Establishing musculoskeletal oncology service in resource constrained country: challenges and solutions

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Abstract
The burden of orthopedic tumor surgery in Pakistan is not known. Similarly the number of procedures being performed for bone and soft tissue surgery are not known. This is even becoming more challenging where the existence of rules and regulations in health care are next to minimal. Furthermore data recording in our country and case registries hardly exist. Despite the lack of information and resources, with high disease burden on community, various providers provide surgical interventions every day in our settings. A lot of tumor surgery is still being done by general surgeons and general orthopedic surgeons who have little knowledge and update about musculoskeletal oncology principles. Lack of subspecialized centers and the high cost of such centers force the patients to visit these surgeons for a highly sophisticated problem like a bone tumor which is the disease of young bones. In this article we will emphasize on the difficulty in establishing an orthopedic tumor service in our part of the world and the consequences including delay in diagnosis, faulty course of management and later decline in functionality, disease progression and increased mortality. We will highlight the principles and stepwise approach of orthopedic tumor surgery and explain the difficulty encountered if these principles are not followed.

Keywords: musculoskeletal, tumors, oncology, developing country

Malignant tumors are an overwhelming challenge to the orthopedic surgeons in a developing country due to inadequate awareness and limited diagnostic and therapeutic facilities. Another issue is the lack of the specialized orthopedic surgeons in tumor surgery in our side of the world, along with a scarcity of relevant published literature.

Orthopedic oncology is a highly sophisticated subspecialized field that requires a long and strenuous fellowship training in diagnosing and managing primary benign and malignant bone and soft tissue tumors. Even tumor surgery fellowships in North America typically produce no more than 12–15 new tumor surgeons each year[1–3]. Although general orthopedic surgeons maybe qualified to undertake surgical intervention of these tumors, but it is advisable to involve an orthopedic oncologist from the start to minimize the chances of bad outcome.

As primary bone sarcomas usually affect young individuals, an amputation could mean a lifelong dependency and disability hence decreased quality of life. In a society like ours where young people are responsible for entire households, it is imperative to restore their maximum physical capability and avoid amputation. This is now possible with a combination of recent innovations in understanding tumor biology and limb-salvage techniques supplemented by improved techniques in histopathology, radiology, radiotherapy, and medical oncology.

Because of lack of awareness on part of both patients and health care providers, interventions in tumor surgery are provided by many general orthopedic surgeons every day. This includes biopsy and tumor removal followed by referral to oncologist for chemotherapy and radiation. Coming from the developing country settings, knowing the prevalence and the scope of surgical disease is critical to planning further options. This stems largely from lack of awareness and financial gains to be credited after every procedure.

Impact of untreated or wrongly treated tumors, on disability, premature morbidity and mortality, presents a great challenge. Owing to an overall lack of knowledge and experience in this highly specialized field, surgeons end up performing surgeries with misplaced and miscalculated incisions, avoidable amputations or just treating patients on the basis of Tru-cut biopsy results from nonaccredited histopathology laboratory and untimely referrals to oncologists. A part of this problem lies with the patients who delay a visit to the surgeon or simply ignore the advice of referral to a specialist.

In efforts to avoid the above, development of treatment strategies have occurred in the past several years. Management starts stepwise from the time patient present in clinic, relevant imaging, staging, planned biopsy and incision site and size followed by appropriate treatment options. Interruption in this stepwise
approach would lead to increased sufferings, disability, cost involved, and additional burden on the health care systems.

The purpose of this article is to review the principles of management of bone and soft tissue tumors, increase the awareness to this highly subspecialized field and emphasize on the need to develop such service in our part of the world despite the constraints.

Discussion

Bone and soft tissue sarcomas are rare mesenchymal malignancies that arise in 2–4 per 100,000 head of population[4]. Overall survival following treatment of primary sarcoma now approaches 75% at 5 years, and surgery remains the mainstay of treatment[5,6].

