DISEASE:
Familial peripheral male-limited precocious puberty

NAME:	Familial peripheral male-limited precocious puberty
DESCRIPTION:	Familial male limited precocious puberty (FMPP) is a gonadotropin-independent familial form of male-limited precocious puberty, generally presenting between 2-5 years of age as accelerated growth, early development of secondary sexual characteristics and reduced adult height.
ORPHACODE:	3000
SYNONYMS:	FMPP Familial gonadotropin-independent male-limited sexual precocity Male-limited precocious puberty Testotoxicosis
XREF(S):	Orphanet MedDRA MedDRA ICD-10 OMIM MeSH
ANALYTE(S):	LHCGR
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- Premature Ovarian Failure/Primary Ovarian Insufficiency (POF/POI) (32 genes)

Related Laboratories

- Centre de Génétique Humaine - Erasme ULB
- Centrum Medische Genetica - UZ Brussel VUB

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- luteinizing hormone/choriogonadotropin receptor

Related Gene Panels

- Premature Ovarian Failure/Insufficiency (32 genes) - VUB