

Original Article

## Evaluation of Histopathological Subtypes of Ameloblastoma in a Tertiary Institution in Southern Part of Nigeria

Cornelius Azekanabo Iyogun, Gabriel Ehimen Ukegheson, Olufemi Gbenga Omitola

### Abstract

**Background:** Ameloblastoma is the most frequent odontogenic tumour reported in Nigeria. It is a slow growing locally invasive tumor which though rarely undergoes metastasis, but is clinically persistent. **Aim:** The aim of this study was to evaluate the prevalence of the various histopathological subtypes of ameloblastoma seen at the Oral Pathology department of the University of Port Harcourt Teaching Hospital and compare it with other studies in Nigeria and elsewhere. **Materials and Methods:** This is a five year (2008-2013) retrospective study of all histopathologically diagnosed ameloblastoma at the Oral Pathology department of the University of Port Harcourt Teaching Hospital, Port Harcourt. Lesions were categorized using the 2005 World Health Organization classification of odontogenic tumours. **Results:** Within the study period, there were 52 cases of ameloblastoma out of a total of 63 odontogenic tumors seen. The patients' age ranged from 7 – 65 years with a mean age of  $28.5 \pm 14.6$  years. There was equal sex predilection. Solid/multicystic ameloblastoma was the most prevalent 30(57.7%) and mandible was the predominant site with 43(82.7%) cases. Solid/multicystic ameloblastoma were more in females while unicystic ameloblastoma were more in males. Solid/multicystic and unicystic ameloblastoma were predominantly seen in the mandible. There was no case of malignant ameloblastoma or ameloblastic carcinoma. **Conclusion:** Solid/multicystic ameloblastoma was the most prevalent with higher incidence in females. The mandible was the commonest site of occurrence.

**Keywords:** Ameloblastoma; Histological; Prevalence; Unicystic; Multicystic; Odontogenic.

*Cornelius Azekanabo Iyogun, Gabriel Ehimen Ukegheson, Olufemi Gbenga Omitola. Evaluation of Histopathological Subtypes of Ameloblastoma in a Tertiary Institution in Southern Part of Nigeria. International Journal of Oral & Maxillofacial Pathology; 2014;5(4):07-11. ©International Journal of Oral and Maxillofacial Pathology. Published by Publishing Division, Celesta Software Private Limited. All Rights Reserved.*

### Introduction

Ameloblastoma is a slow growing locally invasive tumor which rarely metastasize, but persists clinically.<sup>1-5</sup> It is the most common odontogenic tumour reported in Nigeria,<sup>6-8</sup> South Africa<sup>9</sup> and found to be more common among blacks than whites<sup>10,11</sup>. It is said to be mostly occurring in the 3<sup>rd</sup>-4<sup>th</sup> decades of life although recent reports have found it to be commoner in children and adolescent<sup>6,12</sup> and affects the mandible more than maxilla.<sup>6,8</sup> Maxillary ameloblastoma is indistinguishable from their mandibular counterparts histologically, but may behave more aggressively. This is presumably because of the thin and fragile bone of the maxilla which unlike the thick cortical plates of the mandible, allows a relatively unimpeded spread of the tumour to the surrounding structures, such as the maxillary sinus, nasal cavity, orbit and occasionally the cranial base.<sup>2</sup> This account for the high recurrence rate of this tumour in the maxilla with significant mortality and often management challenges of the tumour in this location.<sup>13</sup>

The 2005 World Health Organization (W.H.O) histological classification of ameloblastomas (benign type) includes four types: solid/multicystic, unicystic, extraosseous/peripheral and desmoplastic.<sup>14</sup> The solid/multicystic is further divided into follicular and plexiform or mixed types. The follicular type can further be subdivided at cellular level into four subgroups; spindle cell type, acanthomatous type, granular cell, and basal cell ameloblastoma. The Plexiform types are anastomosing strands of basal cells often without a conspicuous stellate reticulum type cells in a delicate stroma often showing cystic degeneration.

The unicystic ameloblastoma appears as a cyst on gross examination but may appear multilocular on radiography. It has two main histologic pattern; luminal and mural but sometimes a third group named intraluminal when it appears there are detachment of ameloblastomatous tissue inside the lumen. The extra osseous type appears histologically as the solid/multicystic type. The desmoplastic type has a loose stroma immediately surrounding the odontogenic

epithelium but further away a very dense stroma which may appear on histology with little evidence of odontogenic epithelium.<sup>15</sup> Previous Nigerian studies, on histological subtype of ameloblastoma are few.<sup>15-18</sup> These studies were mainly in the southwest geopolitical zone of Nigeria with none from the south-south zone of Nigeria particularly Port Harcourt the largest city in the zone. This study was therefore designed to evaluate the prevalence of the various histological subtypes of ameloblastoma seen at the Oral Pathology department of the University of Port Harcourt Teaching Hospital (UPTH) in Port Harcourt, south-south region of Nigeria and compare it with other studies.

