

DISEASE:
Common variable immunodeficiency

NAME:	Common variable immunodeficiency
DESCRIPTION:	Common variable immunodeficiency (CVID) comprises a heterogeneous group of diseases characterized by a significant hypogammaglobulinemia of unknown cause, failure to produce specific antibodies after immunizations and susceptibility to bacterial infections, predominantly caused by encapsulated bacteria.
ORPHACODE:	1572
SYNOMYS:	CVID Idiopathic immunoglobulin deficiency Primary antibody deficiency Primary hypogammaglobulinemia

XREF(S):	Orphanet MeSH MedDRA ICD-10 ICD-10 ICD-10 ICD-10 ICD-10 OMIM OMIM OMIM OMIM OMIM OMIM OMIM OMIM OMIM
ANALYTE(S):	CD19 ICOS CR2 TNFRSF13C MS4A1 CD81 PRKCD TNFSF12 TNFSF12 NFKB1 NFKB2 IRF2BP2 TNFRSF13B

CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

Source URL: <http://gentest.healthdata.be/disease/1391>

RELATED CONTENT

Related Genetic Tests

- [Primary immune deficiencies \(gene panel\)](#)

Related Laboratories

- [Centrum Menselijke Erfelijheid - KUL](#)

Related Analytes

- [CD19 molecule](#)
- [CD81 molecule](#)
- [complement C3d receptor 2](#)
- [inducible T cell costimulator](#)
- [interferon regulatory factor 2 binding protein 2](#)
- [membrane spanning 4-domains A1](#)
- [nuclear factor kappa B subunit 1](#)
- [nuclear factor kappa B subunit 2](#)
- [protein kinase C delta](#)
- [TNF receptor superfamily member 13B](#)
- [TNF receptor superfamily member 13C](#)
- [TNF superfamily member 12](#)

Related Gene Panels

- Primary immune deficiencies (444 genes) - KUL

Source URL: <http://gentest.healthdata.be/disease/1391>