

SPECTRUM OF MUSCULOSKELETAL TUMOURS IN CHILDREN IN A TERTIARY HOSPITAL IN NIGERIAOde Michael B¹, Shitta A. H², Amupitan I³, Mancha D. G⁴, Onche I. I⁵**HOW TO CITE THIS ARTICLE:**

Ode Michael B, Shitta A. H, Amupitan I, Mancha D. G, Onche I. I. "Spectrum of Musculoskeletal Tumours in Children in a Tertiary Hospital in Nigeria". Journal of Evolution of Medical and Dental Sciences 2015; Vol. 4, Issue 48, June 15; Page: 8337-8343, DOI: 10.14260/jemds/2015/1210

ABSTRACT: BACKGROUND: Musculoskeletal tumours in children are a source of concern for the orthopaedic surgeon and paediatric oncologist. In Nigeria and other parts of sub Saharan Africa, where infectious diseases like malaria, tuberculosis and malnutrition are the leading causes of illness among children, little attention is given to paediatric oncology. This study was aimed at determining the spectrum of these tumours in children at our centre, and arriving at the common types of these tumours and the sites involved. **METHODOLOGY:** This was a retrospective study analysis of the pathology records from the cancer registry and the case notes of children who were diagnosed clinically and histologically, and managed for musculoskeletal tumours at the Jos University Teaching hospital Nigeria, from 2000 to 2009. **RESULTS:** A total of 90 musculoskeletal tumours in children were identified. The ages of the children ranged from 11 months to 18 years with a mean age of 10.9 years +/-5.1 years. The male female ratio was 1.3: 1.70(77.7%) tumours were malignant while, 20(22.2%) were benign. For the malignant variety, Rhabdomyosarcoma made up, 54(77.1%), Osteosarcoma 8(11.4%), Fibrosarcoma 5(7.1%). The benign tumour varieties were; Osteochondroma 5(25%), Giant cell tumour 2(10%), Angio fibroma 2(10%), Osteoid osteoma 2(10%). The anatomic regions affected were; head and neck 27(30.0%), trunk 20(22.2%), thigh 10(11.1%), legs 10(11.1%). **CONCLUSION:** Malignant tumours made up most of the musculoskeletal tumours in children. Rhabdomyosarcoma was the commonest malignant tumour and osteochondroma the commonest benign. The head and neck was the anatomical site mostly affected.

KEYWORDS: Musculoskeletal, Tumours, children, Nigeria.

INTRODUCTION: Tumours in children can be challenging for the clinician, and are an unpleasant reality in clinical practice.⁽¹⁻³⁾ Children can be plagued by a wide variety of tumours and malignant conditions ranging from the, hematological malignancies to solid tumours of the intra cranial, intra thoracic and intra-abdominal compartments to those involving the musculoskeletal system.⁽⁴⁻⁷⁾ The occurrence of these tumours in these children can be very devastating as with tumours even in adults. The inability of children to clearly characterize their symptoms.⁽⁸⁾ can delay the presentation of these conditions to the clinician and even when they present, can result in a misdiagnosis or a delayed diagnosis. Musculoskeletal tumours on the other hand when they appear may lead the parents to consider other clinical entities first because such conditions are usually associated with adults more frequently than children.

This can result in delayed presentation. An early diagnosis of these conditions whether benign or malignant and prompt treatment would lead to a better prognosis and outcome of care. Early presentation and improved management modalities have shown an increased survivalship in children with tumours both benign and malignant.^(5,9-12) In Nigeria, sub-Saharan Africa and other developing countries where the burden of disease in children is mainly from infective and nutrition

ORIGINAL ARTICLE

related illnesses such as malaria and tuberculosis⁽¹³⁻¹⁶⁾ attention to conditions such as tumours tends to take the backstage. The management of tumours also require a multidisciplinary and multimodal approach,⁽¹⁷⁻²⁰⁾ the use of specialized diagnostic equipment both laboratory and imaging and a range of treatment modalities,^(5,21-24) and this further adds to the cost of care, which further limits the quality of care obtainable in resource poor settings. These conditions have however been shown to be present among children in these environments.^(3,5,6,25) and thus, more attention needs to be paid to these conditions by means of enlightenment and greater allocation of the limited health resources in developing countries towards the management of these conditions. The poor attention to paediatric surgical oncology in this region, ignorance on the part of parents and poor access to healthcare, results in many of these patients presenting to the hospitals with locally advanced or metastatic tumours,^(3,5,25,26) with the attendant poor prognosis this stage of presentation portends. This study was aimed at elucidating the musculoskeletal tumours common in our hospital, the various histologic varieties as well as the age of occurrence and the sites more commonly involved.

