

DISEASE:
Short stature due to GHSR deficiency

NAME:	Short stature due to GHSR deficiency
DESCRIPTION:	Short stature due to GHSR deficiency is a rare, genetic, endocrine growth disease, resulting from growth hormone secretagogue receptor (GHSR) deficiency, characterized by postnatal growth delay that results in short stature (less than -2 SD). The pituitary gland is typically without morphological changes, although anterior pituitary gland hypoplasia has been reported.
ORPHACODE:	314811
SYNOMYS:	Ghrelin receptor deficiency Short stature due to growth hormone secretagogue receptor deficiency
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	GHSR
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RELATED CONTENT

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- Short Stature (gene panel)

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

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- growth hormone secretagogue receptor

Related Gene Panels

- Short Stature (46 genes) - IPG

Source URL: <http://gentest.healthdata.be/disease/1780>