DISEASE:
Pallister-Hall syndrome

NAME:	Pallister-Hall syndrome
DESCRIPTION:	Pallister-Hall syndrome (PHS), a pleiotropic autosomal dominant malformative disorder, is characterized by hypothalamic hamartoma, pituitary dysfunction, bifid epiglottis, polydactyly, and, more rarely, renal abnormalities and genitourinary malformations.
ORPHACODE:	672
SYNONYMS:	Hypothalamic hamartoblastoma syndrome
XREF(S):	Orphanet OMIM MeSH ICD-10
ANALYTE(S):	GLI3
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RELATED CONTENT

Related Genetic Tests

- Renal or urinary tract malformation (CAKUT) (gene panel)

Related Laboratories

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

Related Analytes

- GLI family zinc finger 3

Related Gene Panels

- Cakut (congenital anomalies of the kidney and urinary tract-1) (69 genes) - IPG