METHODOLOGY: This was a retrospective study analysis of the pathology records from the cancer registry and the case notes of children who were diagnosed clinically and histologically, and managed for musculoskeletal tumours at the Jos University Teaching Hospital from 2000 to 2009. Data was obtained from these sources regarding the tumour type, sex, age at presentation and the anatomical site involved. Children with nonsolid tumours, tumours in the chest or within the abdomen and pelvis and those who had incomplete records were excluded from the study. The data was analysed for simple means and percentages using Epi - info statistical software.

RESULT: A total of 90 musculoskeletal tumours in children were identified. The ages of the children ranged from 11 months to 18 years with a mean age of 10.9 years +/-5.1 years. The male female ratio was 1.3: 1.70(77.7%) tumours were malignant while, 20(22.2%) were benign. Of the malignant variety, Rhabdomyosarcoma made up, 54(77.1%), Osteosarcoma 8(11.4%), Fibrosarcoma 5(7.1%). Table 1. The benign tumour varieties were; Osteochondroma 5(25%), Giant cell tumour 2(10%), Angio fibroma 2(10%), Osteoid osteoma 2(10%) Table 2. The anatomic regions affected were; head and neck 27(30.0%), trunk 20(22.2%), thigh 10(11.1%), legs 10(11.1%) Table 3, Fig. 1.

DISCUSSION: 90 musculoskeletal tumours in children were identified in this study. Malignant tumours made up 77.7% of these musculoskeletal tumours while the benign tumours made up 22.2%. The commonest histologic type of malignant tumour was Rhabdomyosarcoma which made up 54(77.1%) of the malignant tumours. This finding is similar to that obtained by other investigators.^(1,3) Rhabdomyosarcoma has been found to be amongst the commonest malignant musculoskeletal tumours affecting children.^(1,17,27,28) Israelsen et al in their work found Osteosarcoma and Ewings sarcoma the commonest varieties.⁽²⁹⁾ Osteosarcoma was the next most commonly occurring malignant tumour in children here 8(11.4%).

The above findings are in keeping with a review article by Arndt et al who found Osteosarcoma, Rhabdomyosarcoma and Ewings sarcoma to be the most commonly occurring malignant musculoskeletal tumours in children and adolescents.⁽¹⁷⁾ This is fact is also stated by Heare et al in their study.⁽¹²⁾ Osteosarcoma is a primary malignant bone tumour that has a peak incidence in adolescents and late adulthood.⁽³⁰⁾ It is commonly associated with bone growth during pubertal

ORIGINAL ARTICLE

growth spurts.^(30,31) It commonly affects adolescents and is one of the commonest musculoskeletal malignancies in adolescents.⁽³²⁾ Only one case of malignant fibrous histiocytoma and dermatofibrosarcoma were found in this study. Though Ewings sarcoma is among the common musculoskeletal tumours in children,^(12,17,33) no case of Ewings sarcoma was seen in this study. Other investigators in this part of the world have not reported Ewings sarcoma as a common cause of musculoskeletal malignancy.^(1,3,5,27) Of the benign tumours, Osteochondroma was the most commonly occurring (25%). Next commonly occurring benign tumours were Giant cell tumour (10%), Osteoid osteoma (10%). Similar findings are also noted in studies by other researchers.^(34,35)

The most frequently affected anatomical site in this study was the head and neck region (30%) and this site is also seen as being the most affected in works done by several other investigators.^(27,36,37) The trunk was next in line of involvement (22.2%) and the thighs and legs being involved in 11.1% each. The male female ratio in this study was 1.3: 1 which was also similar to that by Tanko et al and Adewuyi et al.^(1,3) The mean age of presentation was 10.9 yrs+/-5.1. Adewuyi and colleagues.⁽³⁾ had a mean age of 8.7years for rhabdomyosarcoma. From the above study the malignant tumours make up a larger percentage of musculoskeletal tumours in children. Ewings Sarcoma is found to be a rare occurrence in this environment. A prospective multicenter study covering a wide region of Nigeria would give a clearer picture of these tumours in children. This information can be employed in health care planning and population enlightenment to enhance early presentation and help improved care towards these conditions.

CONCLUSION: Malignant tumours made up most of the musculoskeletal tumours in children in our center. Rhabdomyosarcoma was the commonest malignant tumour and osteochondroma the commonest benign tumour. The head and neck was the anatomical site mostly affected.

