DISEASE:
Feingold syndrome type 1

NAME:	Feingold syndrome type 1
DESCRIPTION:	A rare, genetic congenital malformation syndrome characterized by digital anomalies (shortening of the 2nd and 5th middle phalanx of the hand, clinodactyly of the 5th finger, syndactyly of toes 2-3 and/or 4-5, thumb hypoplasia), microcephaly, facial dysmorphism (short palpebral fissures and micrognathia), gastrointestinal atresia (primarily esophageal and/or duodenal), and mild-to-moderate learning disability.
ORPHACODE:	391641
SYNONYMS:	Brunner-Winter syndrome type 1 Digital anomalies with short palpebral fissures and atresia of esophagus or duodenum type 1 FGLDS1 FS1 MMT type 1 MODED syndrome type 1 Microcephaly-digital anomalies-normal intelligence syndrome type 1 Microcephaly-intellectual disability-tracheoesophageal fistula syndrome type 1 Microcephaly-oculo-digito-esophageal-duodenal syndrome syndrome type 1 ODED syndrome type 1 Oculo-digito-esophageal-duodenal syndrome type 1

XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	MYCN
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

RELATED CONTENT

Related Genetic Tests

- Feingold syndrome

Related Laboratories

- Centrum Medische Genetica - UZ Gent

Related Analytes

- MYCN proto-oncogene, bHLH transcription factor