DISEASE:
X-linked hyper-IgM syndrome

NAME:	X-linked hyper-IgM syndrome
ORPHACODE:	101088
SYNONYMS:	HIGM1 Hyper-IgM syndrome due to CD40 ligand deficiency Hyper-IgM syndrome due to CD40L deficiency Hyper-IgM syndrome type 1 XHIGM
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	CD40LG
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

RELATED CONTENT

Related Genetic Tests

- Immune deficiency, X-linked, with hyperIgM
- Primary immune deficiencies (gene panel)

Related Laboratories

- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- CD40 ligand

Related Gene Panels

- Immunogenetics (21 genes)
- Primary immune deficiencies (444 genes) - KUL