Historically, mainstay of managing these tumors was amputation. But as bone tumors are the disease of young bones, amputation increases the disability for those patients who have survival long enough to justify complex surgery. At our side of the world, society follows a joint family system where the entire family depends on one breadwinner. So when the latter is disabled, the entire family is paralyzed, both socially and economically. Thus it is imperative to restore their maximum physical capability and avoid amputation.

Surgery to resect the tumor followed by reconstructions to preserve function, mobility, and esthetics (limb-sparing surgery) has now replaced amputation as the primary form of surgical intervention[7–9].

The major therapeutic goals are long-term survival, avoidance of local recurrence, maximizing function, and minimizing morbidity. Landmark trials conducted in the 1970s and 1980s at the National Cancer Institute showed equivalent survival outcomes between limb amputation and limb-sparing surgery combined with radiotherapy[10,11].

Good outcome in limb saving procedures depends on multiple factors which was extensively studied and proved in literature like tumor size, depth, histologic grade, anatomic site, and margin size[12,13]. Older age has been reported to be associated with lower survival rates[14,15]. Older patients tend to present with larger and higher grade tumor which possibly result in increased local recurrences[16].

At the other side, there are bad prognostic factors, yet avoidable. Examples include low degree of suspicion in plain radiographs, particularly if associated with history of trauma, or if suspected lesion is seen, doing the biopsy by nonexperienced personnel in the field of musculoskeletal tumors, late referral to the orthopedic oncologist and cancer center, and decreased awareness and knowledge by the patients, their relatives and even the physicians in account for the rarity of this disease.

Definite diagnosis is mandatory before any attempt at surgical intervention. In reality, this sometimes is difficult and patients with malignant bone disease can be misdiagnosed as having benign lesions. Tumors of an osteolytic behavior in their early stages can simulate giant cell tumor (Fig. 1). We recommend an open biopsy or frozen section diagnosis peroperatively. Such cases they present challenge even to the experienced orthopedic oncologist due to the violation of the tumor and the delay between the initial intralesional procedure and the proper definitive operation.

Another category of misdiagnosis reported is the radiographic findings misinterpreted as bone infection. They do not often go for biopsy and when healing is delayed it is attributed to the natural course of the disease[17].

A biopsy should be planned as carefully as the definitive procedure and should be done only after clinical and radiographic examinations are done. In principle, the same group that will be undertaking definitive treatment should perform the biopsy. Avoid transverse incisions because they increase the challenge for the limb salvage surgeon (Fig. 2).

It is worthy to mention here the principles of orthopedic tumors management and biopsy. A biopsy should be planned as carefully as the definitive procedure and should be done only after clinical and radiographic examinations are done. Biopsy incision and tract is considered contaminated with tumor cells and should be in the excised specimen. Transverse incisions should be avoided because they are extremely difficult or impossible to excise with the specimen. If a drain is used, it should exit in line with the incision so that the drain track also can be easily excised en bloc with the tumor[18].

Our limited knowledge on this issue arises from the epidemiological factors that included neglect, unawareness of the problem, low socio economic status and financial burden involved in seeking treatment, and limited diagnostic facilities. On top of that is the lack of subspecialized centers and orthopedic tumor surgeons. This leads to increasing disability and impairing the quality of life of our patients. By and large, such tumors carry poor prognosis with high morbidity and mortality.

Cause of delay in seeking medical advice was neglect by the patient and family due to financial constraints, culture, lack of access to health care facilities, consultation with traditional bone settlers and even misdiagnosis by general orthopedic surgeons.

Giving the rarity of these tumors, along with their wide range of occurrence in any part of the body, they are resected by physicians other than orthopedic oncologists with nonstandardized techniques and without keeping a suspicious of malignancy in mind and safe margins[19–21].

