Belgian Genetic Tests database

DISEASE:
Trisomy 18

NAME:	Trisomy 18
DESCRIPTION:	Trisomy 18 is a chromosomal abnormality associated with the presence of an extra chromosome 18 and characterized by growth delay, dolichocephaly, a characteristic facies, limb anomalies and visceral malformations.
ORPHACODE:	3380
XREF(S):	Orphanet
CREATED:	03 Sep 2019 - 10:44
CHANGED:	03 Sep 2019 - 10:44

Source URL: http://gentest.healthdata.be/disease/3854

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