

White Paper on the Efficacy of the Osseo-Restore™ Appliance to Effect Skeletal Patency and Growth in The Anterior Maxilla

<sup>a</sup> Moulton D. Abernathy, Jr, DDS, DFBI

<sup>b</sup> Timothy G. Bromage,\* BA, MA, PhD

<sup>c</sup> Stephen Deal, DDS, BS

<sup>d</sup> Steve Galella, DDS, IBO

<sup>e</sup> W.C. Simmons, DDS, BS

<sup>a</sup> Private General Dentistry Practice, Parsons, Tennessee; Facial Beauty Institute Diagnostics, Collierville, Tennessee

<sup>b</sup> Department of Molecular Pathobiology, New York University College of Dentistry, New York; Professor

<sup>c</sup> Director, Stephen Deal DDS, Little Rock, Arkansas; Director, Facial Beauty Institute, Collierville, Tennessee

<sup>d</sup> Manager, Steve A. Galella, DDS, PLLC, Director, Facial Beauty Institute, Collierville, Tennessee

<sup>e</sup> Director, Cris Simmons DDS, Seattle, Washington; Director, Facial Beauty Institute, Collierville, Tennessee

\* Corresponding author:

Timothy G. Bromage  
Department of Molecular Pathobiology  
New York University College of Dentistry  
345 East 24th Street  
New York, NY 10010  
USA

Tel: +1 212 998 9597 (office)

+1 212 289 4534 (home)

Fax: +1 212 995 4445

Email: [tim.bromage@nyu.edu](mailto:tim.bromage@nyu.edu)

## **Abstract**

**Introduction.** Bone growth remodeling by bone deposition and resorption has a deep scientific research history, which in the present day requires to be put to purpose for understanding remodeling phenomena occurring over the subnasal region during orthodontic treatment;

**Methods.** A conebeam CT superimposition technique rendering an assessment of bone growth remodeling with an appreciation for its principles, combined with an understanding of the role of mechanical forces aids our understanding of craniofacial growth and development;

**Results.** Mechanical forces elicited by an orthopedic appliance in the biological range will stimulate normal growth remodeling in the growing child and compensatory remodeling in the adult non-grower;

**Conclusions.** Anterior alveolar remodeling may be specifically targeted to treat maxillary deficiencies provided that treatment forces do not exceed the biological range.

## **Highlights**

Bone growth remodeling principles are central to an understanding of the developing human face.

During growth teeth drift by classic deposition-resorption remodeling mechanisms.

Changes in bone mass occur by seeking strain thresholds that remain patent through life.

Orthopedic appliances form bone over anterior maxillary roots when forces remain in the biological range.

Properly designed and managed orthopedic appliances can be safely applied to reverse maxillary deficiencies.

Recent claims on social media, e.g. <sup>1</sup>, draw attention to orthodontic treatments using appliances that putatively eliminate bone in the subnasal region of the maxilla and expose roots of the anterior dentition. Claims such as this without scientific research obfuscate and contribute nothing but unbridled fear. Whilst acknowledging that proper research on the topic takes some time, we wish to respond by saying, 1) we have begun this research in earnest for the sake of documenting variation in morphogenetic responses to such treatments, and 2) to address immediately in this white paper the issue. To achieve the latter goal, we shall briefly review the history of bone growth remodeling research and then its application to the craniofacial complex. We will then dwell on the role of bone growth remodeling for promoting integration of the craniofacial complex and, in this context, recognize the role that remodeling plays in the repositioning of teeth during growth in the grower and non-grower. Bone strain as a goal to be achieved by bone remodeling processes is invoked as an axiom of hard tissue biology, which is violated when orthodontic treatment forces exceed the capacity of the bone to adapt.

While first experiments in bone growth took place in the 18th Century, the modern synthesis of its concepts began with Brullé and Hugeny <sup>2</sup>, which then became the focus of research by eminent figures in the morphological, histological, and experimental sciences for about 100 years; e.g., <sup>3-25</sup> among others.

Bone growth remodeling is the fundamental mechanism bearing on skeletal morphogenesis, which involves coordinated surface patterns of bone deposition and resorption.\* Prompted by displacement of bones by craniofacial orocapsular matrices, the coupled processes of bone deposition and resorption during growth achieve the changes in size and shape that are necessary to enclose their respective matrices and support their muscle attachments <sup>26-28</sup>.