Proper management of these tumors require a multidisciplinary approach involving a qualified orthopedic oncology
surgeon who is familiar with limb-salvage procedures supplemented by improved techniques in histopathology, radiology, radiotherapy, and medical oncology. Early referral of these patients plays a vital role for better outcome. Because this is such an uncommon disease, it is so helpful to have guidelines like the National Comprehensive Cancer Network (NCCN) guidelines, because you are likely not going to see so many patients within your practice. Guideline states that when a patient younger than 40 years old presents to you with a bone pain, and you see an abnormal suspicious lesion on plain radiographs you should refer them to an orthopedic oncologist for biopsy and biopsy should be performed at the treating institution. If the patient is elder than 40 years, then they should be worked up for potential bone metastasis[22]. Davis and colleagues compared the outcomes of patients treated primarily in a cancer center versus those treated at noncancer centers who were referred after an unplanned excision. They found that the rate of a local recurrence was higher in the unplanned surgical excision group, particularly patients with a residual tumor in the resected specimen[23]. Goodlad and colleagues reported 95 resections in patients initially treated in noncancer centers from a series of 236 patients with soft tissue sarcomas. They found that 59% of those patients who had undergone unplanned resections had inadequate margins after the resection[24].

At our institute, 135 operated patients with soft tissue sarcomas evaluated for outcomes, in terms of local recurrence and metastasis rate, of reexcision following unplanned excision of the tumor at prereferral hospital, results compared with those of first-time planned surgery. We reported that local recurrence, metastasis, and mortality rates were higher in patients who underwent unplanned resections (21.4% vs. 14.3%, 13.7% vs. 8.3%, 13.7% vs. 9.5%, respectively)[18]. Lewis et al[21] showed that disease-specific metastasis-free survival rate was lower in patients who underwent resection as compared with those who underwent planned primary surgery.

The clinicians and the pathologists handling management responsibility must have high index of suspicion as to the nature of bone lesion in order to establish the diagnosis of bone tumors. This applies specifically to orthopedic tumors because behavior of such tumors is less aggressive than other visceral tumors in body so the aim of the tumor surgeon is usually toward limb salvage and better quality of life.

Limb-sparing surgery is the technique of choice for surgical management of limb sarcomas. In comparison to amputation, limb-sparing surgery has the same overall survival rate, higher patient satisfaction, lower energy expenditure for walking and a lower cost to the community[26]. Innovative techniques are available that may result in a functional limb[27,28].

Reconstruction techniques at our side of the world are technically demanding and very expensive for our patients who pay the whole medical course and services out of their own pocket. We reviewed 40 consecutive pediatric patients, aged 16 years or younger, with locally aggressive or malignant bone tumors treated with tumor resection, autoclaving and reimplantation of the orthotopic auto graft. Vascularized or nonvascularized fibular graft was used as a biological adjunct and fixation done with plate. We recommended a low cost alternate using the patient's own autoclaved tumor bone for skeletal reconstruction. It consists of excision, sterilization, and reimplantation. Having the advantages of biological reconstruction with a “custom fit” segment, providing anatomic site for muscles and tendons reattachment, avoiding immunological response or transmission risks, no bone banking required, cheap, convenient for the surgeon, less operating time comparing to other reconstructive procedures and having higher incidence of integration and healing than allografts. We have used it successfully in both pediatric and adult populations[29,30].

Bone and tissue bank forms a very essential back-up service for any musculoskeletal oncology service. Unfortunately we do not have a custom of organ donation by the deceased; hence bone banking is virtually nonexistent. Secondly, using implants in
pediatric tumor reconstruction is an extremely costly solution in our society. Growing implants may not provide all solutions to difficult pediatric problems. We have successfully used and published the use of fresh parental fibular allograft in reconstruction after limb salvage surgery. This innovative technique is again with low cost, and with minimal morbidity. We hardly saw any tissue reaction in any of our patients[31].

Data management in the form of a formal tumor registry is also being practiced within our section of orthopedics. This has helped us put out numerous publications in the field of musculoskeletal oncology[12–47]. We feel that now this experience needs to be replicated on a national level and we are making efforts in this regard.