REFERENCES:

1. Tanko NM, Echejoh GO, Manasseh NA, Mandong MB, Uba AF. Paediatric solid tumours in Nigerian children: a changing pattern? African journal of paediatric surgery: AJPS. 2009; 6 (1): 7-10.
2. Akinde OR, Abdulkareem FB, Daramola AO, Anunobi CC, Banjo AA. Morphological pattern of childhood solid tumours in Lagos University Teaching Hospital. Nigerian quarterly journal of hospital medicine. 2009; 19 (4): 169-74.
3. Adewuyi SA, Musa H, Samaila MO, Ogunrinde GO, Ameh EA, Popoola OB. Pattern of paediatric solid cancers seen in radiotherapy and oncology department, Ahmadu Bello University Teaching Hospital, Zaria - Nigeria. The Nigerian postgraduate medical journal. 2013; 20 (2): 120-4.
4. Stefan DC. Patterns of Distribution of Childhood Cancer in Africa. Journal of tropical pediatrics. 2015.
5. Hadley LG, Rouma BS, Saad-Eldin Y. Challenge of pediatric oncology in Africa. Seminars in pediatric surgery. 2012; 21 (2): 136-41.
6. Missaoui N, Khouzemi M, Landolsi H, Jaidene L, Abdelkrim SB, Abdelkader AB, et al. Childhood cancer frequency in the center of Tunisia. Asian Pacific journal of cancer prevention: APJCP. 2011; 12 (2): 537-42.

ORIGINAL ARTICLE

7. Tantawy AA, El Sherif NH, Ebeid FS, El-Desouky ED. Survival analysis after diagnosis with malignancy of Egyptian adolescent patients: a single-center experience. *Journal of pediatric hematology/oncology*. 2014; 36 (6): e346-52.
8. Junnila JL, Cartwright VW. Chronic musculoskeletal pain in children: part I. Initial evaluation. *American family physician*. 2006; 74 (1): 115-22.
9. Bhatia S, Landier W. Evaluating survivors of pediatric cancer. *Cancer journal*. 2005; 11 (4): 340-54.
10. Goldsby RE, Taggart DR, Ablin AR. Surviving childhood cancer: the impact on life. *Paediatric drugs*. 2006; 8 (2): 71-84.
11. Landier W, Bhatia S. Cancer survivorship: a pediatric perspective. *The oncologist*. 2008; 13 (11): 1181-92.
12. Heare T, Hensley MA, Dell'Orfano S. Bone tumors: osteosarcoma and Ewing's sarcoma. *Current opinion in pediatrics*. 2009; 21 (3): 365-72.
13. Bhutta ZA, Sommerfeld J, Lassi ZS, Salam RA, Das JK. Global burden, distribution, and interventions for infectious diseases of poverty. *Infectious diseases of poverty*. 2014; 3: 21.
14. Valadas E, Gomes A, Sutre A, Brilha S, Wete A, Hanscheid T, et al. Tuberculosis with malaria or HIV co-infection in a large hospital in Luanda, Angola. *Journal of infection in developing countries*. 2013; 7 (3): 269-72.
15. Mharakurwa S, Mutambu SL, Mberikunashe J, Thuma PE, Moss WJ, Mason PR, et al. Changes in the burden of malaria following scale up of malaria control interventions in Mutasa District, Zimbabwe. *Malaria journal*. 2013; 12: 223.
16. Murray CJ, Ortblad KF, Guinovart C, Lim SS, Wolock TM, Roberts DA, et al. Global, regional, and national incidence and mortality for HIV, tuberculosis, and malaria during 1990-2013: a systematic analysis for the Global Burden of Disease Study 2013. *Lancet*. 2014; 384 (9947): 1005-70.
17. Arndt CA, Rose PS, Folpe AL, Laack NN. Common musculoskeletal tumors of childhood and adolescence. *Mayo Clinic proceedings*. 2012; 87 (5): 475-87.
18. Weber KL, Peabody T, Frassica FJ, Mott MP, Parsons TW, 3rd. Tumors for the general orthopedist: how to save your patients and practice. *Instructional course lectures*. 2010; 59: 579-91.
19. Ogilvie CM, Crawford EA, Slotcavage RL, King JJ, Lackman RD, Hartner L, et al. Treatment of adult rhabdomyosarcoma. *American journal of clinical oncology*. 2010; 33 (2): 128-31.
20. Federman N, Bernthal N, Eilber FC, Tap WD. The multidisciplinary management of osteosarcoma. *Current treatment options in oncology*. 2009; 10 (1-2): 82-93.
21. Puhaindran ME, Pratt J, Manoso MW, Healey JH, Mintz DN, Athanasian EA. Predictive value of magnetic resonance imaging in determining presence of residual disease after marginal excision of unsuspected soft tissue sarcomas of the hand. *The Journal of hand surgery*. 2010; 35 (9): 1479-84.
22. Nystrom LM, Morcuende JA. Expanding endoprosthesis for pediatric musculoskeletal malignancy: current concepts and results. *The Iowa orthopaedic journal*. 2010; 30: 141-9.
23. Ilaslan H, Schils J, Joyce M, Marks K, Sundaram M. Radiofrequency ablation: another treatment option for local control of desmoid tumors. *Skeletal radiology*. 2010; 39 (2): 169-73.

