



Pesquisa Brasileira em Odontopediatria e Clinica Integrada 2018, 18(1):e3854 DOI: http://dx.doi.org/10.4034/PBOCI.2018.181.ed3 ISSN 1519-0501

GUEST EDITORIAL

Soft-tissue 3D Facial Imaging in Children and Adolescents: Towards the Definition of New Reference Standards

Chiarella Sforza¹, Claudia Dolci¹, Gianluca MartinoTartaglia¹, Virgilio Ferruccio Ferrario¹

¹Professor, Functional Anatomy Research Center, Department of Biomedical Sciences for Health, Faculty of Medicine, Università degli Studi di Milano, Milan, Italy.

In the last decades, head and face imaging has shifted from two-dimensional (2D) representations (conventional radiography, photography) to three-dimensional (3D) techniques that can better depict the complex morphology of this part of the body, since they can provide a large number of additional anthropometric information [1-3]. 3D imaging systems can be divided into volumetric (computed tomography, cone beam computed tomography, magnetic resonance imaging) and optical surface instruments (laser scanning, moiré techniques, stereophotogrammetry, patterned light techniques) [4]. These last are safe and not invasive, and provide a 3D representation of the external (cutaneous) facial surface.

Among all instruments, stereophotogrammetry is becoming the most diffused and used for both adults and children. Stereophotogrammetry is fast (typical scan time 2 ms), and it provides 3D photographs coupling a 3D mesh of the analyzed surface with a color facial image (texture). The technique takes photographs of the face from at least two different positions with two or more cameras (or set of cameras) at the same time. Using a previous calibration of the instrument, these coordinated photographs are combined to form a computerized stereoscopic reconstruction of the face [3,4]. Stereophotogrammetric systems have proven to be repeatable and able to provide accurate measurements [4,5].

Alongside with stereophotogrammetry, some laboratories and research groups are using laser scanners. The device illuminates the object with a low-intensity laser, and digital cameras capture the images. During data acquisition, either the laser light or the face rotate to sample the entire surface; triangulation geometry provides the depth information [6]. A multicentric study showed that laser scanning and stereophotogrammetric acquisitions can provide superimposable data, that can be efficiently shared among laboratories [7].

3D optical surface assessments can be of great value in clinical applications, at all ages of life, but especially for children where the concerns about radioprotection are maximized [3].



A PubMed search using the key-words "3D, face, human" retrieved 1628 full text papers published between 1965 and 2017; if "babies or children or child or adolescent" were added as keywords, the retrieved papers were 399 (https://www.ncbi.nlm.nih.gov/pubmed/, accessed on November, 4th 2017). Interestingly, the majority of these papers were published in the last 7 years (59% and 67%, respectively), thus confirming a growing interest in this topic, with an increment specifically higher for people in their first decades of life.

The fields of application of 3D facial imaging are diverse, ranging from those more frequently encountered by dentists to more rare disorders that anyway possess specific craniofacial characteristics of dental interest. Examples include the evaluation and treatment planning of children with occlusal abnormalities [8,9], orofacial clefts [10], or cranial deformities [11], as well as the early diagnosis of congenital and genetic disorders (Down's syndrome, ectodermal dysplasia, Glut1 deficiency syndrome, Moebius syndrome, velocardiofacial syndrome) or acquired pathologies like fetal alcohol syndrome that modify facial phenotype [12-18].

The use of new 3D techniques, with a better characterization of the anthropometric facial features, implies that these specific clinical purposes should be accompanied by the definition of appropriate reference values, that may be considered the new Bolton reference standards [19]. The formulation of reference standards requires the collection of cross-sectional (usually) and/or longitudinal (hopefully) data from normal subjects of comparable age, sex and ethnicity [3,20]. Indeed, even for genetic pathologies like Down's syndrome, where the original disorders is mostly the same all over the different ethnical groups may possess specific characteristics with a resulting particular phenotype [14,21]. Therefore, reference values should be collected in healthy people sharing the highest possible number of common factors with the patients.