### Materials and Methods

This was a 5-year (2008-2013) retrospective study of all histologically diagnosed cases of ameloblastoma reported at the Oral Pathology department of UPTH, Port Harcourt. Ethical approval for this study was obtained from the Ethical Committee of the hospital. Formalin fixed tissues and paraffin embedded blocks of patients were processed and prepared for staining with haematoxylin & eosin. All slides were subsequently reviewed by two Oral Pathologists (CA Iyogun and OG Omitola) to reconfirm the diagnosis. The diagnoses were in accordance with the 2005 W.H.O classification of ameloblastoma. Biodata comprising of age, sex, and site were obtained from the patients records in the department. Data obtained were coded and analysis was performed using SPSS version 17.0 (Chicago, Illinois, USA). Chi square test was used to assess the statistical significance between different proportions. Significant level was fixed at >0.05.

### Results

In the five year study period, there were 52 cases of ameloblastoma out of a total of 63 odontogenic tumors seen. The patients age range from 7–65 years with a mean age of 28.5±14.6 years. Solid/multicystic ameloblastoma was the most prevalent histological subtype accounting for 30(57.7%) cases follow by unicystic ameloblastoma with 20(38.4%) cases. Peripheral and desmoplastic ameloblastoma had 1(1.9%) case each (Table 1).

The mandible was the predominant site for ameloblastoma with 43(82.7%) cases while the maxilla had 6(11.5%). The right and left posterior sides of the mandible were

predominantly affected with 19(36.5%) and 15(28.8%) cases respectively. There was one peripheral and one palatal cases accounting for 1.9% each. One case had no data with respect to site on the jaw (Table 2).

Histological subtypes		Frequency
Solid/Multicystic		30 (57.7%)
Unicystic	Mural	17 (32.7%)
	Luminal	2 (3.8%)
	Intramural	1 (1.9%)
Peripheral		1 (1.9%)
Desmoplastic		1 (1.9%)
Total		52 (100.0%)

Table 1: Frequencies of histopathological subtypes of Ameloblastoma

Side of Jaw	Frequency
Right	19 (36.5%)
Left	15 (28.8%)
Anterior mandible	5 (9.6%)
Right and left mandible	6 (11.5%)
Total	45 (86.5%)
Not recorded	7 (13.5%)
	52 (100.0%)
Site on jaw	Frequency
Mandible	43 (82.7%)
Maxilla	6 (11.5%)
Peripheral	1 (1.9%)
Palate	1 (1.9%)
Total	51 (98.1%)
Not recorded	1 (1.9%)
	52 (100%)

Table 2: Frequencies of side and site of jaw involved

The peak incidence of ameloblastoma was from the second to third decades of life. Also, the peak incidence of solid multicystic ameloblastoma and mural type of unicystic ameloblastoma was the second to third decades. This was not statistically significant (P-value 0.494) (Table 3).

There was equal sex predilection of 26 cases each but among the solid/multicystic type of ameloblastoma, there were 19(36.6%) females and 11(21.1%) males giving a female to male ratio of 1.7:1. There were 13 males (25.1%) and 7(13.4%) among unicystic ameloblastoma giving a male to female ratio of 1.85: 1. This value was not statistically significant (P-value 0.120) (Table 4).

As regard the relationship of the histological subtypes and site, solid/multicystic was

predominantly in the mandible with 24(47.2%) cases while the maxilla had only 4(7.8%) cases. Unicystic subtypes were also predominantly in the mandible with

19(37.3%) cases. This observation was statistically significant with P-value of 0.0000 (Table 5).

