Open Peer Review on Qeios

Odontoleukodystrophy

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Odontoleukodystrophy. ORPHA:77295

Leukodystrophy with oligodontia is characterised by progressive ataxia beginning during infancy, a pyramidal syndrome and dental agenesis. The syndrome has been described in four children born to consanguineous parents. The mode of transmission is autosomal recessive.