Enlow <sup>29</sup> benefited from these historical precedents in bone growth remodeling research and began publishing numerous papers and books alone and with colleagues that documented the principles of bone growth remodeling that would be used to chart the normal pattern of human facial growth and development. *These principles are central to an understanding of the anteriorly-facing surfaces of the developing human face, in particular that of the subnasal region, which is the topic of this communication and much speculation in the orthodontic community.*

Enlow <sup>29,30</sup> emphasized the *direction* in which bone surfaces were oriented to explain, for instance, that the outer flared metaphyseal cortex of a growing long bone was necessarily resorptive because this surface faced obliquely *away* from the growth direction. The metaphyseal endosteal aspect was explained to be depository because this surface faced obliquely *toward* the growth direction. Enlow <sup>26,31,32</sup> popularized these concepts in relation to the growth of the craniofacial skeleton, which emphasized the concept of “cortical” or “osseous” drift (bone growth movement through tissue space) and the “V principle” (gross morphological bone restructuring, shaping and enlargement) as the morphogenetic responses to displacement-based compensatory bone growth remodeling. Enlow <sup>26,31,32</sup> also proposed the employ of "Part - Counterpart" analysis from 2D lateral radiographs, the superimposition method of which permits

the identification of deposition-resorption growth fields and thus a description of not only what happens, but how it happens in the anteriorposterior and superoinferior directions. The method has the benefit of demonstrating what compensatory remodeling phenomena are occurring over the subnasal region in both the growing child and the adult non-grower.

Today we have an enhanced understanding of bone formation processes from Boyde and Jones<sup>33</sup> and a wealth of knowledge on the molecular and cellular physiology of developing bone; e.g.,<sup>34-36</sup>. This improved understanding underpins what we now know is the high level of anatomical integration among the parts of the growing oral region<sup>37</sup> that are consistent with Part - Counterpart analysis.

This integration is not something static, but rather dynamic throughout life. In growing individuals the remodeling of the periosteal subnasal alveolar region is resorptive, growing rearward in compensation for displacement of the maxilla anteriorly. The alveolar periosteal bone is not eliminated because bone formation occurs on the contralateral surfaces of the root sockets and behind the teeth on the palatal alveolar cortex; *teeth are not passively displaced with the jaws as they grow, but rather they drift by classic deposition-resorption remodeling mechanisms within their sockets*. In the adult non-grower the same is true, and though this is not strictly occurring as an ontogenetic growth program, it is primary growth elicited as a bone's adaptive response to mechanical loading.

It is axiomatic that an alteration in the mechanical forces on *any* bone of the body will cause that bone to respond. Bone construction and reconstruction is regulated by sensing of the ambient strain environment<sup>38</sup>, so much so that strain is said to be the 'goal' of bone size and shape change<sup>39</sup>. A bone that fails to experience its strain threshold will reduce its mass, and a bone that experiences peak strains above its threshold will gain mass<sup>40</sup>. Both reductions and gains in mass do so by seeking the appropriate strain threshold for that bone, a process that remains patent throughout life.

If mechanical forces originating from an orthopedic appliance are used to generate anteriorward tooth movement, the only mechanisms available for such movement are those of the classic deposition-resorption remodeling mechanisms discussed here. Mechanical forces in the biological range will cause cortical drift of the tooth roots through bone tissue space to accommodate the desired position directed by the force, and further, the adjacent alveolar bone may also increase in mass in order to balance the perceived strains and to bring the bone back to its appropriate threshold. Posterior teeth encroaching anteriorly because of applied forces will cause growth that includes bone formation on the periosteal subnasal surface. The contralateral surfaces within the anterior aspects of the root sockets will be resorptive, and thus by cortical drift the roots will shift or rotate anteriorly; rotation occurs because the anchoring bands are typically positioned on the first permanent molars at mid-crown height, and forces driven forward at the level of height of contour may obliquely reorient the anterior crowns whilst leaving the root tips near to their original position. 10 3, 3 14