It is clear that this task is actually very complex, and only a word-wide collaboration can succeed in the goal. There have already been some examples of multicentric data collection, but in most cases this was limited to adults [7,22]. For instance, data provided by Farkas et al. [222] were obtained in 1470 healthy adult subjects (18 to 30 years), equally divided in the two sexes. The largest group was made by European Caucasians, with minor contributions from Middle-East, Asia and Africa; data were compared to literature values for North American whites. Indeed, most of the values were obtained by conventional anthropometry, but the method used in this study could be taken as reference: the late professor Farkas invited colleagues who had published on the topic, defined the measurement protocol and the characteristics of the sample, and collected data. The original information was combined into a general framework of great interest: the paper is the most cited Farkas' one, with 230 citations (retrieved in Scopus https://www-scopus-com.pros.lib.unimi.it:2050/search/form.uri?display=basic, accessed on November, 12th 2017).

As far as children and adolescents are concerned, some research groups are working towards this goal. One initiative is FaceBase (https://www.facebase.org/, accessed on November, 12th 2017), a consortium launched in 2009 and funded by NIDCR (National Institute of Dental and Craniofacial



Research, USA), where scientists can find data supporting researches about craniofacial development and malformation. Alongside with genetic, molecular and biological data, the database also hosts facial images. Currently, there are photogrammetric scans of more than 700 healthy Caucasian American children from 3 to 12 years of age. The dataset is available to the research community, and it aims to foster and facilitate worldwide cooperation and collaboration. Weinberg et al. [3] recently published the results of an inter-centers collaboration based on FaceBase data repository, with data from 2,454 subjects of both sexes and aged from 3 years onwards. From the same source, Kesterke et al. [20] analyzed 1,555 healthy persons from 3 to 25 years of age.

In Europe, more than 2,500 German children aged 3-6 years were photographed in 3D by Moller et al. [23], while Bugaighis et al. [24] collected data on 80 Caucasian children from the North East of England aged 8–12 years. Forty-five Czech children were longitudinally studied between 12 and 15 years of age by Koudelová et al. [25] using stereophotogrammetry. Welsh children aged 11.8 years on average were analyzed by Kau et al. [6] using a laser scan. In our laboratory we had been collecting 3D data for the quantitative description of soft-tissue facial morphology for more than 25 years, providing reference values for more than 2,000 Italian children from 4 years of age onward [26,27].

Reference values for children belonging to countries and ethnic groups outside Northern America and Europe are actually very scanty. In Asia, Mori et al. [28] scanned in 3D the nasiolabial characteristics of Japanese boys and girls aged 5 to 6 years, while Al-Khatib et al. [29] studied the 3D characteristics of the nose of Malaysian adolescents aged 13 to 17 years. In Africa, Sforza et al. [30,31] published reference values for more than 650 Northern Sudanese people from 4 years of age into young adulthood.

Babies in the age range of 0-6 years have almost been neglected so far. Indeed, collaboration from very young persons (and from their families in some occasions) can be difficult, and the organization of data collection very demanding, even if the actual time necessary for the stereophotogrammetric or laser scans is very reduced. About one hundred Scottish babies aged around 3 months of age were imaged by both White et al. [32] and Hood et al. [33]. Other investigators started data collections at 3 [3,20,23] or 4 years of age [27].

In conclusion, 3D optical, surface imaging is becoming an important instrument for diagnosis, treatment planning and follow up of children, who should not receive ionizing radiations unless strictly necessary. The instruments are fast, safe, and neither painful nor fastidious. Their use for clinical applications should be accompanied by a systematic data collection in healthy young people, spanning from birth to late adolescence into young adulthood. Researchers all over the world should collaborate towards this goal, with the definition of these new facial standards.

References



1. Brons S, van Beusichem ME, Bronkhorst EM, Draaisma J, Bergé SJ, Maal TJ, et al. Methods to quantify soft-tissue based facial growth and treatment outcomes in children: a systematic review. PLoS One 2012; 7(8):e41898. doi: 10.1371/journal.pone.0041898.

2. Knoops PG, Beaumont CA, Borghi A, Rodriguez-Florez N, Breakey RW, Rodgers W, et al. Comparison of three-dimensional scanner systems for craniomaxillofacial imaging. J Plast Reconstr Aesthet Surg 2017; 70(4):441-9. doi: 10.1016/j.bjps.2016.12.015.

3. Weinberg SM, Raffensperger ZD, Kesterke MJ, Heike CL, Cunningham ML, Hecht JT, et al. The 3D facial norms database: part 1. A web-based craniofacial anthropometric and image repository for the clinical and research community. Cleft Palate Craniofac J 2016; 53(6):e185-e197.

