DISEASE:
Arthrochalasia Ehlers-Danlos syndrome

NAME:	Arthrochalasia Ehlers-Danlos syndrome
DESCRIPTION:	A form of Ehlers-Danlos syndrome (EDS) characterized by congenital bilateral hip dislocation, severe generalized joint hypermobility with recurrent joint dislocations and subluxations, hyperextensible and/or fragile skin.
ORPHACODE:	1899
SYNONYMS:	Arthrochalasia EDS Arthrochalasis multiplex congenita EDS VII Ehlers-Danlos syndrome type 7 Ehlers-Danlos syndrome, arthrochalasia type aEDS
XREF(S):	Orphanet OMIM ICD-10 OMIM
ANALYTE(S):	COL1A2 COL1A1
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Related Laboratories

- Centrum Medische Genetica - UZ Gent

Related Analytes

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

Related Gene Panels

- Ehlers Danlos / arthrochalasis type VIIA –VIIB (ex 5-8) (2 genes) - UGent
- Ehlers-Danlos syndrome -UGent