THE CENTRALAFRICAN JOURNAL OF MEDICINE

Vol. 50, Nos. 7/8 **CONTENTS** July-August 2004 **ORIGINAL ARTICLES** Vaginal hysterectomy. A five year prospective descriptive study J Shava, N L Nene, L Mpande61 Pattern of cleft lip and plate in Benin City. Nigeria..... ON Obuekwe, O Akapata 65 Traumatic asphyxia during stadium stampede ... REVIEW ARTICLE I.T. Gangaidzo Forgotten diseases: Relapsing fever NOTES AND NEWS Instructions to Authors Central African Journal of Medicine 75

Pattern of cleft lip and plate in Benin City, Nigeria

ON OBUEKWE, O AKAPATA

Abstract

Objective: To study the pattern of cleft lip and/or palate (CL/P) in Benin City, Nigeria and to compare the findings with reports from other parts of the world. The results of this study may be used to improve the welfare of affected patients.

Design: A descriptive study.

Setting: The University of Benin Teaching Hospital (UBTH), Benin City, Edo State, Nigeria.

Subjects: 103 ethnic Nigerian patients with cleft lip and/or palate.

Main Outcome Measures: Consecutive ethnic Nigerian patients presenting with CL/P were studied. Patients' data collected included age, sex, ethnic group, type of CL/P and the laterality pattern. Descriptive statistics were generated for all the variables recorded.

Results: There were 49.5% females and 50.5% males; 95.1% were children and 4.9% were adults. The ethnic groups most often affected were Urhobo 29.1%, Bini 27.1% and Ibo 17.4%. The combined cleft lip and palate 60.2% was the commonest type of cleft. There were 28.2% patients with cleft lip only. More (7.8%) females were affected by the isolated cleft palate deformity. Twenty five patients had exceeded the time of repair. Their reasons were ignorance (36%) and financial constraints (64%)

Conclusion: The data from this study is consistent with studies from other parts of the world. However, a high proportion of patients did not have access to early treatment due to ignorance and financial constraints. Public enlightenment and financial assistance for the indigent patient is recommended.

Cent Afr J Med 2004;50(7/8):65-9

Introduction

The cleft lip and/or palate (CL/P) is the most common congenital craniofacial malformation. ¹⁻³ The aetiology of CL/P is complex and heterogeneous as causes linked to

environment, genetics and gene-environment interaction are known. Although the CL/P deformity has a worldwide frequency of one in 700, the incidence of CL/P varies from region to region and among races. Most studies have shown that the incidence is highest in Orientals and

Correspondence to:
Dr O N Obuekwe
Department of Oral and Maxillofacial Surgery
University of Benin Teaching Hospital
Benin City
Edo State, Nigeria
E mail: oxobuekwe@yahoo.com

Mongoloids, intermediate in Caucasians and lowest in blacks. ¹²⁻¹⁴ Concerning the incidence of the deformity, there is a higher rate of combined cleft lip and palate (CLP) compared to the cleft palate (CP). ¹⁵⁻¹⁷ Studies of laterality patterns in infants with external birth defects found that the left side was significantly more involved for CL/P. ¹⁷⁻¹⁹ Other congenital anomalies are associated with CL/P. ²⁰⁻²² Certain parental factors like age and occupation have been implicated in the incidence of CL/P. ^{23,24}

Gender differences exist in the occurrence of CL/P, with a higher overall incidence in males compared to females. ^{19,25-27} However, most studies have shown a preponderance of females for CP,^{25,27} while males are more often affected by the combined cleft lip and palate (CLP). ^{19,26,27} Also black males are more often involved in bilateral cases. ^{14,28}

The aim of this study is to document the pattern of CL/P as seen in Benin City, Edo State, Nigeria and to compare the results with those from other parts of the world. The results of this study may be used to improve the welfare of affected patients.