Figure 1-2 are conebeam CT scans that illustrate these phenomena on a growing 9 yrs and 4 mo old child treated with a Osseo-Restore™ removable appliance for 10 mo, and an individual

whose growth was complete and that received an Osseo-Restore™ fixed appliance for 4 mo and then Controlled Arch™ System orthodontics for 12 mo. The superimposition technique follows that of Enlow<sup>31</sup>, which is fundamentally based upon growth boundaries, with all landmarks being implemented in the 3D rendering software, Anatomage (Santa Clara, CA). This method allows before-and-after growth/treatment differences to be visualized, measured, and related specifically to bone deposition-resorption remodeling activities. The child in Figure 1 had been diagnosed as exceeding growth in the vertical dimension relative to that of their horizontal development. As judged by the protrusive nature of the midface in the older blue profile in relation to the earlier profile in warm colors, treatment with the removable appliance primarily stimulated growth of the maxilla anteriorly. Normal bone resorption over the nasoalveolar clivus for this age accompanied a small downward relocation of the subnasal region, but otherwise forming bone predominated, which is not the typical remodeling pattern for a growing child<sup>26,31,32</sup>. In Figure 2, of the individual whose growth had ceased, orthodontic forces uprighted several teeth and caused an inferior and outward relocation of the maxillary alveolus by ca. 1-3 mm. A modest amount of resorption on outer alveolar and deposition over palatal surfaces of the alveolar bone will have made this relocation possible. A small inferior relocation of the tooth-bearing portion of the mandible also occurred.

Should mechanical treatment forces exceed the biological range, the rate of bone formation will not outpace resorption. Accelerated bone repair can reach 1.7  $\mu\text{m}/\text{day}$ <sup>41</sup>, but bone resorption is capable of removing bone by up to 25  $\mu\text{m}/\text{day}$ <sup>42</sup>. Given such a discrepancy, the periosteal alveolar cortex will be eliminated and the subnasal roots will be exposed within, for instance, only one-three months if the biological range of forces is chronically exceeded. Nevertheless, caution must be exercised when suggesting that tooth roots have been exposed from conebeam CT scans. Labial alveolar cortical bone in places can range from ca. 100-600  $\mu\text{m}$  in thickness<sup>43</sup>, and x-ray voxel edge effects (i.e., voxels not fully containing bone) will fail to visualize and reconstruct the bones' mineralization completely, particularly as the bone thickness diminishes to the set voxel size of the scanner. In addition, newly formed bone mineralization density, while increasing rapidly in the first weeks of formation, takes months-to-years to be nearly complete<sup>44-46</sup> and thus this bone will be underrepresented and poorly visualized in conebeam CT scans.

## Conclusion

The clinical implications contained within craniofacial growth and development studies may be immediately appreciated in the application of specialized dental appliances, e.g., the Osseo-Restore™ appliance, which stimulates signals for modification of anterior alveolar bone within biologically safe boundaries as described above. Properly designed and managed orthopedic appliances such as this capable of stimulating appropriate signal systems, which we are only now beginning to understand, can be predictably and safely applied to potentially reverse epigenetic maxillary deficiencies.

\*Because of disparities of terminology in the literature, the term remodeling is qualified here to describe 'growth remodeling', often referred to as modeling, versus 'secondary remodeling',

which often refers to Haversian replacement of bone or the repair/maintenance of bone removed because of fracture or requirements of mineral homeostasis.

