



Genetics aspect of hypodontia in twins (Case report)

Yuke Rustan, Hendrarlin Soenawan

Department of Pedodontics, Faculty of Dentistry, University of Indonesia



Introduction

The term 'hypodontia' refers to absence of one or more tooth in developmental stage, either in primary or permanent dentition. Permanent dentition is more frequently affected than primary dentition, with incidence varies from 1.6% to 9.6% in the general population. A study conducted in Asian population indicates that the most frequent absent is mandibular incisor¹. Fervently genetic influence in hypodontia has been established by many studies performed on families and twins². This case report present 9 years old girls, twins, who has no mandibular lateral incisors but they did not aware about this.

Case Report

A 9 years old girl came to Pediatric clinic for general check up. During intra oral examination, mandibular lateral incisors was found missing. To confirm the condition, an orthopantomogram was taken, it was confirmed as a hypodontia. From alloanamnesis with the parents, the girl is twins. Next week, her sibling came to the Pediatric clinic. She has no mandibular lateral incisor as well. Orthopantomogram confirming the condition as a hypodontia.

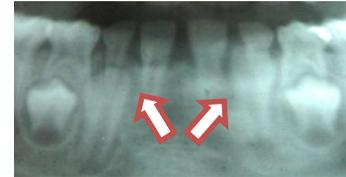
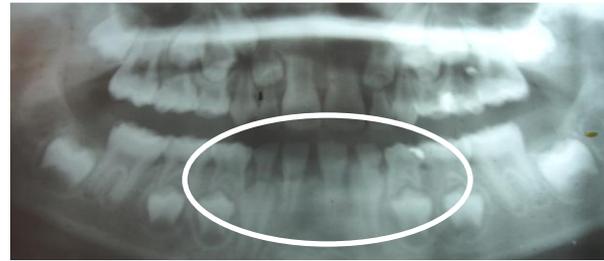


Fig1. Twin A orthopantomogram and intra oral photographs

Discussion

Known from the literature, the diagnose of hypodontia in permanent teeth was concluded on 12-14 years old children. The condition is re-confirm with orthopantomogram³. Several assumption came for this condition such as infections, toxins and chemotherapy. However, most cases were found connected with genetic factors. The heritability of congenital missing tooth has been shown in many studies. The importance of genetic factors is shown by appearance of multiple cases among families and higher concordance in monozygotic than dizygotic twins^{4,5}. Those of particular interest in tooth development are the homeobox genes *Msx1* and *Msx2* which is known as transcription factors and controlling gene expression. In humans, genetic analysis of a twins has demonstrated a mutation in the *Msx1* gene⁵.

Conclusion

In this case, both twins and parents didn't realize about the teeth missing and didn't complain about the teeth alignment. So, there is no treatment done but plan to do annual regular check up. Tooth development is a very complex process and involving many "players." The *Msx1* gene is one of them⁵. Reports done with confirmation that environmental or epigenetic factors can modify phenotype expression and involved in the multifactorial etiology of hypodontia.

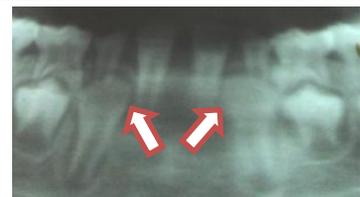
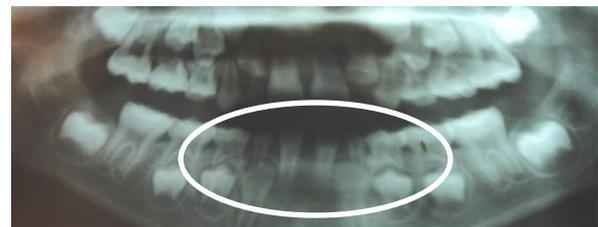


Fig2. Twin B orthopantomogram and intra oral photographs

1. Davis PJ. Hypodontia and hyperdontia of permanent teeth in Hong Kong school children community. *Dent oral epidemiology* 1987; 15: 218-20.
2. Larmour CJ, Mossey PA, Thind BS, et al. *Hypodontia-A retrospective review of prevalence and etiology*. Part 1.
3. Arte S, Pirinen S. *Hypodontia*. *Orphanet encyclopedia*. May 2004. Downloaded from <http://www.orpha.net/data/patho/GB/uk-hypodontia.pdf>
4. [Kaminsky ZA](#), [Tang J](#), [Wang SC](#). DNA methylation profiles in monozygotic and dizygotic twins. *Nat Genet*. 2009 Feb;41(2):240-5.
5. Suarez BK, Spence MA. The Genetics of Hypodontia. *J Dent Res* July-August 1974 : 781-5.