Materials and Methods

This study was carried out at the Department of Oral and Maxillofacial Surgery, University of Benin Teaching Hospital, (UBTH) Edo State, Nigeria. Over a four year period, consecutive ethnic Nigerian patients presenting with CL/P were studied. Patients' data collected included age, sex, ethnic group, and type of CL/P. The laterality pattern was also recorded. The places of delivery of the patients were recorded as well as the referring personnel. Patients presenting late for treatment and the reasons for delay were documented. Parents' data documented were age, occupation and the parent(s) (or person) accompanying the patient. Descriptive statistics were generated for all the variables recorded.

Results

During the study period, 103 patients with CL/P were recorded. There were 51 (49.5%) females and 52 (50.5%) males giving a female to male ratio of about 1:1 (Table I). There were 98 (95.1%) children and five (4.9%) adults. The children were aged one day to nine years (mean age eight months) with most 66 (64.0%) in the zero to six month age group (Table I).

Table I: Age and sex distribution of patients with CL/P.

Age range		Sex _			
(in months)	Male	Female	Number	Percent	
0-6	34	32	66	64.0	
7 – 12	8	9	17	16.5	
13 – 18		3	3	2.9	
19 – 24	1	1	2	2.0	
> 24	9	6	15	14.6	
Total	52	51	103	100	

The adults were aged 18 years to 34 years with a mean age of 23.3 years. The ethnic groups most often affected were Urhobo 30 (29.1%), Bini 28 (27.1%) and Ibo 18 (17.4%). These are shown in Table II.

Table II: Distribution of patients by ethnic groups.

Ethnic group	Number	Percent		
Urhobo	. 30	29.1		
Bini	28	27.1 17.4		
lbo	18			
Esan	7	6.8		
Etsako	7	6.8		
lka	4 -	3.9		
ljaw	3	2.9		
Hausa	1	1.0		
Yoruba	1	1.0		
lbibio	1	1.0		
Ighara	1	1.0		
Ora	1	1,0		
Akoko Edo	1	1.0		
Total	103	100		

Most 77 (74.8%) patients were delivered in hospitals while 20 (19.4%) were delivered at home. Six (5.8%) were delivered in maternity homes. The general medical practitioner referred 39 (37.8%) patients while the paediatrician referred 20 (19.4%) patients. Other sources of referral are shown in Table III.

Table III: Sources of referrals of patients.

Source	Number	Percent
Obstetrician	11	10.7
Midwife	9	8.7
Paediatrician	20	19.4
Surgeon	7	6.8
General practitioner	39	37.8
Relations	8	7.9
Others	9	8.7
Total	103	100

The combined cleft lip and palate 62 (60.2%) was the most common type of cleft, of which 17 (16.5%) cases involved the left side in males, seven (6.8%) in females and 26 (25.2%) were bilateral. There were 29 (28.2%) patients with cleft lip only. Of these 16 (15.5%) involved the left side, nine (8.7%) the right side and four (3.9%) were bilateral. More (eight or 7.8%) females were affected by the isolated cleft palate deformity. These are shown in Table IV. Most 102 (99.0%) cases of CL/P were nonsyndromic while one (1.0%) case was syndromic (van der Woude). Four (3.9%) patients had associated congenital anomalies (cardiac 1.0%, inguinal hernia 1.0%, syndactyl 1.0%, varus deformity 1.0%). Twenty five (24.3%) patients had exceeded the expected time for repair. Their reasons were ignorance, nine (36%) cases and financial constraints, 16 (64%).