4. Sforza C, de Menezes M, Ferrario V. Soft- and hard-tissue facial anthropometry in three dimensions: what's new. J Anthropol Sci 2013; 91:159-84. doi: 10.4436/jass.91007.

5. Andrade LM, Rodrigues da Silva AMB, Magri LV, Rodrigues da Silva MAM. Repeatability study of angular and linear measurements on facial morphology analysis by means of stereophotogrammetry. J Craniofac Surg 2017; 28(4):1107-11. doi: 10.1097/SCS.00000000003554.

6. Kau CH, Zhurov A, Richmond S, Bibb R, Sugar A, Knox J, et al. The 3-dimensional construction of the average 11-year-old child face: a clinical evaluation and application. J Oral Maxillofac Surg 2006; 64(7):1086-92.

7. Kau CH, Richmond S, Zhurov A, Ovsenik M, Tawfik W, Borbely P, et al. Use of 3-dimensional surface acquisition to study facial morphology in 5 populations. Am J Orthod Dentofacial Orthop 2010; 137(4 Suppl):S56.e1-9. doi: 10.1016/j.ajodo.2009.04.022.

8. Godt A, Bechtold TE, Schaupp E, Zeyher C, Koos B, Baas E, et al. Correlation between occlusal abnormalities and parameters investigated by three-dimensional facial photography. Angle Orthod 2013; 83(5):782-9. doi:10.2319/111412-874.1.

9. Primožič J, Richmond S, Kau CH, Zhurov A, Ovsenik M. Three-dimensional evaluation of early crossbite correction: a longitudinal study. Eur J Orthod 2013; 35(1):7-13. doi: 10.1093/ejo/cjq198.

10. Ort R, Metzler P, Kruse AL, Matthews F, Zemann W, Grätz KW, et al. The reliability of a threedimensional photo system- (3dMDface-) based evaluation of the face in cleft lip infants. Plast Surg Int 2012; 2012:138090. doi: 10.1155/2012/138090.

11. Schaaf H, Pons-Kuehnemann J, Malik CY, Streckbein P, Preuss M, Howaldt HP, et al. Accuracy of threedimensional photogrammetric images in non-synostotic cranial deformities. Neuropediatrics 2010; 41(1):24–9. doi: 10.1055/s-0030-1255060.

12. Dellavia C, Catti F, Sforza C, Tommasi DG, Ferrario VF. Craniofacial growth in ectodermal dysplasia. An 8 year longitudinal evaluation of Italian subjects. Angle Orthod 2010; 80(4):733-9. doi: 10.2319/101909-584.1.

13. Lewyllie A, Roosenboom J, Indencleef K, Claes P, Swillen A, Devriendt K, et al. A comprehensive craniofacial study of 22q11.2 deletion syndrome. J Dent Res 2017; 96(12):1386-91. doi: 10.1177/0022034517720630.

14. Kruszka P, Porras AR, Sobering AK, Ikolo FA, La Qua S, Shotelersuk V, et al. Down syndrome in diverse populations. Am J Med Genet A 2017; 173(1):42-53. doi: 10.1002/ajmg.a.38043.

15. Mutsvangwa TE, Meintjes EM, Viljoen DL, Douglas TS. Morphometric analysis and classification of the facial phenotype associated with fetal alcohol syndrome in 5- and 12-year-old children. Am J Med Genet A 2010; 152A(1):32-41. doi: 10.1002/ajmg.a.33137.

16. Pucciarelli V, Bertoli S, Codari M, De Amicis R, De Giorgis V, Battezzati A, et al. The face of Glut1-DS patients: a 3D craniofacial morphometric analysis. Clin Anat 2017; 30(5):644-52. doi: 10.1002/ca.22890.

17. Sforza C, Dellavia C, Dolci C, Donetti E, Ferrario VF. A quantitative three-dimensional assessment of abnormal variations in the facial soft tissues of individuals with Down syndrome. Cleft Palate Craniofac J 2005; 42(4):410-6.

Sforza C, Grandi G, Pisoni L, Di Blasio C, Gandolfini M, Ferrario VF. Soft tissue facial morphometry in subjects with Moebius syndrome. Eur J Oral Sci 2009; 117(6):695-703. doi: 10.1111/j.1600-0722.2009.00685.x.
Broadbent Sr. BH, Broadbent Jr. BH, Golden WH. Bolton standards of dentofacial developmental growth. Saint Louis: Mosby; 1975.

