Laron syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Laron syndrome. ORPHA:633

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.