DISEASE:
Autosomal dominant Robinow syndrome

NAME:	Autosomal dominant Robinow syndrome
DESCRIPTION:	The more common type of Robinow syndrome (RS) characterized by mild to moderate limb shortening and abnormalities of the head, face and external genitalia.
ORPHACODE:	3107
XREF(S):	Orphanet OMIM OMIM ICD-10 OMIM
ANALYTE(S):	DVL3 WNT5A DVL1 FZD2
CREATED:	13 May 2019 - 01:02
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RELATED CONTENT

Related Genetic Tests

- Short Stature (gene panel)
- cleft lip with/whitout cleft palate (virtual gene panel)

Related Laboratories

- Centre de Génétique Médicale UCL
- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)

Related Analytes

- dishevelled segment polarity protein 1
- dishevelled segment polarity protein 3
- frizzled class receptor 2
- Wnt family member 5A

Related Gene Panels

- Cleft lip and palate / dysmorphic facial features / craniofacial anomalies (255 genes)) - UCL
- Short Stature (46 genes) - IPG