DISEASE:
Hydrolethalus

NAME:	Hydrolethalus
DESCRIPTION:	Hydrolethalus (HLS) is a severe fetal malformation syndrome characterized by craniofacial dysmorphic features, central nervous system, cardiac, respiratory tract and limb abnormalities.
ORPHACODE:	2189
XREF(S):	Orphanet ICD-10 OMIM OMIM MeSH
ANALYTE(S):	HYLS1 KIF7
CREATED:	13 May 2019 - 01:02
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RELATED CONTENT

Related Genetic Tests

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

Related Laboratories

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

Related Analytes

- HYLS1 centriolar and ciliogenesis associated
- kinesin family member 7

Related Gene Panels

- Ciliopathy (120 genes) - UGent
- Cleft lip and palate / dysmorphic facial features / craniofacial anomalies (255 genes)) - UCL