DISEASE:
Oculodentodigital dysplasia

NAME:	Oculodentodigital dysplasia
DESCRIPTION:	A rare congenital malformation syndrome characterized by craniofacial, ocular, dental, digital anomalies and neurologic symptoms.
ORPHACODE:	2710
SYNONYMS:	Meyer-Schwickerath syndrome ODDD syndrome Oculodentoosseous dysplasia
XREF(S):	Orphanet OMIM OMIM MedDRA ICD-10
ANALYTE(S):	GJA1
CREATED:	13 May 2019 - 01:02
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RELATED CONTENT

Related Genetic Tests

- Cataract (gene panel)
- Oculo Dento Digital Dysplasia
- Primary lymphedema / fetal hydrops (gene panel)
- Spastic Paraplegia (gene panel)

Related Laboratories

- Centre de Génétique Médicale UCL
- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)
- Centrum Medische Genetica - UZ Gent

Related Analytes

- gap junction protein alpha 1

Related Gene Panels

- Cataract - UGent
- Spastic Paraplegia (89 genes) - IPG
- test