ORIGINAL ARTICLE

24. Stein-Wexler R. MR imaging of soft tissue masses in children. *Magnetic resonance imaging clinics of North America*. 2009; 17 (3): 489-507, vi.
25. Ekenze SO, Ekwunife H, Eze BI, Ikefuna A, Amah CC, Emodi IJ. The burden of pediatric malignant solid tumors in a developing country. *Journal of tropical pediatrics*. 2010; 56 (2): 111-4.
26. Eyesan SU, Obalum DC, Nnodu OE, Abdulkareem FB, Ladejobi AO. Challenges in the diagnosis and management of musculoskeletal tumours in Nigeria. *International orthopaedics*. 2009; 33 (1): 211-3.
27. Akhiwu WO, Igbe AP, Aligbe JU, Eze GI, Akang EE. Malignant childhood solid tumours in Benin City, Nigeria. *West African journal of medicine*. 2009; 28 (4): 222-6.
28. Mandong BM, Ngbea JA. Childhood rhabdomyosarcoma: a review of 35 cases and literature. *Nigerian journal of medicine: journal of the National Association of Resident Doctors of Nigeria*. 2011; 20 (4): 466-9.
29. Israelsen RB, Illum BE, Crabtree S, Randall RL, Jones KB. Extremity sarcoma surgery in younger children: ten years of patients ten years and under. *The Iowa orthopaedic journal*. 2011; 31: 145-53.
30. Ottaviani G, Jaffe N. The epidemiology of osteosarcoma. *Cancer treatment and research*. 2009; 152: 3-13.
31. Burningham Z, Hashibe M, Spector L, Schiffman JD. The epidemiology of sarcoma. *Clinical sarcoma research*. 2012; 2 (1): 14.
32. Mirabello L, Troisi RJ, Savage SA. Osteosarcoma incidence and survival rates from 1973 to 2004: data from the Surveillance, Epidemiology, and End Results Program. *Cancer*. 2009; 115 (7): 1531-43.
33. Damron TA, Ward WG, Stewart A. Osteosarcoma, chondrosarcoma, and Ewing's sarcoma: National Cancer Data Base Report. *Clinical orthopaedics and related research*. 2007; 459: 40-7.
34. Abdulkareem FB, Eyesan SU, Akinde OR, Ezembakwe ME, Nnodu OE. Pathological study of bone tumours at the National Orthopaedic Hospital, Lagos, Nigeria. *West African journal of medicine*. 2007; 26 (4): 306-11.
35. Obalum DC, Eyesan SU, Ezembakwe ME, Abdulkareem FB. Pattern of osteochondromas in Lagos, Nigeria. *Nigerian quarterly journal of hospital medicine*. 2008; 18 (2): 69-71.
36. Missaoui N, Landolsi H, Jaidene L, Anjorin A, Abdelkader AB, Yaacoubi MT, et al. Pediatric rhabdomyosarcomas in Tunisia. *Asian Pacific journal of cancer prevention: APJCP*. 2010; 11 (5): 1325-7.
37. Mandong BM, Kidmas AT, Manasseh AN, Echejoh GO, Tanko MN, Madaki AJ. Epidemiology of soft tissue sarcomas in Jos, North Central Nigeria. *Nigerian journal of medicine: journal of the National Association of Resident Doctors of Nigeria*. 2007; 16 (3): 246-9.

