DISEASE:
Trisomy 13

NAME:	Trisomy 13
DESCRIPTION:	Trisomy 13 is a chromosomal anomaly caused by the presence of an extra chromosome 13 and is characterized by brain malformations (holoprosencephaly), facial dysmorphism, ocular anomalies, postaxial polydactyly, visceral malformations (cardiopathy) and severe psychomotor retardation.
ORPHACODE:	3378
SYNONYMS:	Patau syndrome
XREF(S):	Orphanet
CREATED:	24 Jul 2019 - 10:28
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RELATED CONTENT

Related Genetic Tests

- Non Invasive Prenatal Testing (NIPT) of trisomies 13, 18 et 21 and sex chromosomes
- Non Invasive Prenatal Testing (NIPT) of trisomies 13, 18 et 21 and sex chromosomes
- Non Invasive Prenatal Testing (NIPT) of trisomies 13, 18 et 21 and sex chromosomes

Related Laboratories

- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)
- Centrum Medische Genetica - UZ Gent
- Centrum Menselijke Erfelijkheid - KUL