DISEASE:
Laron syndrome

NAME:	Laron syndrome
DESCRIPTION:	Laron syndrome is a congenital disorder characterized by marked short stature associated with normal or high serum growth hormone (GH) and low serum insulin-like growth factor-1 (IGF-I) levels which fail to rise after exogenous GH administration.
ORPHACODE:	633
SYNONYMS:	Complete growth hormone insensitivity GH receptor deficiency Growth hormone receptor deficiency Laron-type dwarfism Primary GH insensitivity Primary GH resistance Primary growth hormone insensitivity Primary growth hormone resistance Short stature due to growth hormone resistance
XREF(S):	Orphanet OMIM MeSH ICD-10

ANALYTE(S):	GHR
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RELATED CONTENT

Related Genetic Tests

- Short Stature (gene panel)
- Short stature/ Growth retardation/ (gene panel)

Related Laboratories

- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)
- Centrum Medische Genetica - UZ Antwerpen

Related Analytes

- growth hormone receptor

Related Gene Panels

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