ORIGINAL ARTICLE

Malignant tumours	Frequency	Percentage
Dermatofibro sarcoma	1	1.4%
Fibrosarcoma	5	7.1%
Liposarcoma	1	1.4%
Malignant firous histiocyoma	1	1.4%
Osteosarcoma	8	11.4%
Rhabdomyosarcoma	54	77.1%
Total	70	100.0%

Table 1: Malignant tumours

Benign tumours	Frequency	Percentage
Angiofibroma	2	10.0%
Chondroma	2	10.0%
Chondromyxoid fibroma	1	5.0%
Fibroma	1	5.0%
Fibrous dysplasia	1	5.0%
Fibrous histiocyoma	3	15.0%
Giant cell tumours	2	10.0%
Osteiod osteoma	2	10.0%
Osteoblastoma	1	5.0%
Osteochondroma	5	25.0%
Total	20	100.0%

Table 2: Benign tumours

Location	Frequency	Percentage
Arm	2	2.2%
Axilla	5	5.6%
Foot	5	5.6%
Forearm	2	2.2%
Groin	5	5.6%
Hand	4	4.4%
Head and neck	27	30.0%
Leg	10	11.1%
Thigh	10	11.1%
Trunk	20	22.2%
Total	90	100.0%

Table 3: Anatomic location of the Tumours

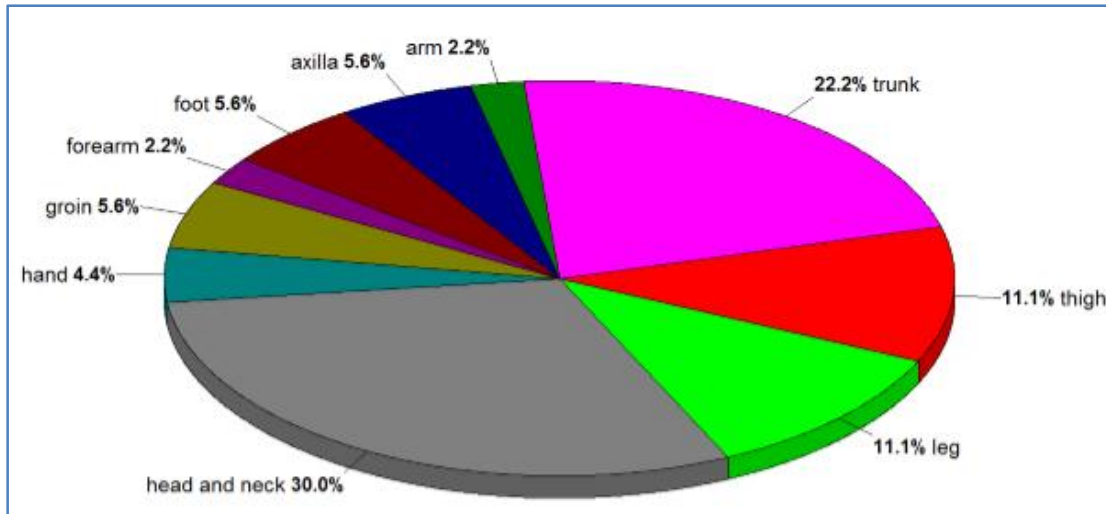


Fig. 1: Anatomic location of Tumours

AUTHORS:

1. Ode Michael B.
2. Shitta A. H.
3. Amupitan I.
4. Mancha D. G.
5. Onche I. I.

PARTICULARS OF CONTRIBUTORS:

1. Lecturer/Consultant Orthopaedic Surgeon, Department of Orthopaedics, JOS University Teaching Hospital, JOS, Plateau State, Nigeria.
2. Lecturer/ Consultant Paediatric Surgeon, Department of Surgery, JOS University Teaching Hospital, JOS, Plateau State, Nigeria.
3. Lecturer/Consultant Orthopaedic Surgeon, Department of Orthopaedics, JOS University Teaching Hospital, JOS, Plateau State, Nigeria.

FINANCIAL OR OTHER

COMPETING INTERESTS: None

4. Lecturer/Consultant Orthopaedic Surgeon, Department of Orthopaedics, JOS University Teaching Hospital, JOS, Plateau State, Nigeria.
5. Professor/ Consultant Orthopaedic Surgeon, Department of Orthopaedics, JOS University Teaching Hospital, JOS, Plateau State, Nigeria.

NAME ADDRESS EMAIL ID OF THE CORRESPONDING AUTHOR:

Dr. Ode Michael B,
Department of Orthopaedics,
JOS University Teaching Hospital,
PMB 2076, JOS, Plateau State,
Nigeria-930001.
E-mail: odemb2014@gmail.com

Date of Submission: 18/05/2015.
Date of Peer Review: 19/05/2015.
Date of Acceptance: 08/06/2015.
Date of Publishing: 13/06/2015.