We emphasize that the experts who received advanced training in Orthopedics tumor surgery should deal such cases. Orthopedics societies should guide surgeons to refer such cases to the concerned to save the lives and limbs of our patients.

Development of expert manpower resources was the first challenge. One of our surgeons (senior author) went abroad on Faculty Development Award provided by the hospital. He went to Mayo Clinic, Rochester, MN and then to Rizzoli Institute, Bologna, Italy. This experience helped him learn the latest techniques of limb salvage surgery. Since then he has been in various musculoskeletal oncology centers around the world to keep pace with latest developments in this field. Similarly, our histopathologist, went to Cleveland Clinic and then Mayo Clinic to gain deeper insight into pathologic diagnosis of bone and soft tissue disorders. Likewise we have teamed up with a radiologist who specializes in musculoskeletal magnetic resonance imaging and he is our partner in orthopedic tumor board. These key manpower developments have helped us build a team of experts who collectively decide the course of action required for a particular sarcoma patient.

It was after a long effort that we were able to establish a musculoskeletal tumor board many years back. This paved the way for combined management of such challenging cases. There was no example existing in the whole country when we started this activity. Now at least we have 3 orthopedic tumor boards running in different institutions in different parts of our country. We now have a regular monthly scheduled multidisciplinary tumor board meeting separately for adult and pediatric patients. This board includes the orthopedic tumor surgeon, senior medical and radiotherapy oncologists, pathologist, radiologist, residents and medical students as well. This board provides insights and feedback and approved beneficial for the patients, and for all members. Above all we believe that it is the right of every patient to be presented in such highly professional meeting and we invite cases from other institutions all over the country to be discussed in this board.

Conclusions

This review tried to highlight some of the causes of neglect in malignant bone and soft tissue tumors in our side of the world. We require a concrete effort from the Orthopedics societies and the government to create awareness among general physicians and surgeons to know the consequences of such neglect and early referral to the orthopedics tumor surgeons to save loss of extremities. Considering their very small numbers, we need to train more orthopedic oncology surgeons in Pakistan. They need to join hands in developing and maintaining a national Orthopedics tumor registry. This will help produce publications and reflect on our work periodically. Tumor board is very effective in increasing the knowledge and experience and above all improves patients’ outcome.

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Author contribution

O.H.A.H.: initial draft, writing of final manuscript. A.Z.: editing. Z.N.: contribution of clinical pictures and editing. M.U.: supervision of all steps of manuscript writing.

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References

[1] Miller BJ, Rajani R, Leddy L, et al. How much tumor surgery do early-career orthopaedic oncologists perform?. Clin Orthop Relat Res. 2015;473:695–702.

[2] White J, Toy P, Gibbs P, et al. The current practice of orthopaedic oncology in North America. Clin Orthop Relat Res 2010;468:2840–53.

[3] Enneking WF. An abbreviated history of orthopaedic oncology in North America. Clin Orthop Relat Res 2000;374:115–24.

[4] Charoenlap C, Imanishi J, Tanaka T, et al. Outcomes of unplanned sarcoma excision: impact of residual disease. Cancer Med 2016;5:980–8.

[5] Choong PF, Rudiger HA. Prognostic factors in soft-tissue sarcomas: what have we learnt? Expert Rev Anticancer Ther 2008;8:139–46.

[6] Clark JC, Dass CR, Choong PF. A review of clinical and molecular prognostic factors in osteosarcoma. J Cancer Res Clin Oncol 2008;134:281–97.

[7] Choong PF, Sim FH. Limb-sparing surgery for bone tumors: new developments. Semin Surg Oncol 1997;13:64–9.

[8] Federman N, Bernthal N, Eilber FC, et al. The multidisciplinary management of osteosarcoma. Curr Treat Options Oncol 2009;10:82–93.

[9] Papagelopoulos PJ, Galanis EC, Vlastou C, et al. Current concepts in the evaluation and treatment of osteosarcoma. Orthopedics 2000;23:858–67; quiz 868–9.

