Sella Turcica Bridging as a Predicator of Dentofacial Anomalies: A Cephalometric Analysis

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ABSTRACT

Introduction: Sellar changes are associated with several dentofacial anomalies. Clinicians should be aware of different morphological varaiants of sella turcica (ST).

Objective: To find the prevalence of sella turcica bridging and to analyze the absence or presence of bridging with a spectrum of dentofacial anomalies.

Materials & Method: 710 case records were selected from the database; out of which 473 subjects met the inclusion criteria. 280 lateral cephalogram revealed a normal shape of ST. Among them, 71 subjects were selected by interval sampling which were taken as the control group. Among initial 473 subjects, 78 samples showed ST bridging and were taken as the study group. Dental casts and radiographs (panoramic and cephalometric) were evaluated to find any dentofacial anomalies. Subjects were divided on the basis of dentofacial anomalies such as abnormal tooth position, size, shape, number, malocclusion, sagittal skeletal relationship. Chi-square test and binomial logistic regression analysis were done to find the association and correlation among the variables.

Result: In the study group, 32 subjects showed a complete bridging and 46 subjects a partial bridging of the sella turcica. There was no significant association between ST bridging and age, gender or ethnic groups. However, there was significant association (p=0.001) between the presence of bridging and dentofacial anomalies when compared with the control group.

Conclusion: The occurrence of ST bridging is 16.49% with no association to age, gender and ethnic groups in a Nepalese sample. Sella turcica bridging can be used as a diagnostic tool for early prediction of dentofacial anomalies.

Key-words: bridging, dentofacial anomalies, sella turcica

INTRODUCTION

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Sella turcica (ST) is a saddle-shaped structure located on the intracranial surface of the sphenoid bone in the middle cranial fossa.¹ ST can be divided anatomically into anterior wall (tuberculum sellae), floor, and posterior wall (dorsum sellae). Axelsson *et al*² classified morphological variants of ST into: oblique anterior wall, double contour of the floor, irregularity (notching) in the posterior part of the dorsum sellae, pyramidal shape of the dorsum sella and ST bridging. Kucia *et al*³ added three other variants: hypertrophic posterior clinoid process, hypotrophic posterior clinoid process, and oblique contour of the floor.

ST is an important and commonly used landmark for cephalometric tracings. The relationship of pituitary gland pathology and endocrine disease is the result of their common embryological origin, which are the cranial neural crestal cells of early chondrocranium.^{1,4,5} Same cells are also associated with the genesis of both ST and derivatives of the frontal process/first branchial arch, along with the teeth.^{6,7} Thus the variation in ST shape can be used as an early predictor of abnormalities related to the craniofacial complex.⁸

Bridging is an extensively studied morphological variant of ST. In healthy subjects, its reported incidence ranges from 3.8-13%.^{1,2,9} Numerous authors have mentioned bridging as an important radiological feature in syndromes.¹⁰⁻¹² A study revealed incidence of bridging to be 18.6% in patients with severe craniofacial deviation; whereas, another reported an incidence of 16.7% and 7.3% in patients treated with combined surgical-orthodontic therapy and only orthodontic therapy respectively.¹³ A significant association of ST bridging with dental anomalies such as

palatally displaced canines, mandibular second premolar aplasia and dental transposition has been established in previous studies.^{14,15}

Hence, the present study aims to determine the occurrence of ST bridging in pre-orthodontic Nepalese patient samples and to compare the spectrum of different dentofacial anomalies in patients with and without bridging.

MATERIALS AND METHOD

The retrospective observational study was conducted after receiving ethical clearance from Institutional Review Committee. Case records of 710 Nepalese subjects were collected from the database of Department of Orthodontics, Dhulikhel Hospital. Cephalometric and panoramic radiographs of good quality were included. The exclusion criteria were poor visibility of ST on cephalogram, craniofacial anomaly or syndrome, and history of trauma. Dental casts and extra-oral radiographs were analyzed to detect dentofacial anomalies such as abnormal tooth position, size, shape, number, malocclusion and sagittal dental relationship (Angle Class I, II, III) following International Classification of Diseases to Dentistry & Stomatology given by World Health Organisation.¹⁶

Among the initially screened case records, 473 subjects (66.61%) met the inclusion criteria. The selected radiographs were then segregated according to age, gender and ethnic groups (Newar, Brahmin, Chhetri, Mongoloid, and Madhesi). The selected cephalograms were classified into six different morphological variants of ST based on the classification by Axelsson *et al*² (Figure 1). The classification of ST on cephalograms was done independently by an oral radiologist and an orthodontist. The initial grouping of 473 lateral cephalograms resulted in 280 (59.19%) STs with a normal shape. Out of these, 71 (25.35%) subjects (19 males and 52 females) were selected by interval sampling and taken as the control group. Among the initial 473 sample of subjects, 115 subjects (24.31%) showed other morphological variants of ST.

