DISEASE:
Léri-Weill dyschondrosteosis

NAME:	Léri-Weill dyschondrosteosis
DESCRIPTION:	A rare, genetic skeletal dysplasia marked by disproportionate short stature and the characteristic Madelung wrist deformity.
ORPHACODE:	240
SYNONYMS:	Léri-Weill syndrome
XREF(S):	Orphanet ICD-10 OMIM MeSH
ANALYTE(S):	SHOX
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

RELATED CONTENT

Related Genetic Tests

- Leri-Weill dyschondrosteosis / ISS
- Leri-Weill dyschondrosteosis / SHOX-related short stature
- Leri-Weill dyschondrosteosis / SHOX-related short stature
- Leri-Weill dyschondrosteosis / SHOX-related short stature
- Short Stature (gene panel)
- Short stature/ Growth retardation/ (gene panel)

Related Laboratories

- Centre de Génétique Humaine - CHU Sart-Tilman
- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)
- Centrum Medische Genetica - UZ Antwerpen
- Centrum Medische Genetica - UZ Gent

Related Analytes

- SHOX homeobox

Related Gene Panels

- Growth retardation/short stature (genepanel) - UZA
- Short Stature (46 genes) - IPG