### Author Contributions

1  
2  
3  
4  
5

### Acknowledgments

### References

1. Meehan R. Dental Malpractice: Anterior Growth Guidance Appliance (AGGA: FAGGA) and ALF Hearst CT News Blogs. Norwalk, CT: Hearst Communications Inc.; 2020: p. Blog.
2. Brullé M, Huguény M. Expériences sur le développement des os dans les mammifères et les oiseaux, faites au moyen de l'alimentation par la garance. *Annales des sciences naturelles Zoologie* 1845;4:283-357 (et planche 216-217).
3. Angle EH. Bone-Growing. *Dental Cosmos* 1910;52:261-267.
4. Baer MJ. Patterns of growth of the skull as revealed by vital staining. *Human Biology* 1954;26:80-126.
5. Barnicot NA. The supravital staining of osteoclasts with neutral-red: Their distribution on the parietal bone of normal growing mice, and a comparison with the mutants grey-lethal and hydrocephalus-3. *Proceedings of the Royal Society of London, Series B* 1947;134:467-485.
6. Bjork A. Facial growth in man, studied with the aid of metallic implants. *Acta Odontologica Scandinavica* 1955-1956;13:9-34.
7. Brash JC. The growth of the jaws and palate *The Growth of the Jaws, Normal and Abnormal, in Health and Disease*. London: The Dental Board of the United Kingdom; 1924. p. 23-66.
8. Brodie AG. On the growth pattern of the human head from the third month to the eighth year of life. *American Journal of Anatomy* 1941;68:209-262.
9. Craven AH, Holland MS. Growth in width of the head of the Macaca Rhesis monkey as revealed by vital staining. *American Journal of Orthodontics* 1956;42:341-362.
10. Dubreuil G, Lacoste A. Les phénomènes de résorption dans les os de la voûte crânienne. Leur importance; leurs conséquences. *C R Association des Anatomistes* 1922;17:123-133.
11. Gans BJ, Sarnat BG. Sutural facial growth of the Macaca rhesus monkey: A gross and serial roentgenographic study by means of metallic implants. *American Journal of Orthodontics* 1951;37:827-841.

12. Gegenbaur C. Ueber die Bildung des Knochengewebes. *Jenaische Zeitschrift für Medizin und Naturwissenschaft* 1864;1:343-369.
13. Humphrey GM. On the growth of the jaws. *Transactions of the Cambridge Philosophical Society* 1871;11:1-5.
14. Keith A. *Menders of the Maimed: The Anatomical & Physiological Principles Underlying the Treatment of Injuries to Muscles, Nerves, Bones & Joints*. London: H. Frowde; Hodder & Stoughton; 1919.
15. Keith A, Campion GG. A contribution to the mechanism of growth of the human face. *International Journal of Orthodontia, Oral Surgery and Radiography* 1922;8:607-633.
16. Kölliker A. Die normale Resorption des Knochengewebes und ihre Bedeutung für die Entstehung der typischen Knochenformen. Leipzig: Vogel; 1873.
17. Kölliker A. *Embryologie ou Traité Complet du Développement de l'Homme et des Animaux Supérieurs*. Traduction faite sur la deuxième édition allemande par Schneider, A. Paris: C. Reinwald; 1882.
18. Lieberküehn N. Ueber den Abfall der Geweihe und seine Aehnlichkeit mit dem cariösen Process. *Archiv für Anatomie, Physiologie und Wissenschaftliche* 1861 (in Brash 1924):748-759.
19. Moore AW. Head growth of the macaque monkey as revealed by vital staining, embedding, and undecalcified sectioning. *American Journal of Orthodontics* 1949;35:654-671.
20. Schour I. Measurements of bone growth by alizarine injections. *Proceedings of the Society for Experimental Biology and Medicine* 1936;34:140-141.
21. Scott JH. The analysis of facial growth. I. The anteroposterior and vertical dimensions. *American Journal of Orthodontics* 1958;44:507-512.
22. Scott JH. The analysis of facial growth. II. The horizontal and vertical dimensions. *American Journal of Orthodontics* 1958;44:585-589.
23. Stricker S. *Manual of Human and Comparative Histology*. London: The New Sydenham Society; 1870 (with translation by Power, M.).
24. Tomes CS. *A System of Dental Surgery*. London: John Churchill; 1859.
25. Young RW. Postnatal growth of the frontal and parietal bones in white males. *American Journal of Physical Anthropology* 1957;15:367-386.
26. Enlow DH. *The Human Face: An Account of the Postnatal Growth and Development of the Craniofacial Skeleton*. New York: Harper and Row Pub., Inc.; 1968.
27. Moss ML, Salentijn L. The capsular matrix. *American Journal of Orthodontics* 1969;16:474-490.
28. Thilander B. Basic mechanisms in craniofacial growth. *Acta Odontologica Scandinavica* 1995;53:144-151.
29. Enlow DH. A Study of the Post-Natal Growth and Remodeling of Bone. *The American Journal of Anatomy* 1962;110:79-101.
30. Enlow DH. *Principles of Bone Remodeling: An Account of Postnatal Growth and Remodeling Processes in Long Bones and the Mandible*. Springfield: Charles C. Thomas; 1963.
31. Enlow DH. *Handbook of Facial Growth*. Toronto: W. B. Saunders Company; 1975.
32. Enlow DH, Hans MG. *Essentials of Facial Growth*. Ann Arbor, MI: Needham Press, Inc.; 2008.
33. Boyde A, Jones S. Scanning electron microscopic studies of the formation of mineralized tissues. In: Slavkin HC, Bavetta LA, editors. *Developmental Aspects of Oral Biology*. New York: Academic Press, Inc.; 1972. p. 243-274.