Radiographs were classified according to the sagittal skeletal relationship using Steiner's analysis. Skeletal bases were grouped according to ANB angle value: Class I skeletal base: angle 2-4°, Class II: >4° and Class III: <0°. Wits appraisal was determined when limitations of ANB angle occurred. The extent of bridging was determined by measuring the length and diameter of ST on cephalometric radiographs.² The measurements were done manually by one observer (oral radiologist) in dark room. The contour of pituitary fossa was traced from the tip of the dorsum sellae to the tuberculum sellae on a transparent acetate sheet using 0.5mm lead pencil. Length of the sellae was

measured as the distance between tuberculum sella and tip of the dorsum sellae. Diameter of ST was measured from the tuberculum sellae to the most posterior point on the inner wall of the pituitary fossa. Length and diameter of the sella were compared to score the extent of bridging, resulting into the following groups:¹⁴

Class I: No bridging (length of sella greater than or equal to three-fourths of the diameter)

Class II: Partial bridging (length of sella lesser than three-fourths of the diameter)

Class III: Complete bridging (Figure 2).

Intra-observer bias was eliminated by retracing randomly selected twenty cephalograms on two different occasions within two weeks under similar conditions. Data were analyzed using SPSS Version 21. Study and control groups were segregated according to different age, gender, ethnic groups and dentofacial anomalies. Statistical comparisons included chi-square test to find the association of dentofacial anomalies in study and control groups of different age, gender and ethnicity. The statistical significance was set at p-value < 0.05. Binominal logistic regression analysis was used to correlate sagittal skeletal relationships in both groups.

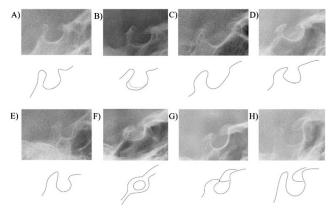
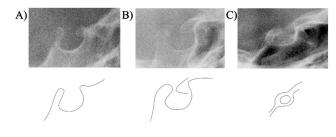
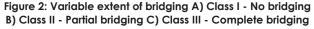


Figure 1: Morphological variants of sella turcica seen in lateral cephalogram A) normal B) double contour of the floor C) oblique anterior wall D) irregularity (notching) of dorsum sella E) pyramidal shape of dorsum sella F,G) complete bridging H) partial bridging





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RESULT

The test group consisted of 78 (16.49%) subjects with 30 males and 48 females exhibiting radiological bridging; among them 32 subjects (41.02%) showed complete and 46 subjects (58.97%) showed partial bridging. Data regarding the distribution of bridging for different age, gender and ethnic groups are shown in Table 1. The chi-square test was not significant, suggesting no association between bridging and age (p=0.305), gender (p=0.089), ethnicity (p=0.559). In the test group, 139 anomalies were seen; whereas in the control group, 69 anomalies were seen. In both control and test groups, most common

anomaly were related to tooth position, followed by malocclusion (Table 2). The study showed significant association (p=0.001) between dental anomalies with test and control groups, suggesting that the occurrence of dental anomalies increase with bridging of sella turcica (Table 3).

Distribution of no bridging, partial bridging and complete bridging among different skeletal malocclusions is shown in Table 4. Binominal logistic regression analysis showed the presence of bridging in skeletal Class II at almost six times the rate (CI: 2.16-17.797; p=0.001) compared to skeletal Class III (Table 5)..

Table 1: Occurrence of ST bridging in different age, e	ethnicity, gender and their association
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	Age		Ethnicity			Gender			
Years	Test Group (N=78)	Control Group (N=71)	Туре	Test Group (N=78)	Control Group (N=71)	Туре	Test Group (N=78)	Control Group (N=71)	
≤10	1(1.3%)	0(0%)	Newar	35(44.9%)	23(32.4%)	Male	30(38.5%)	19(26.8%)	
11-20	39(50%)	29(40.8%)	Brahmin	21(26.9%)	24(33.8%)				
21-30	35(44.9%)	41 (57.7%)	Chhetri	15(19.2%)	14(19.7%)	Female	48(61.5%)	52(73.2%)	
21.40	2/2.0071	1/1 407)	Mongoloid	6(7.7%)	9(12.7%)				
31-40 3(3.8%)	1 (1.4%) Madhesi	Madhesi	1(1.3%)	1(1.4%)					
p-Value	0.3	05 (NS)	0.559 (NS)				0.089 (NS)		

NS: Not significant

Table 2: Dentofacial anomalies with ST bridging (test group) and without bridging (control group)

