

DISEASE:
Leukocyte adhesion deficiency type II

NAME:	Leukocyte adhesion deficiency type II
DESCRIPTION:	Leukocyte adhesion deficiency type II (LAD-II) is a form of LAD (see this term) characterized by recurrent bacterial infections, severe growth delay and severe intellectual deficit.
ORPHACODE:	99843
SYNONYMS:	CDG syndrome type IIc CDG-IIc CDG2C LAD-II Rambam-Hasharon syndrome SLC35C1-CDG
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	SLC35C1
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

Source URL: <http://gentest.healthdata.be/disease/3799>

RELATED CONTENT

Related Genetic Tests

- Congenital disorders of glycosylation (79 genes)

Related Laboratories

- Centrum Menselijke Erfelijheid - KUL

Related Analytes

- solute carrier family 35 member C1

Related Gene Panels

- Congenital disorders of glycosylation (79 genes) - KUL

Source URL: <http://gentest.healthdata.be/disease/3799>