34. Baron R, Rawadi G, Roman-Roman S. Wnt Signaling: A key regulator of bone mass. *Current Topics in Developmental Biology* 2006;76:103-127.
35. Kini U, Nandeesh BN. Physiology of bone formation, remodeling, and metabolism. In: al. Fe, editor. *Radionuclide and Hybrid Bone Imaging*. Berlin: pringer-Verlag; 2012.
36. Rowe P, Koller A, Sharma S. Physiology, Bone Remodeling StatPearls [Internet]. Treasure Island (FL): StatPearls Publishing LLC.; 2020.
37. Ackermann RR. Ontogenetic integration of the hominoid face. *Journal of Human Evolution* 2005;48:175-197.
38. Rubin CT, Lanyon LE. Osteoregulatory nature of mechanical stimuli: Function as a determinant for adaptive bone remodeling. *Journal of Orthopaedic Research* 1987;5:300-310.
39. Lanyon LE. Functional strain as a determinant for bone remodeling. *Calcified Tissue International* 1984;36:556-561.
40. Christen P, Ito K, Ellouz R, Boutroy S, Sornay-Rendu E, Chapuriat RD et al. Bone remodelling in humans is load-driven but not lazy. *Nature Communications* 2014;5:4855.
41. Isaacson BM, Potter BK, Bloebaum RD, Epperson RT, Kawaguchi BS, Swanson BA et al. Link between clinical predictors of heterotopic ossification and histological analysis in combat-injured service members. *The Journal of Bone and Joint Surgery, American* 2016;98:647-657.
42. Hadjidakis D, Androulakis I. Bone remodeling. *Annals of the New York Academy of Sciences* 2006;1092:385-396.
43. Kim HJ, Yu SK, Lee MH, Lee HJ, Kim HJ, Chung CH. Cortical and cancellous bone thickness on the anterior region of alveolar bone in Korean: a study of dentate human cadavers. *The Journal of Advanced Prosthodontics* 2012;4:146-152.
44. Boivin G, Meunier PJ. Changes in Bone Remodeling Rate Influence the Degree of Mineralization of Bone. *Connective Tissue Research* 2002;43:535-537.
45. Neel EAA, Aljabo A, Strange A, Ibrahim S, Coathup M, Yopung AM et al. Demineralization–remineralization dynamics in teeth and bone. *International Journal of Nanomedicine* 2016;11:4743-4763.
46. Roschger P, Paschalis EP, Fratzl P, Klaushofer K. Bone mineralization density distribution in health and disease. *Bone* 2008;42:456-466.