Туре	No.	•	Laterality			Sex	
		%	Left (%)	Right (%)	Bilateral (%)	Male (%)	Female (%)
Cleft lip	29	28.2	16 (15.5)	9 (8.7)	4 (3.9%)	15 (14.6)	14 (13.6)
Cleft lip and palate	62	60.2	24 (23.3)	12 (11.7)	26 (25.2)	33 (32.0)	29 (28.1)
Cleft palate	12	11.6	-	_ '	_ ` `	4 (3.9)	8 (7.8)
Total	103	100	40 (38.8)	21 (20.4)	30 (29.1)	52 (50.5)	51 (49.5)

Ninety five mothers (92.2%), 89 fathers (86.4%), four church mission workers (3.9%) and a husband (1.0%) accompanied the patients. Four (3.9%) patients (adults) reported alone. The mothers ranged in age from 19 to 43 years with a mean age of 32.0 years while the fathers were aged from 26 to 55 years with a mean age of 38.4 years. Regarding parental occupation, most mothers were housewives (40 or 38.8%), traders (28 or 27.1%) and fashion designers (five or 4.6%). The fathers were mainly professional drivers (28 or 27.1%), traders (18 or 17.5%) and civil servants (16 or 15.5%). A diverse group of jobs made up the rest of the parental occupations. Most 89 (86.4%) patients were of the low socio-economic group while 10 (13.6%) were of the middle class.

Discussion

Edo State (population 2 172 608)²⁹ is located in the southern part of Nigeria. The UBTH is the largest health facility within a 250 kilometre radius and has over four states as its catchment area. The dominant ethnic groups of Edo State in descending order are Bini, Esan and Etsako. The Ibo followed by the Urhobo tribe are the dominant ethnic group in the adjacent Delta state (population 2 590 491).²⁹ The Ibo are not only indigenous to the catchment area but Ibos from other states are widely distributed in Nigeria due to their extensive engagement in trading. The pattern of CL/P shows a large variation among different countries. 10 A sub-division has to be made for the different ethnic groups within a base population as the incidence can differ among races. 10,11 This is consistent with the objectives of this study which documented CL/P on different ethnic groups but exclusively among Nigerian blacks. The ethnic groups often involved are the Urhobo (29.1%) of the Niger Delta area of Nigeria. The Bini of Edo State ranked second (27.1%) on the list of ethnic groups affected and when the other ethnic groups within the state are added, indigenes of Edo State ranked highest as the single state most commonly affected by CL/P. This may be due to the location of the study centre (Benin City) and for reasons of proximity, most patients report here. A study has shown that proximity to referral centres can influence response of patients sent to a specialist.30

Most of the patients in this study were children (95.1%). The unexpected birth of a baby with a cleft deformity is a traumatic and shocking experience, generating anxiety for parents.³¹ Most parents are likely to seek help early for their

children with CL/P. A near equal sex distribution was observed in this study. Most reports on the epidemiology of CL/P show that overall the deformity is commoner in males. 19-22 However, some studies have shown near equal gender distribution 17,19 or a preponderance of females. 32 Also, the anatomic location of the cleft differs among the sexes. The isolated cleft palate is said to be commoner among females. 21 This is consistent with the findings from this study.

The incidence of CL/P is influenced by the sources of data (e.g. hospital record, birth certificates) and these could lead to under reporting unless the registration system is good. ¹⁰ Also, questionnaires sent to different doctors can produce reliable data. ¹⁰ This study relied on patients referred from other sources (Table III) and may not reflect the true figures. A study has shown that fram patients referred to a specialist did not report to him. ³³

This study found the left, sided (23.3%)-cleft lip and palate (60.2%) as the commonest cleft deformity and more often, males (32.0%) were affected. This is consistent with most studies. ^{17,19-22} Differences in the lateral distribution of specific birth defects may be due to subtle differences in morphogenesis on the left and right sides of the embryo brought about by establishment of left-right asymmetry prior to organogenesis. ¹⁸ Some studies have found the cleft lip as the commonest cleft deformity. ^{31,24} Most (99.0%) of the CL/P seen in this study were the non-syndromic type. This is consistent with other studies. ^{16,34,35} Other studies have shown that congenital anomalies are associated with CL/P to varying degrees. ^{17,35}

In our centre, cleft lip is repaired at three to four months and cleft palate at 12 to 18 months. Sandberg et al. have stated that with advances in neonatology and paediatric anaesthesia, cleft surgery should be performed during the neonatal period. Even with our arbitrary upper limit of six months for cleft lip repair and two years for cleft palate repair, 25 (24.3%) patients were deemed to have presented late, citing ignorance and financial constraints as reasons. Late repair of CL/P may have a negative impact on parentinfant bonding and infant growth and development.¹ Although the analysis of parental occupation did not reveal any occupational exposures that may lead to clefting in the offspring,²⁸ it did reveal that most parents (89 or 86.4%) were of the low socio-economic group. This group of parents will benefit from enlightenment programmes and financial assistance concerning their children with CL/P.

