Dentin dysplasia

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Dentin dysplasia. ORPHA:1653

Dentin dysplasia (DD) is a rare disorder belonging to the group of hereditary dentin defects (see this term) and is characterized by abnormal dentin structure and root development resulting in abnormal tooth development. It encompasses two subtypes: DD type I and DD type II (see these terms).