Belgian Genetic Tests database

DISEASE:
Hereditary coproporphyria

NAME:	Hereditary coproporphyria
DESCRIPTION:	Hereditary coproporphyria is a form of acute hepatic porphyria (see this term) characterized by the occurrence of neuro-visceral attacks and, more rarely, by the presence of cutaneous lesions.
ORPHACODE:	79273
XREF(S):	Orphanet MeSH MedDRA ICD-10 OMIM
ANALYTE(S):	CPOX
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

Source URL: http://gentest.healthdata.be/disease/3280

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