In conclusion, this study has shown that the ethnic roups most often involved in orofacial clefting were the Jrhobos, Binis and Ibos. It also showed that the lemographic and laterality pattern is similar to reports from other parts of the world. However, a high proportion of parents did not have access to early treatment due to ignorance and financial constraints. Public enlightenment is necessary to highlight that the treatment of this deformity is available. The government, charity and non-governmental organizations are also enjoined to assist these patients, especially the indigent ones. Since this study documented the pattern of CL/P on patients referred to our unit, under reporting is possible. A multicentre study involving many hospitals is, therefore, recommended.

References

- 1. Sandberg DJ, Magee WP, Denk MJ. Neonatal cleft lip and palate repair. AORN 2002;75:490-8.
- 2. Williams Y. Working with parents to promote health. *J Child Health Care* 1998;2:182-6.
- 3. Hibbert SA, Field JK. Molecular basis of familial cleft lip and palate. *Oral Dis* 1996;2:238-41.
- 4. Bianchi F, Calzolari E, Ciulli L, Cordier S, Gualandi F, Pierini A, et al. Environment and genetics in the aetiology of cleft lip and cleft palate with reference to the role of folic acid. Epidemiol Prev 2000;24:21-7.
- Lorente C, Cordier S, Goujard J, Ayme S, Bianchi F, Calzolari E, et al. Tobacco and alcohol use during pregnancy and risk of oral clefts. Occupational Exposure and Congenital Malformation Working Group. Am J Public Health 2000;90:415-9.
- 6. Carinci F, Pezzetti F, Scapoli L, Martinelli M, Carinci P, Tognon M. Genetics of non-syndromic cleft lip and palate: a review of international studies and data regarding the Italian Population. Cleft Palate Craniofac J 2000;37:33-40.
- 7. Vieira AR, Orioli IM, Castilla EE, Cooper ME, Marazita ML, Murray JC. MSX1 and TGFB3 contribute to clefting in South America. *J Dent Res* 2003;82:289-92.
- 8. Chung KC, Kowalski CP, Kim HM, Buchman SR. Maternal cigarette smoking during pregnancy and the risk of having a child with cleft lip/palate. *Plast Reconstr Surg* 2000;105:485-91.
- 9. Murray JC. Gene/environment causes of cleft lip and/or palate. *Clin Genet* 2002;61:228-56.
- 10. Derijcke A, Eerens A, Carel C. The incidence of oral clefts: a review. Br J Oral maxillofac Surg 1996;34:488-94.
- 11. Croen LA, Shaw GM, Wasserman CR, Tolarova MM Racial and ethnic variations in the prevalence of orofacial clefts in California, 1983-1992. Am J Med Genent 1998;79:42-7.
- 12. Menegotto BG, Salzano FM. Epidemiology of oral clefts in a large South American sample. *Cleft Palate Craniofac J* 1991;28:373-7.

