Deafness-oligodontia syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Deafness-oligodontia syndrome. ORPHA:3230

Deafness-oligodontia syndrome is characterised by sensorineural hearing loss and oligodontia/hypodontia. It has been described in two pairs of siblings and in one isolated case. Dizziness was reported in one of the pairs of siblings. Transmission appears to be autosomal recessive.