[10] Rosenberg SA, Tepper J, Glattstein E, et al. The treatment of softtissue sarcomas of the extremities: prospective randomized evaluations of (1)
limb-sparing surgery plus radiation therapy compared with amputation and (2) the role of adjuvant chemotherapy. Ann Surg 1982;196:305–15.

[11] Yang JC, Chang AE, Baker AR, et al. Randomized prospective study of the benefit of adjuvant radiation therapy in the treatment of soft tissue sarcomas of the extremity. J Clin Oncol 1998;16:197–203.

[12] Liu CY, Yen CC, Chen WM, et al. Soft tissue sarcoma of extremities: the prognostic significance of adequate surgical margins in primary operation and reoperation after recurrence. Ann Surg Oncol 2010;17:2102–11.

[13] Billingsley KG, Lewis JJ, Leung DH, et al. Multifactorial analysis of the survival of patients with distant metastasis arising from primary extremity sarcoma. Cancer 1999;85:389–95.

[14] Gronchi A, Casali PG, Mariani L, et al. Status of surgical margins and prognosis in adult soft tissue sarcomas of the extremities: a series of patients treated at a single institution. J Clin Oncol 2003;21:96–104.

[15] Biau DJ, Ferguson PC, Turcotte RE, et al. Adverse effect of older age on the recurrence of soft tissue sarcoma of the extremities and trunk. J Clin Oncol 2011;29:4029–33.

[16] Lahat G, Dhuka AR, Lahat S, et al. Factors that influence the outcome of limb salvage surgery for soft tissue sarcoma. Ann Surg Oncol 2009;16:2579–86.

[17] Leon DG, Lee SY, Kim JW. Bone primary sarcomas undergone unplanned intralesional procedures—the possibility of limb salvage and their oncologic results. J Surg Oncol 2006;94:592–8.

[18] Canale ST, Beaty JH. General principles of tumors. In: Canale ST, ed. Campbell’s Operative Orthopaedics (Vol 1, Part 8), 12th ed. Amsterdam; The Netherlands: Elsevier; 2013:794–5.

[19] Umer HM, Umer M, Qadir I, et al. Impact of unplanned excision on prognosis of patients with extremity soft tissue sarcoma. Sarcoma 2013;2013.

[20] Venkatesan M, Richards Cj, McCulloch TA, et al. Inadvertent surgical resection of soft tissue sarcomas. Eur J Surg Oncol 2012;38:346–51.

[21] Noria S, Davis A, Kandel R, et al. Residual disease following unplanned excision of a soft-tissue sarcoma of an extremity. J Bone Joint Surg Am 1996;78:650–5.

[22] Bower JE, Bak K, Berger A, et al. Screening, assessment, and management of fatigue in adult survivors of cancer: an American Society of Clinical oncology clinical practice guideline adaptation. J Clin Oncol 2014;32:1840–50.

[23] Davis AM, Kandel RA, Wunder JS, et al. The impact of residual disease on local recurrences in patients treated by initial unplanned resection for soft tissue sarcomas of the extremity. J Surg Oncol 1997;66:61–7.

[24] Goodlad JR, Fletcher CD, Smith MA. Surgical resection of primary soft tissue sarcoma. Incidence of residual tumour in 95 patients needing reexcision after local excision. J Bone Joint Surg Am 1996;78B:658–1.

[25] Lewis JJ, Leung D, Espan J, et al. Effect of resection in extremity soft tissue sarcoma. Ann Surg 2008;231:655–3.

[26] Choong PF, ed. Principles of limb-sparing surgery in bone and soft tissue sarcoma. Cancer Forum; 2010: The Cancer Council Australia.

[27] Kong GY, Rudiger HA, Ek ET, et al. Reconstruction after external hemipelvectomy using tibia-hindfoot rotationplasty with calcaneo-sacral fixation. Int Semin Surg Oncol 2008;5:1.

[28] Sm IW, Tse LF, Ek ET, et al. Salvaging the limb salvage: management of complications following endoprosthetic reconstruction for tumours around the knee. Eur J Surg Oncol 2007;33:796–802.