- 13. Leck I. The geographical distribution of neural tube defects and oral clefts. *Br Med Bull* 1984;40:390-5.
- Tolarova M. A study of the incidence, sex ratio, laterality and clinical severity in 3 660 probands with facial clefts in Czechosloakia. Acta Chir Palst 1987;29:77-87.
- Natsume N, Kawai T, Kohama G, Teshima T, Kochi S, Ohashi Yet al. Incidence of cleft lip or palate in 303 738 Japanese babies born between 1994 and 1995. Br J Oral Maxillofac Surg 2000;38:605-7.
- 16. Liang J, Wang Y, Miao L, Zhu J, Zhou G, Wu Y. Nonsyndromic cleft lip with or without palate in Chinese population: analysis of 3 766 cases. Hua Xi Yi Ke Da Xue Xue Bao 2000;31:408-10.
- 18. Paulozzi LJ, Larry JM. Laterality patterns in infants with external birth defects. *Teratology* 1999;60:265-71
- 19. Orkar KS, Ugwu BT, Momoh JT. Cleft lip and palate. The Jos experience. *East Afr Med J* 2002;79:510-3.
- 20. Rajabian MH, Sherkat M. An epidemiological study of oral clefts in Iran: an analysis of 1 669 cases. *Cleft Palate Craniofac J* 2000; 37:191-6.
- 21. Mcheik JN, Lacombe D, Bondonny J M, Vergnes P. Cleft lip and palate. Epidemiologic analysis: report of 60 cases. *Ann Chir Plast Esthet* 2000;45:425-9.
- 22. Wantia N, Rettinger G. The current understanding of cleft lip malformations. Facial Plast Surg 2002;18:147-53.
- 23. Vieira AR, Orioli IM, Murray JC. Maternal age and oral clefts: a reappraisal. *Oral Surg Oral Med Oral Pathol Oral Radiol Endod* 2002;94:530-5.
- 24. Schnitzer P.G, Olshan AF, Erickson JD. Paternal occupation and risk of birth defects in offspring. *Epidemiology* 1995;6:577-83.
- Coupland MA, Coupland AI. Seasonality, incidence and sex distribution of cleft lip and palate births in Trent Region ('73 - '82). Cleft Palate J 1988;25:33-7.
- 26. al-Bustan SA, el-Zawahri MM, al-Adsani AM, Bang RL, Ghunaim I, Maher BS, et al. Epidemiologic and genetic study of 121 cases of oral clefts in Kuwait. Orthod Craniofac Res 2002;5:154-60.
- Nazer J, Hubner ME, Catalan J, Cifuentes L. Incidence of cleft lip and palate in the University of Chile Maternity Hospital and in Maternity Chilean participating in the Latin American collaborative study of cogenital malformations (ECLAMC). Rev Med Chil 2001;129:285-93.
- 28. Stoll C, Alembik Y, Dott B, Roth MP. Epidemiology and genetic study in 207 cases of oral clefts in Alsace, North Eastern France. *J Med Genet* 1991;28:325-9.
- 29. National population commission.1991 population census of the federal Republic of Nigeria: analytical report at the national level. April 1998;25-29.

- Dobbs F. Referrals to Irish accident and emergency departments. Ir Med J 1995;88:54-5. Young, O' Riordan M, Goldstein JA, Robin NH.
- palate or both want to know? Cleft Palate Craniofac J 2001;38:55-8. Olasoji O, Arotiba T, Dogo D. Experience of unoperated cleft lip and palate patients in a Nigerian

What information parents of newborns with cleft lip,

teaching hospital. Trop Doct 2002; 32:33-6. Bourguet C, Gilchrist V, McCord G. The consultation and referral process. A report from NEON. North

Eastern Ohio Network Research Group. J Fam Pract

1998;46:47-53.

- 34. Steinwachs EF, Amos C, Johnston D, Mulliken J, Stal S, Hecht JT. Nonsyndromic cleft lip and palate is not associated with cancer or other birth defects. Am J Med Genet 2000;90:17-24.

 - Natsume N, Niimi T, Furukawa H, Kawai T, Ogi N, Suzuki Y et al. Survey of congenital anomalies associated with cleft lip and/or palate in 701 181 Japanese people. Oral Surg Oral Med Oral Pathol Oral Radiol Endod 2001;91:157-61.



This work is licensed under a Creative Commons
Attribution – NonCommercial - NoDerivs 3.0 License.

To view a copy of the license please see: http://creativecommons.org/licenses/by-nc-nd/3.0/

