Belgian Genetic Tests database

DISEASE:
Autosomal recessive severe congenital neutropenia due to CXCR2 deficiency

NAME:	Autosomal recessive severe congenital neutropenia due to CXCR2 deficiency
DESCRIPTION:	A rare, genetic, primary immunodeficiency disorder characterized by recurrent bacterial infections (including septic thrombophlebitis and subacute bacterial endocarditis) and neutropenia without lymphopenia or warts, resulting from recessively inherited mutations in CXCR2.
ORPHACODE:	420699
XREF(S):	Orphanet
ANALYTE(S):	CXCR2
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Source URL: http://gentest.healthdata.be/disease/2516

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