[29] Umer M, Umer HM, Qadir I, et al. Autoclaved tumor bone for skeletal reconstruction in paediatric patients: a low cost alternative in developing countries. BioMed Res Int 2013;2013.

[30] Bohm P, Fritz J, Theide S, et al. Reimplantation of extracorporeal irradiated bone segments in musculoskeletal tumor surgery: clinical experience in eight patients and review of the literature. Langenbecks Arch Surg 2003;387:355–65.

[31] Kadir MM, Askari R, Baz S. Use of fresh parental fibular allograft for reconstruction of skeletal defects after limb salvage surgery. J Pak Med Assoc 2014;64:S151.

[32] Sepah YJ, Umer M, Qureshi A, et al. Lymphangiosarcoma of the arm presenting with lymphedema in a woman 16 years after mastectomy: a case report. Cases J 2009;2:6887.

[33] Anjum N, Rashid H, Umer M. Use of spare parts in musculoskeletal oncology. Orthopaedia cases. In: Orthopaedia—Collaborative Orthopaedic Knowledgebase. 2010. Available at: www.orthopaedia.com/x/wwAiAg. Accessed January 19, 2012.

[34] Ansari TZ, Masood N, Parekh A, et al. Four year experience of sarcoma of soft tissues and bones in a tertiary care hospital and review of literature. World J Surg Oncol 2011;9:51.

[35] Qadir I, Umer M, Baloch N. Functional outcome of limb salvage surgery with mega-endoprosthetic reconstruction for bone tumors. Arch Orthop Trauma Surg 2012;132:1227–32.

[36] Fabbri N, Tiwari A. Extraskeletal osteosarcoma: clinicopathologic features and results of multimodal management. J Clin Oncol 2010;28:11.

[37] Qadir I, Umer M, Umer HM, et al. Managing soft tissue sarcomas in a developing country: are prognostic factors similar to those of developed world? W J Surg Oncol 2012;10:188.

[38] Abbas K, Umer M. Complex biological reconstruction after wide excision of osteogenic sarcoma in lower extremities. Plast Surg Int 2013;2013.

[39] ur Rashid H, Abbas K, Umer M. Single stage oncologic resection and reconstruction: a step toward development of sarcoma service in resource constrained country. Med Prast Surg 2013;3:134–41.

[40] Askari R, Umer M, e-Azam M, et al. Our experience with Van Nes rotationplasty for locally advanced lower extremity tumours. J Pak Med Assoc 2014;64:12.

[41] Haroon S, Uddin N, Hafeez K, et al. Chondromyxoid fibroma; experience of 36 cases of an intriguing entity. J Pak Med Assoc 2014;64(12 Suppl 2):S175–9.

[42] Umer M, Abbas K, Khan S, et al. Locking compression plate in musculoskeletal oncology “a Friend in Need”. Clin Orthop Surg 2013;5:321–6.

[43] Mohib Y, Umer HM. Operative management of patients with non-spinal metastatic bone disease. Does it actually improve quality of life? J Pak Med Assoc 2014;64:S116.

[44] Ahmed W, Mohib Y, Rashid RH, et al. Clinic based biopsy vs. theatre biopsy of bone and soft tissue extremity tumours: comparable diagnostic modalities. J Pak Med Assoc 2015;65:S207.

[45] Ahmed W, Kadir MM, Mohib Y, et al. Clinic-based biopsy with a small curette: is it a reliable and safe option in extremity tumours? J Pak Med Assoc 2015;65:S205.

[46] Tarig MI, Din NU, Ahmed A, et al. Challenges and pitfalls in diagnosis of Parosteal Osteosarcoma: a clinicopathologic study of 23 cases. J Solid Tumors 2016;6:17.

[47] Umer M, Ali M, Rashid RH, et al. Outcomes of internal hemipelvectomy for pelvic tumors: a developing country’s prospective. Int J Surg Oncol 2017;2:e07.