Histological subtypes	Age group						
	0-9	10-19	20-29	30-39	40-49	50-59	60-69
Solid/Multicystic	1(1.9%)	7(13.4%)	10(19.1)	5(9.5%)	3(5.8%)	4(7.6%)	0(0.0%)
Unicystic	Mural	0(0.0%)	8(15.3%)	4(7.7%)	2(3.8%)	1(1.9%)	0(0.0%)
	Luminal	0(0.0%)	1(1.9%)	0(0.0%)	0(0.0%)	1(1.9%)	0(0.0%)
	Intramural	0(0.0%)	1(1.9%)	0(0.0%)	0(0.0%)	0(0.0%)	0(0.0%)
Peripheral	0(0.0%)	0(0.0%)	0(0.0%)	0(0.0%)	1(1.9%)	0(0.0%)	0(0.0%)
Desmoplastic	0(0.0%)	0(0.0%)	1(1.9%)	0(0.0%)	0(0.0%)	0(0.0%)	0(0.0%)
Total	1(1.9%)	17(30.6%)	15(28.7%)	7(13.3%)	6(11.5%)	4(7.6%)	2(3.8%)

Table 3: Relationship of Histopathological subtypes to age group (P-value 0.494)

Histological subtypes	Gender of patients		
	Male	Female	Total
Solid/Multicystic	11(21.1%)	19(36.6%)	30(57.7%)
Unicystic	Mural	12(23.2%)	5(9.6%)
	Luminal	0(0.0%)	2(3.8%)
	Intramural	1(1.9%)	0(0.0%)
Peripheral	1(1.9%)	0(0.0%)	1(1.9%)
Desmoplastic	1(1.9%)	0(0.0%)	1(1.9%)
Total	26(50.0%)	26(50.0%)	52(100.0%)

Table 4: Relationship of Histopathological subtype to gender (P-value 0.120)

Histological subtypes	Site on jaw			
	Mandible	Maxilla	Palate	Peripheral
Solid/Multicystic	24(47.2%)	4(7.8)	0(0.0%)	1(2.0%)
Unicystic	Mural	16(31.4%)	1(2.0)	0(0.0%)
	Luminal	2(3.9%)	0(0.0%)	0(0.0%)
	Intramural	1(2.0%)	0(0.0%)	0(0.0%)
Peripheral	0(0.0%)	0(0.0%)	1(2.0%)	0(0.0%)
Desmoplastic	0(0.0%)	1(2.0%)	0(0.0%)	0(0.0%)
Total	43(84.3%)	6(11.8%)	1(2.0%)	1(2.0%)

Table 5: Relationship of histopathological subtypes to site on the jaws (P-value 0.000)

		Frequency
Solid / Multicystic	Follicular	14 (26.9%)
	Plexiform	9 (17.3%)
	Granular	1 (1.9%)
	Basal Cell type	3 (5.8%)
	Acanthomatous	2 (3.8%)
	Hemangiomas	1 (1.9%)
	Unicystic	20 (38.4%)

Table 6: Frequencies of histopathological subtypes of Ameloblastoma

Follicular ameloblastoma was the most common cytologic type with 14 cases followed by the plexiform type with nine cases. The basal cell type had three cases while the acanthomatous type had two cases and there was only one case of granular cell type in this study (Table 6).

There was no case of malignant ameloblastoma or ameloblastic carcinoma in this study.

### Discussion

Ameloblastoma is the commonest odontogenic tumor in Nigeria.<sup>4</sup> It is also the commonest in North Africa,<sup>19</sup> and in people of Asian descent<sup>20</sup> and India.<sup>21</sup> In this study, 52 cases of ameloblastoma were seen accounting for 82.5% of odontogenic tumors found in the period under review. This is in agreement with various reports from sub-Saharan Africa<sup>7,18</sup> that ranges from 70 to above 80%. However, this is in contradiction with reports from Europe and America where odontoma is the most prevalent odontogenic tumour. This may be due to underreporting of odontoma (which is generally an asymptomatic lesion) in developing world.<sup>22-</sup>

<sup>25</sup> From this study, the youngest patient was seven years while the oldest patient was 65 years with the mean age of  $28.5 \pm 14.6$  years. This is in agreement with the reports from Brazil<sup>12</sup> (31 years), Nigeria<sup>26</sup> (31.7+15.6) years and Jamaica<sup>27</sup> (29.1 years). Our finding of peak incidence within the 2<sup>nd</sup> and 3<sup>rd</sup> decade of life is in agreement with that of Santos et al,<sup>12</sup> Adebisi et al,<sup>18</sup> but differs with that of Basse et al,<sup>7</sup> Waldron et al<sup>28</sup> who reported peak incidence within the 4<sup>th</sup> and 5<sup>th</sup> decade of life.

In this study, male/ female ratio was equal, which was comparable to studies in the western part of Nigeria<sup>18,26</sup> who reported 1:1 and 1.1:1 respectively. However, a ratio of 1.6:1 was reported in Kaduna.<sup>29</sup> The unicystic ameloblastoma affected 13(25.1%) male and 7(13.4%) females giving a male to female ratio of 1.85:1. This was in agreement with a similar study in Brazil.<sup>12</sup> Solid/multicystic ameloblastoma was the most prevalent histological subtype accounting for 57.7%. This was far higher than 13.9% reported in an Indian study.<sup>21</sup> In this study, the most prevalent subtype was unicystic ameloblastoma with a prevalence of 35.5%.

