DISEASE:
Oligodontia

NAME:	Oligodontia
DESCRIPTION:	Oligodontia is a rare developmental dental anomaly in humans characterized by the absence of six or more teeth.
ORPHACODE:	99798
SYNONYMS:	Selective tooth agenesis
XREF(S):	Orphanet OMIM OMIM OMIM ICD-10 OMIM OMIM OMIM OMIM OMIM

ANALYTE(S):	SUMO1 AXIN2 EDA EDARADD FGFR1 IRF6 WNT10A MSX1 PAX9 TGFA LRP6 WNT10B
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

RELATED CONTENT

Related Genetic Tests

- Ectrodactyly / cleft lip/palate syndrome type 3 / Ectodermal dysplasia
- cleft lip with/whitout cleft palate (virtual gene panel)

Related Laboratories

- Centre de Génétique Médicale UCL
- Centrum Medische Genetica - UZ Gent

Related Analytes

- axin 2
- ectodysplasin A
- EDAR associated via death domain
- fibroblast growth factor receptor 1
- interferon regulatory factor 6
- LDL receptor related protein 6
- msh homeobox 1
- paired box 9
- small ubiquitin like modifier 1
- transforming growth factor alpha
- Wnt family member 10A
- Wnt family member 10B

Related Gene Panels

- Cleft lip and palate / dysmorphic facial features / craniofacial anomalies (255 genes)) - UCL
- Ectrodactyly / cleft lip/palate / Ectodermal dysplasia - UGent

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Source URL: http://gentest.healthdata.be/disease/3811