DISEASE:
Microcephalic osteodysplastic primordial dwarfism type II

NAME:	Microcephalic osteodysplastic primordial dwarfism type II
DESCRIPTION:	A rare bone disease and a form of microcephalic primordial dwarfism characterized by severe pre- and postnatal growth retardation, with marked microcephaly in proportion to body size, skeletal dysplasia, abnormal dentition, insulin resistance, and increased risk for cerebrovascular disease.
ORPHACODE:	2637
SYNONYMS:	MOPD type II Majewski osteodysplastic primordial dwarfism type II
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	PCNT
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RELATED CONTENT

Related Genetic Tests

- Brain malformations (gene panel)
- Short stature/ Growth retardation/ (gene panel)

Related Laboratories

- Centre de Génétique Humaine - Erasme ULB
- Centrum Medische Genetica - UZ Antwerpen

Related Analytes

- pericentrin

Related Gene Panels

- Brain malformations (34 genes) - ULB
- Growth retardation/short stature (genepanel) - UZA

Source URL: http://gentest.healthdata.be/disease/2070