

ANALYTE:
G6PC3

NAME:	glucose-6-phosphatase catalytic subunit 3
SYMBOL:	G6PC3
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	UGRP
XREF(S):	<u>Orphanet</u> <u>Ensembl</u> <u>Genatlas</u> <u>HGNC</u> <u>OMIM</u> <u>Reactome</u> <u>SwissProt</u>
CREATED:	13 May 2019 - 01:01
CHANGED:	22 Jun 2023 - 16:14

Source URL: <http://gentest-acc.healthdata.be/analyte/2515>

RELATED CONTENT

Related Genetic Tests

- [Congenital disorders of glycosylation \(gene panel\)](#)
- [Congenital malformation \(gene panel - 1721 genes\)](#)
- [Congenital malformation \(gene panel\)](#)
- [Congenital structural heart defects \(gene panel\)](#)
- [Intellectual disability \(virtual gene panel\)](#)
- [Myeloid/lymphoid neoplasms with germline predisposition \(gene panel\)](#)
- [Neurodevelopmental disorders \(1300 genes\)](#)
- [Neurodevelopmental disorders \(gene panel\)](#)
- [Primary immune deficiencies \(gene panel\)](#)
- [Primary immune deficiencies \(gene panel\)](#)
- [« Inherited bone marrow failures syndromes » with or without organ dysfunction](#)

Related Diseases

- [Autosomal recessive severe congenital neutropenia due to G6PC3 deficiency](#)

Related Gene Panels

- [Congenital disorders of glycosylation \(100 genes\) - KUL](#)
- [Congenital malformation \(1721 genes\) - ULB](#)
- [Congenital malformation gene panel - VUB](#)
- [Congenital structural heart defects - UGent](#)
- [Hematologic Familiar Forms - ULG](#)
- [Intellectual disability \(gene panel\)](#)

- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- Primary immune deficiencies (444 genes) - KUL
- Primary immune deficiencies - UGent

Source URL: <http://gentest-acc.healthdata.be/analyte/2515>