Mandible was the predominant site of ameloblastoma with 43(82.7%) of cases while the maxilla had 6(11.5%). This was in agreement with previous study by Gardner.<sup>30</sup> However, Arotiba et al<sup>5</sup> in Ibadan and Adebisi et al<sup>18</sup> in Ile-Ife both in southern Nigeria reported higher prevalence of 91 and 93% respectively. Follicular ameloblastoma was the most common cellular type followed by the plexiform type and distantly by basal cell type. This is in agreement with the finding of Ladeinde et al<sup>16</sup> and Adebisi et al.<sup>18</sup> It is important to note that there were no cases of malignant ameloblastoma or ameloblastic carcinoma. This is similar to the findings in Kano Nigeria.<sup>8</sup>

### Conclusion

Ameloblastoma was found to have equal sex predilection in this study but the solid/multicystic ameloblastoma was the most prevalent in this region and had a higher incidence in females than their male counterpart with mandible being the commonest site. Follicular ameloblastoma was the commonest cellular subtype in this study.

### Acknowledgement

We would like to acknowledge all the staff members of Oral Pathology Department for their support and guidance.

### Author Affiliations

1.Dr. Iyogun CA, Department of Oral Pathology and Biology, Faculty of Dentistry, 2.Dr. Ukegheson GE, Department of Child Dental Health, 3.Dr. Omitola OG, Department of Oral Pathology and Biology, Faculty of Dentistry, University of Port Harcourt/ University of Port Harcourt Teaching Hospital, Port Harcourt, Rivers State, Nigeria.

### Reference

1. Neville BW, Damm DD, Allen CM, Bouquot JE. Textbook of Oral and Maxillofacial Pathology. SauUis; 3<sup>rd</sup> Ed, St Louis; WM Saunders Co, p. 2009.
2. Natri AL, Wiesenfeld D, Radden BG, Everson J, Scully C. Maxillary ameloblastoma: a retrospective study of 13 cases. Br J Oral Maxillofac Surg 1995;33:28-32.
3. Ogunsalu C, E Scipio E, Williams N. Review of six cases of maxillary ameloblastoma from the west indies re-entry cryosurgery as prophylactic surgical intervention. West Indian Med J 2009;58(4):309-403.
4. Odukoya O. Odontogenic tumors: analysis of 289 Nigerian cases. J Oral Pathol Med 1995;24:454-7.
5. Arotiba JT, Ogunbiyi JO, Obiechina AE. Odontogenic tumors: a 15-year review from Ibadan, Nigeria. Br J Oral Maxillofac Surg 1997;35:363-7.
6. Lawal AO, Adisa AO, Popoola BO. Odontogenic tumours in children and adolescents: a review of forty-eight cases. Ann Ib Postgrad Med 2013;11(1):7-11.
7. Basse GO, Osunde OD, Anyanechi CE. Maxillofacial tumors and tumor-like lesions in a Nigerian teaching hospital: an eleven year retrospective analysis. Afr Health Sci 2014;14(1):56-63.
8. Ochicha O, Iyogun CA, Omitola OG, Raphael S and Adebola RA. Odontogenic tumours in Kano; Northan Nigeria. Afr J Oral Health 2014;4(1):9-13.
9. Shear M, Singh S. Age standardized incidence rates of ameloblastoma and dentigerous cyst in Witswatersrand, South Africa. Community Dent Oral Epidemiol 1978;6:195-9.
10. Sawyer DR, Mosadumi A, Page DG, Svirsky JA, Kekere-Ekun AT. Racial predilection of ameloblastoma? A probable answer from Lagos (Nigeria)