20. Kesterke MJ, Raffensperger ZD, Heike CL, Cunningham ML, Hecht JT, Kau CH, et al. Using the 3D Facial Norms Database to investigate craniofacial sexual dimorphism in healthy children, adolescents, and adults. Biol Sex Differ 2016; 7:23. doi: 10.1186/s13293-016-0076-8.



21. Sforza C, Dolci C, Dellavia C, Gibelli DM, Tartaglia GM, Elamin F. Abnormal variations in the facial soft tissues of individuals with Down syndrome: Sudan versus Italy. Cleft Palate Craniofac J 2015; 52(5):588-96. doi:10.1597/14-082.

22. Farkas LG, Katic MJ, Forrest CR, Alt KW, Bagic I, Baltadjiev G, et al. International anthropometric study of facial morphology in various ethnic groups/races. J Craniofac Surg 2005; 16(4):615-46.

23. Moller M, Schaupp E, Massumi-Moller N, Zeyher C, Godt A, Berneburg M. Reference values for threedimensional surface cephalometry in children aged 3-6 years. Orthod Craniofac Res 2012; 15:103-16. doi: 10.1111/j.1601-6343.2012.01541.x.

24. Bugaighis I, Mattick CR, Tiddeman B, Hobson R. Three-dimensional gender differences in facial form of children in the North East of England. Eur J Orthod 2013; 35(3):295-304. doi: 10.1093/ejo/cjr033.

25. Koudelová J, Dupej J, Brůžek J, Sedlak P, Velemínská J. Modelling of facial growth in Czech children based on longitudinal data: Age progression from 12 to 15 years using 3D surface models. Forensic Sci Int 2015; 248:33-40. doi: 10.1016/j.forsciint.2014.12.005.

26. Ferrario VF, Sforza C, Poggio CE, Schmitz JH. Soft-tissue facial morphometry from 6 years to adulthood: a three-dimensional growth study using a new modeling. Plast Reconstr Surg 1999; 103(3):768-78.

27. Sforza C, Grandi G, Binelli M, Dolci C, De Menezes M, Ferrario VF. Age- and sex-related changes in three-dimensional lip morphology. Forensic Sci Int 2010; 200(1-3):182.e1-7. doi: 10.1016/j.forsciint.2010.04.050.

28. Mori A, Nakajima T, Kaneko T, Sakuma H, Aoki Y. Analysis of 109 Japanese children's lip and nose shapes using 3-dimensional digitizer. Br J Plast Surg 2005; 58(3):318-29.

29. Al-Khatib AR, Rajion ZA, Masudi SM, Hassan R, Anderson PJ, Townsend GC. Stereophotogrammetric analysis of nasolabial morphology among Asian Malays: influence of age and sex. Cleft Palate Craniofac J 2012; 49(4):463-71. doi: 10.1597/11-151.

30. Sforza C, Dolci C, Tommasi DG, Pisoni L, De Menezes M, Elamin F. Three-dimensional facial distances of Northern Sudanese persons from childhood to young adulthood. J Craniomaxillofac Surg 2014; 42(5):e318-26. doi: 10.1016/j.jcms.2013.10.013.

31. Sforza C, Dolci C, Gibelli DM, Codari M, Pucciarelli V, Ferrario VF, et al. Age-related and sex-related changes in the normal soft tissue profile of native Northern Sudanese subjects: a cross-sectional study. Br J Oral Maxillofac Surg 2016; 54(2):192-7. doi: 10.1016/j.bjoms.2015.11.015.

32. White JE, Ayoub AF, Hosey MT, Bock M, Bowman A, Bowman J, et al. Three-dimensional facial characteristics of Caucasian infants without cleft and correlation with body measurements. Cleft Palate Craniofac J 2004; 41:593-602.

33. Hood CA, Hosey MT, Bock M, White J, Ray A, Ayoub AF. Facial characterization of infants with cleft lip and palate using a three-dimensional capture technique. Cleft Palate Craniofac J 2004; 41(1):27-35.

Correspondence: Prof. Chiarella Sforza, Department of Biomedical Sciences for Health, University of Milan, via Mangiagalli 31 - I-20133, Milan, Italy. Phone: +39 02 503 15385. Email: chiarella.sforza@unimi.it.