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Full name:	Ehlers-Danlos syndrome -UGent
Abbreviation:	EDS
Laboratory:	Centrum Medische Genetica - UZ Gent
Created:	11 Dec 2019 - 13:31
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Related Diseases

- Arthrochalasia Ehlers-Danlos syndrome
- B3GALT6-related spondylodysplastic Ehlers-Danlos syndrome
- B4GALT7-related spondylodysplastic Ehlers-Danlos syndrome
- Bethlem myopathy
- Brittle cornea syndrome
- Cardiac-valvular Ehlers-Danlos syndrome
- Classical Ehlers-Danlos syndrome
- Classical-like Ehlers-Danlos syndrome type 2
- Congenital muscular dystrophy, Ullrich type
- Dermatosparaxis Ehlers-Danlos syndrome
- Desbuquois syndrome
- Ehlers-Danlos/osteogenesis imperfecta syndrome
- High bone mass osteogenesis imperfecta
- Kyphoscoliotic Ehlers-Danlos syndrome due to FKBP22 deficiency
- Kyphoscoliotic Ehlers-Danlos syndrome due to lysyl hydroxylase 1 deficiency
- Larsen-like syndrome, B3GAT3 type
- Musculocontractural Ehlers-Danlos syndrome
- Myopathic Ehlers-Danlos syndrome
- Periodontal Ehlers-Danlos syndrome
- RIN2 syndrome
- Reunion Island Larsen-like syndrome
- SLC39A13-related spondylodysplastic Ehlers-Danlos syndrome
- Vascular Ehlers-Danlos syndrome

- XYLT1-CDG

Related Analytes

GENE	% OF CODING SEQUENCE SUFFICIENTLY COVERED TO DETECT HETEROZYGOUS MUTATIONS	COPY NUMBER VARIATION	COMMENTS
ADAMTS2	100.00	1	
AEBP1	100.00	1	
B3GALT6	100.00	1	
B3GAT3	100.00	1	
B4GALT7	100.00	1	
C1R	100.00	1	
C1S	100.00	1	
CHST14	100.00	1	
COL12A1	100.00	1	
COL1A1	100.00	1	
COL1A2	100.00	1	
COL3A1	100.00	1	
COL5A1	100.00	1	
COL5A2	100.00	1	
DSE	100.00	1	
FKBP14	100.00	1	
FLNA	100.00	1	
FLNB	100.00	1	

GENE	% OF CODING SEQUENCE SUFFICIENTLY COVERED TO DETECT HETEROZYGOUS MUTATIONS	COPY NUMBER VARIATION	COMMENTS
PLOD1	100.00	1	
PRDM5	100.00	1	
RIN2	100.00	1	
SLC39A13	100.00	1	
TAB2	100.00	1	
XYLT1	100.00	1	
XYLT2	100.00	1	
ZNF469	100.00	1	