DISEASE:
Craniofrontonasal dysplasia

NAME:	Craniofrontonasal dysplasia
DESCRIPTION:	A rare X-linked malformation syndrome characterized by craniofacial abnormalities, grooved nails, intellectual disability and various skeletal and soft tissue abnormalities.
ORPHACODE:	1520
SYNONYMS:	CFND CFNS Craniofrontonasal syndrome
XREF(S):	Orphanet ICD-10 OMIM MeSH
ANALYTE(S):	EFNB1
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RELATED CONTENT

Related Genetic Tests

- Craniosynostosis (gene panel)
- cleft lip with/whitout cleft palate (virtual gene panel)

Related Laboratories

- Centre de Génétique Médicale UCL
- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- ephrin B1

Related Gene Panels

- Cleft lip and palate / dysmorphic facial features / craniofacial anomalies (255 genes)) - UCL
- Craniosynostosis (32 genes) - KUL