Multiple malformation syndrome including cleft lip and palate and cardiac abnormalities due to an interstitial deletion of chromosome 12q

The proband, a male, was the second child of young, healthy parents. The older brother is healthy. The pregnancy progressed normally with spontaneous delivery at 39 weeks. Birth weight was 2030 g, length 44 cm, and head circumference 33 cm. Craniofacial dysmorphism included hypertelorism, upward slanting palpebral fissures, broad nasal bridge, bilateral cleft lip and palate, macrostomia, retrognathia, and deep-set, dorsi rotated auricles. There was overriding of the second and fifth fingers bilaterally, rocker bottom feet, and short big toes. Cardiac findings suggested an atrial septal defect and a superior ventricular septal defect. Recurrent vomiting and haematemesis was observed and the child's development was retarded. He died at 10 weeks and the parents refused necropsy.

Chromosomal analysis of lymphocyte cultures showed an interstitial deletion of the long arm of chromosome 12: 46,XY.del(12)(q13.3→q21). Fibroblast cultures obtained post mortem confirmed this finding. Parental karyotypes were normal.

The phenotype of the child was suggestive of trisomy 18. The cleft lip and palate could be consistent with this diagnosis, as about 25% of patients with trisomy 18 exhibit this feature. It appears that interstitial deletion of the long arm of chromosome 12 is able to mimic the phenotype of trisomy 18 to a considerable extent. To our knowledge, no similar case of interstitial deletion 12q has yet been reported, and recent text books do not provide any appropriate references. Therefore, it is not certain whether or not a constant karyotype/phenotype correlation exists. If this correlation does exist, then this type of interstitial deletion could be described as another example of 'pseudo-trisomy 18'. A further nosological aspect of this case is to add another example to an already lengthy list of syndromes which feature cleft lip and palate.

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References

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