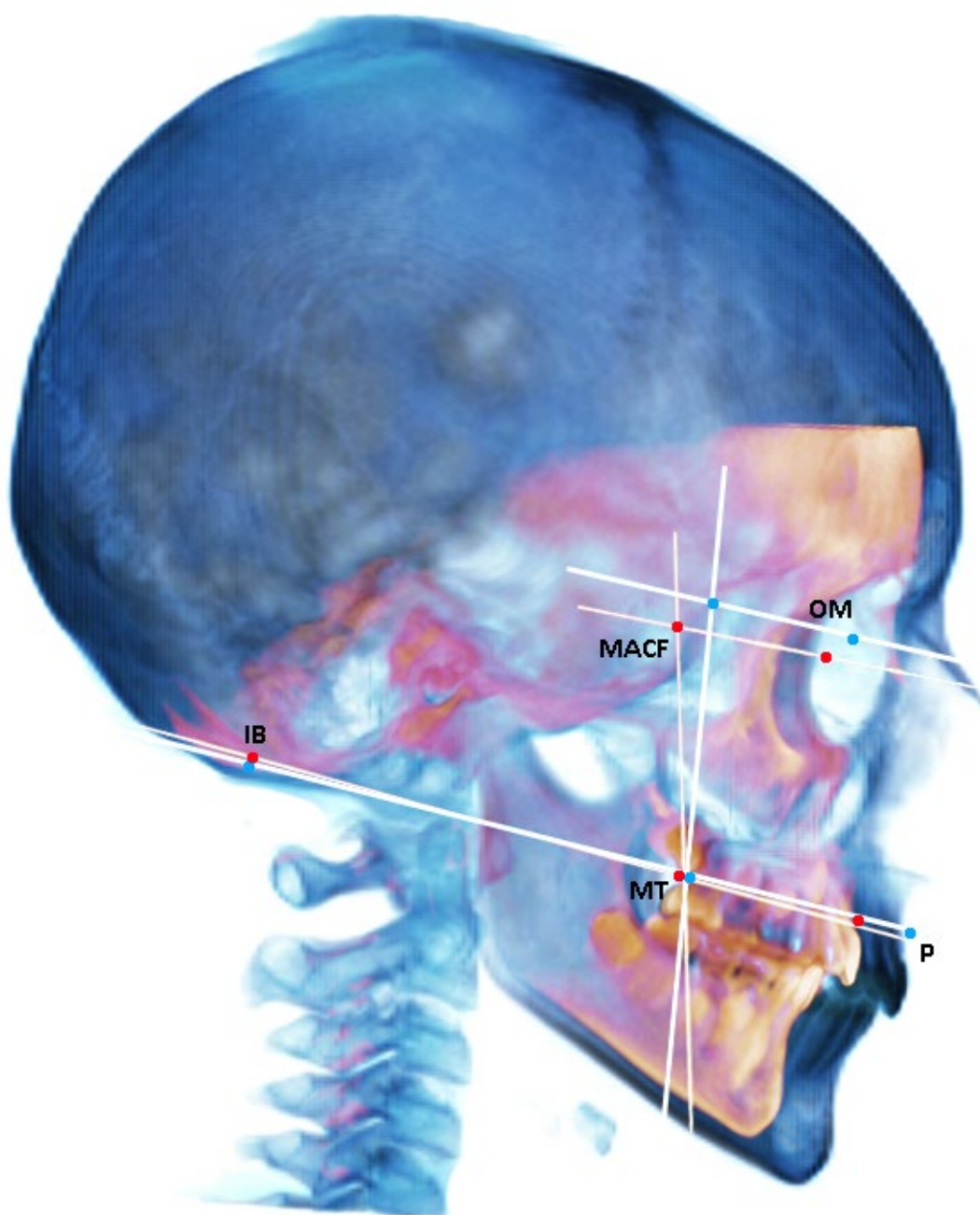


Figure 1

Younger (yellow) and older (blue) conebeam CT scans superimposed according to the method described by Enlow<sup>31</sup>, which describes what surfaces were forming and resorbing during growth. Landmarks include: junction between middle and anterior cranial fossae (MACF), orbital midpoint (OM), inferior brain (IB), maxillary tuberosity (MT), and prosthion (P). The disposition of the older scan indicates that growth of the maxilla was in the anterior direction, without a downward contribution. This result confirms that the therapy stimulated anteriorward growth and displacement and that the subnasal region was characterized by forming bone on its

surface, not resorbing bone as is typical of the growing child <sup>31</sup>, which would have led to rearward and downward growth. Note also the elevation of the orbits in this superimposition, which indicates significant deposits at sutures of the midface, displacing the upper face upward. The mandible has grown in harmony with the maxilla.

