DISEASE:
Sickle cell-hemoglobin D disease syndrome

NAME:	Sickle cell-hemoglobin D disease syndrome
DESCRIPTION:	A rare, genetic hemoglobinopathy characterized by all the characteristics of sickle cell anemia (SCA). Clinical course is similar to SCA, including acute episodes of pain, splenic infarction and splenic sequestration crisis, vaso-occlusive crisis, acute chest syndrome, ischemic brain injury, osteomyelitis and avascular bone necrosis. The genotype is characterized by an HbS allele in combination with the HbD variant, beta121Glu>Gln.
ORPHACODE:	251370
SYNONYMS:	HbSD disease
XREF(S):	Orphanet MedDRA ICD-10
ANALYTE(S):	HBB
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

RELATED CONTENT

Related Genetic Tests

- Beta-globin hemoglobinopathies
- Beta-globin hemoglobinopathies
- Beta-globin hemoglobinopathies (full sequencing)

Related Laboratories

- Centre de Génétique Humaine - CHU Sart-Tilman
- Centre de Génétique Humaine - Erasme ULB
- Centre de Génétique Médicale UCL

Related Analytes

- hemoglobin subunit beta

Source URL: http://gentest.healthdata.be/disease/941