Belgian Genetic Tests database

DISEASE:
Classical Ehlers-Danlos syndrome

NAME:	Classical Ehlers-Danlos syndrome
DESCRIPTION:	A rare inherited connective tissue disorder characterized by skin hyperextensibility, widened atrophic scars, and generalized joint hypermobility.
ORPHACODE:	287
SYNONYMS:	Classical EDS cEDS
XREF(S):	Orphanet OMIM OMIM ICD-10
ANALYTE(S):	COL5A2 COL5A1 COL1A1
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

Source URL: http://gentest.healthdata.be/disease/353

RELATED CONTENT

Related Genetic Tests

- Aneurysm, Thoracic Aortic, familial (gene panel)
- Corneal dystrophy (gene panel)
- Ehlers-Danlos syndroom, EDS (gene panel)
- Osteogenesis Imperfecta (gene panel)

Related Laboratories

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

Related Analytes

- collagen type I alpha 1 chain
- collagen type V alpha 1 chain
- collagen type V alpha 2 chain

Related Gene Panels

- Corneal dystrophy - UGent
- Ehlers Danlos classic type (2 genes) - UGent
- Ehlers-Danlos syndrome -UGent

- Familial Thoracic Aortic Aneurysm (genepanel) - UZA
- Osteogenesis Imperfecta (25 genes) - KUL