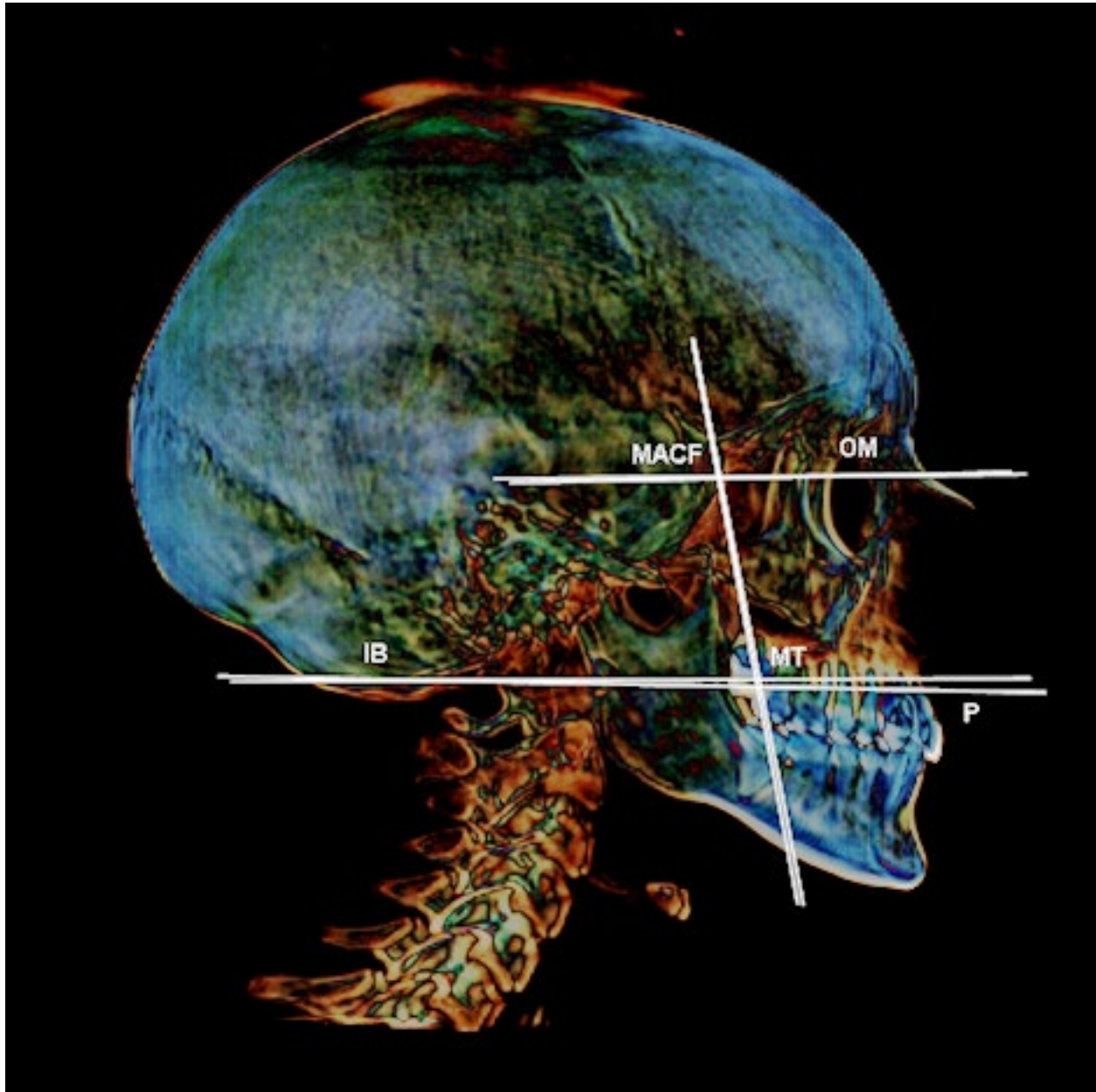


Figure 2

Younger and older conebeam CT scans superimposed according to the method described by Enlow <sup>31</sup>, which describes what surfaces were forming and resorbing during growth (see Fig. 1 for key to labels). The inferiorward relocation of the blue-green-colored subnasal region

indicates that the teeth and bone were relocated with a descending alveolus. The difference in height is indicated by the decent of the upper central incisor in white, and white along the intercuspal row of teeth. This was accompanied by an uprighting of the roots of the maxillary dentition, wherein before treatment is indicated by roots shaded in yellow. The maxillary unerupted third molar also rotated into its correct occluso-cervical orientation just behind MT point.