DISEASE:
Legius syndrome

NAME:	Legius syndrome
DESCRIPTION:	Legius syndrome, also known as NF1-like syndrome, is a rare, genetic skin pigmentation disorder characterized by multiple café-au-lait macules with or without axillary or inguinal freckling.
ORPHACODE:	137605
SYNONYMS:	NF1-like syndrome Neurofibromatosis 1-like syndrome Nonmosaic LGSS Nonmosaic Legius syndrome
XREF(S):	Orphanet OMIM MeSH ICD-10
ANALYTE(S):	SPRED1
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

RELATED CONTENT

Related Genetic Tests

- Neurofibromatosis type 1 / Legius syndrome
- Neurofibromatosis type 1 / Legius syndrome (2 genes)

Related Laboratories

- Centrum Medische Genetica - UZ Gent
- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- sprouty related EVH1 domain containing 1