	Anomalies	Test Group	Control Group	
	Dilaceration	23	08	
Sharpa	Malformed root	02	01	
Shape	Dens invaginatus	00	01	
	Total	25(17.98%)	10(14.49%)	
Size	Microdontia	08	01	
Size	Total	8(5.75%)	1(1.45%)	
	Retained deciduous	03	01	
Number	Supernumerary	04	03	
Number	Congenitally missing	17	12	
	Total	24(17.26%)	16(23.18%)	
	Impacted teeth	33	21	
Position	Buccally/ palatally erupted	10	03	
	Total	43(30.93%)	24(34.78%)	
Malocclusion	Rotated tooth	33	16	
	Diastema/ spacing	06	02	
	Total	39(28.05%)	18(26.08%)	
	Total	139	69	

Table 3: Association of dental anomalies with test and control groups

Presence or absence of	Test	group	Control group		
anomalies	Complete bridging	Partial bridging	(No Bridging)	p-Value	
Anomalies	15(46.9)	13(28.3)	35(49.2%)		
More than one anomalies	15(46.9%)	32(69.6%)	16(22.5%)	0.001*	
Absent	2(6.25%)	1 (2.2%)	20(28.2%)		
Total	32(100%)	46(100%) 71(100%)			

*Significant at p<0.05

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Eulersk of building	Type of skeletal malocclusion						
Extent of bridging	Class I	Class II	Class III	Total			
No bridging	31(43.66%)	25(35.21%)	15(21.13%)	71			
Partial bridging	4(8.7%)	38(82.61%)	4(8.7%)	46			
Complete bridging	6(18.75%)	24(75%)	2(6.25%)	32			
Total	41	87	21	149			

Table 4: Distribution of	of ST bridging among (different skeletal malocclusions
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Delation	D Wald	-16		95% CI			
Relation	В	Wald	df Odds ratio	Odds railo	Lower	Upper	p-Value
Class I	-	27.512	2	-	-	-	0.000*
Class II	-0.215	0.127	1	0.806	0.247	2.638	0.722
Class III	1.825	11.501	1	6.200	2.160	17.797	0.001*

*Significant at p<0.05

DISCUSSION

Development of craniofacial complex is a complex process with cranial neural crestal cells coordinating a complex series of morphogenetic steps.¹⁷ These cells migrate dorsolaterally and along with the ectoderm form two important facial primordial processes, namely frontonasal process and first branchial arch; which in turn form the proximal maxilla and mandible.¹⁸ After migration, they undergo sequential and reciprocal interactions, differentiating into dental epithelial progenitor cells.67 Derivatives of them frame the shape of anterior wall of the ST and anterior portion of the pituitary gland (adenophysis). Kjaer¹⁹ concluded that, malformations in the anterior wall of ST are associated with the abnormalities in frontonasal, maxillary and palatine fields. Thus, any alteration in the morphology of ST may be associated with pituitary gland disorders and dentofacial anomalies.^{3,20}

Numerous studies have suggested bridging as the most common variant of ST morphology.^{1,3,4,13,14} Bridging can be appreciated at the age of six, although it can also develop later.^{10,15} In the literature, bridging has been mentioned as both normal and pathological variant of ST.¹⁴ Many studies have determined bridging in relation to dentofacial anomalies, but those studies were limited to anomalies like palatally displaced canine, mandibular second premolar aplasia, dental transposition or a particular type of malocclusion.^{4,13-15,21} Hence, we conducted this study to evaluate possible dentofacial anomalies given by International Classification of Diseases to Dentistry and Stomatology¹⁶ seen in the subjects with ST bridging, and to correlate them to subjects having normal STs. The present study revealed 59.2% subjects presented with normal ST shape, which is similar to the findings in the available literature.^{5,12} The presence of bridging (complete/partial) was found in 16.5% with no correlation to age and gender. These findings are also similar to the previous studies.^{2,13,14,22} Our study reported no significant association between ethnicity and the bridging. This study also reported the occurrence of dentofacial anomalies increased with bridging, compared to the subjects with normal ST shape. This finding supports previous studies correlating dentofacial anomalies with ST bridging.^{1,4,13,14} Additionally, skeletal Class II was more common in ST bridging. However, this finding was not supported by previous studies, where skeletal Class III was more common in subjects with bridging.^{12,23} This difference can be explained by the fact that, skeletal Class II is more common than skeletal Class III in Nepalese population.^{24,25}

CONCLUSION

Findings of the present study support crucial relationship between cranial and dentofacial growth and development of the pituitary gland including sella turcica. Consequently, lateral cephalometric radiographs should be observed carefully; any alteration of sella turcica morphology especially bridging can be used as a diagnostic tool to predict as well as initiating preventive and early treatment of dentofacial anomalies.

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