DISEASE:
Dermatosparaxis Ehlers-Danlos syndrome

NAME:	Dermatosparaxis Ehlers-Danlos syndrome
DESCRIPTION:	A form of Ehlers-Danlos syndrome (EDS) characterized by extreme skin fragility and laxity, a prominent facial gestalt, excessive bruising and, sometimes, major complications due to visceral and vascular fragility.
ORPHACODE:	1901
SYNONYMS:	Dermatosparaxis EDS Ehlers-Danlos syndrome type 7C Human dermatosparaxis EDS VIIC dEDS
XREF(S):	Orphanet OMIM ICD-10
ANALYTE(S):	ADAMTS2 ADAMTSL2
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RELATED CONTENT

Related Genetic Tests

- Ehlers-Danlos syndroom, EDS (gene panel)
- Epidermolysis bullosa (gene panel)

Related Laboratories

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

Related Analytes

- ADAM metallopeptidase with thrombospondin type 1 motif 2
- ADAMTS like 2

Related Gene Panels

- Ehlers-Danlos syndrome -UGent
- Epidermolysis bullosa and bladder diseases (60 genes) - KUL
- Recessive Ehlers-Danlos Syndrome (11 genes) - UGent