximal tibia, Enneking stage 2B.

Only one of the three patients who initially presented with a pathological fracture was alive, without recurrence at 54 months. She was a 12-year-old girl who had a pathological fracture of the left proximal tibia and was treated with above-knee amputation (AKA).

**Discussion**

Our 1-year and 5-year overall survival rates of 62.7% and 38.1% are better than those in previous published studies from developing countries.\[15-17\] Muthuphei and Mariba’s\[15\] SA study in 2000 had 1- and 5-year survival rates of 25.8% and 7.5%, while Noor et al.\[17\] reported 1- and 5-year survival of 53.6% and 8% in Cambodia.\[15,17\] Our 5-year survival figure is worse than the corresponding figures for developed countries, which average >50%.\[18-20\] Several factors may account for our relatively poor outcomes. Our patients presented on average 4.5 months after the onset of symptoms, with Enneking stages 2B and 3 accounting for a combined 93.4% compared with 6.6% with stage 2A. The delays may be attributable to poor accessibility of health facilities, especially for individuals living in rural areas. The higher the Enneking stage on presentation the poorer the prognosis, as has been reported in the literature\[18\] and shown by our results. Although only 4.9% of patients admitted to first attending a traditional healer, most patients would have consulted them, as evidenced by the presence of ‘traditional’ scarifications. Almost a third of the patients came from outside our catchment area, which could have led to delays in referral. Limited access to MRI in our hospital, leading to a further average delay of 3 weeks, exacerbated the situation. Biopsies were done approximately a week after the MRI and other investigations were completed, resulting in an overall mean time to diagnosis of 23.5 weeks. In contrast, the median time to diagnosis in developed countries has been reported as 9 weeks.\[21\] Patients from rural areas depend on doctors in district hospitals having a high index of suspicion of ‘painful limbs’ and doing initial investigations (X-rays) of the affected limb. Improving awareness through education of the community at large, and district doctors in

**Fig. 1.** Overall Kaplan-Meier survival estimate in years.

**Fig. 2.** Overall Kaplan-Meier survival estimates by sex.

**Fig. 3.** Overall Kaplan-Meier survival estimates by time that elapsed before patients presented at Charlotte Maxeke Johannesburg Academic Hospital.

**Fig. 4.** Overall Kaplan-Meier survival estimates by Enneking stage.
particular, could increase early referral to specialised tumour units and hence the cure rate.

Amputees had poorer survival rates than patients who underwent limb salvage, because they had more advanced disease. Amputation was offered as ‘palliative surgery’ to relieve malaise and for pain control. More amputees than patients who had undergone limb salvage defaulted from follow-up at outpatient clinics (orthopaedic and oncology), despite multidisciplinary emphasis in hospital on surveillance for recurrence, which could explain their poor outcome. Preservation of the limb with limb salvage procedures, and hence a better self-image, could be a factor explaining the good attendance by these patients.

The majority of our patients (95.1%) presented with a painful swollen limb and the rest with pathological fractures. Histological conventional osteosarcoma accounted for 93.4% of cases, while the rest (periosteal, parosteal and telangiectatic) accounted for 6.6%, as shown in Table 1. The incidence of pathological fractures is reported to be 5 - 10% in the literature. Historically, pathological fractures were treated by amputation because of damage to the microcirculation that would result in contamination of adjacent soft tissue and metastasis. Several authors have since shown that there is no difference in terms of survival between patients who have amputation and those who have limb salvage, because the tumour encapsulated the neurovascular structures. She received neoadjuvant and post-adjuvant chemotherapy and was offered AKA because the tumour encapsulated the neurovascular structures. She was alive and disease free 54 months after AKA. Most of our patients presented with advanced disease precluding limb salvage.

Just over a quarter of our patients presented with metastases on admission. Our 5-year survival of 3.75% for this cohort of patients is very low compared with the mid-20s - 30% in developed countries. One patient, who was unfit for surgery, died few days after admission and the other patient, who underwent AKA, 2 months after admission. Only one patient with a pathological fracture who survived was a 12-year-old girl with high-grade osteosarcoma of the left distal tibia. She received neoadjuvant and post-adjuvant chemotherapy and was offered AKA because the tumour encapsulated the neurovascular structures. She was alive and disease free 54 months after AKA. Most of our patients presented with advanced disease precluding limb salvage.

One of the four patients who refused chemotherapy and surgery was traceable and alive 24 months after discharging herself. She was a 20-year-old woman with high-grade osteosarcoma of the left proximal tibia, Enneking stage 2B. She was being looked after by relatives at home.

Study limitations
Although 96.7% of our patients were traceable (or known to have died) at the time of the study, ~20% of them had defaulted from follow-up. This is a major limiting factor in our study. This high default rate could be due to difficulty in accessing our facility, and also to the fact that ~30% of our patients came from other provinces and countries. The majority of those who had defaulted cited lack of funds as the reason for not attending. Attempted telephonic contact of the two patients from other countries was unsuccessful as the numbers were not in use. Another limiting factor is the retrospective nature of the study, despite the fact that the data were collected prospectively using a comprehensive tumour form.

Conclusion
Treatment of osteosarcoma is challenging, demanding and time-consuming, and requires meticulous record keeping. Limited resources hamper a multidisciplinary approach in developing countries. In our setting, both patients and their families need extensive education and counselling to improve access to modern treatments.

Despite better overall 1-year and 5-year survival rates than in other developing countries, our 5-year overall survival rates are suboptimal compared with developed countries, mainly because of delayed referral and logistical constraints to rapid work-up. We strongly recommend improved professional and public awareness of the importance of rapid referral, which is vital for the survival of these patients.

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Author contributions.
LL, ZAL, RDK and ML participated in the planning and design of the study and in interpretation of the results. LL wrote the first draft, and RK and ML edited it and supervised the project. All authors revised, edited and approved the final draft.

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