- and Richmond, Virginia (USA). *J Oral Med* 1985;40:27-31.
11. Owens BM, Schuman NJ, Mincer HH, Turner JE, Oliver FM. Dental odontomas : a retrospective study of 104 cases. *J Clin Pediatr Dent* 1997;21:261-4.
  12. Pereira FAC, Melo LA, Gurgel CAS, Cangussu MCT, Azevedo RA, Santos JN. Clinicopathological and demographic characteristics of ameloblastoma in a population from Bahia, Brazil. *Rev Odonto Cienc* 2010;25(3):250-5.
  13. Bredenkamp JK, Zimmerman MC, Mickel RA. Maxillary ameloblastoma- A potential lethal neoplasm. *Arch Otolaryngol Head Neck Surg* 1989;115:99-103.
  14. Barnes L, Everson JW, Reichart P, Sidransky D. Pathology and genetics of head and neck tumors. Lyon IARC Press: 2005; p. 284-327.
  15. Hertog D, Bloemena E, Aartman IHA, van-der-Waal I. Histopathology of ameloblastoma of the jaws; some critical observations based on a 40 years single institution experience. *Med Oral Pathol Oral Cir Bucal* 2012;17(1):e76-82.
  16. Ladeinde AL, Ajayi OF, Ogunlewe MO, Adeyemo WL, Arotiba GT, et al. Odontogenic tumors: A review of 319 cases in a Nigerian teaching hospital. *Oral Surg Oral Med Oral Pathol Oral Radiol Endod* 2005;99:191-5.
  17. Arotiba GT, Ladeinde AL, Arotiba JT, Ajike SO, Ugboko VI, Ajayi OF. Ameloblastoma in Nigerian children and adolescents: a review of 79 cases. *J Oral Maxillofac Surg.* 2005;63(6):747-51.
  18. Adebisi KE, Ugboko VI, Omoniyi-Esan GO, Ndukwe KC, Oginni FO. Clinicopathological analysis of histological variants of ameloblastoma in a suburban Nigerian population. *Head & Face Medicine* 2006;2:42-50.
  19. Tawfik MA, Zyada MM: Odontogenic tumors in Dakahlia, Egypt: Analysis of 82 cases. *Oral Surg Oral Med Oral Pathol Oral Radiol Endod* 2010;109:e67-e73.
  20. Reichart PA, Philipsen HP, Sonner S: Ameloblastoma: Biological profile of 3677 cases. *Eur J Cancer B Oral Oncol* 1995;31B:86-99.
  21. Gill S, Chawda J, Jani D. Odontogenic tumors in Western India (Gujarat): Analysis of 209 cases. *J Clin Exp Dent.* 2011;3(2):e78-83.
  22. Adebisi KE, Odukoya O, Taiwo EO. Ectodermal odontogenic tumors: analysis of 197 Nigerian cases. *Int J Oral Maxillofac Surg* 2004;33(8):766-70.
  23. Fernandes AM, Duarte EC, Pimenta FJ, Souza LN, Santos VR, Mesquita RA, de Aguilá MC. Odontogenic tumor: a study of 340 cases in a Brazilian population. *J Oral Pathol Med* 2005;34(10):583-7.
  24. Guerrisi M, Piloni MJ, Keszler A. Odontogenic tumors in children and adolescents. A 15-year retrospective study in Argentina. *Med Oral Patol Oral Cir Bucal* 2007;12:e180-5.
  25. Sriram G, Shetty RP. Odontogenic tumors: a study of 250 cases in an Indian teaching hospital. *Oral Surg Oral Med Oral Pathol Oral Radiol Endod* 2008;105:e14-e21.
  26. Ladeinde AL, Ogunlewe MO, Bamgbose BO, Adeyemo WL, Ajayi OF, Arotiba GT, Akinwande JA. Ameloblastoma: Analysis of 207 cases in a Nigeria teaching hospital. *Quintessence Int* 2006;37:69-74.
  27. Ogunsalu C, West W, Lewis A, Williams N. Ameloblastoma in Jamaica predominantly unicystic: analysis of 47 patients over a 16-year period and a case report on re-entry cryosurgery as a new modality of treatment for the prevention of recurrence. *West Indian Med J* 2011;60(2):240-6.
  28. Waldron CA, El-Mofty SK. A Histopathologic study of 116 Ameloblastoma with special reference to the Desmoplastic variant. *Oral Surg Oral Med Oral Pathol* 1987;63:441-451.
  29. Olaitan AA, Adeola DS, Adekeye EO. Ameloblastoma: Clinical features and management of 315 cases in Kaduna, Nigeria. *J Craniomaxillofac* 1993;21:351-5.
  30. Gardner DG. A pathologist's approach to the treatment of ameloblastoma. *J Oral Maxillofac Surg* 1984;42:161-6.

#### Corresponding Author

Dr. Iyogun CA,  
 Department of Oral Pathology and Biology,  
 Faculty of Dentistry,  
 University of Port Harcourt Teaching  
 Hospital,  
 Port Harcourt, Rivers State, Nigeria.  
 Email: [iyoguncornelius@yahoo.com](mailto:iyoguncornelius@yahoo.com)