DISEASE:
3M syndrome

NAME:	3M syndrome
DESCRIPTION:	A rare primordial growth disorder characterized by low birth weight, reduced birth length, severe postnatal growth restriction, large head size, a spectrum of minor anomalies (including facial dysmorphism) and normal intelligence.
ORPHACODE:	2616
SYNONYMS:	3-M syndrome Yakut short stature syndrome
XREF(S):	Orphanet OMIM OMIM OMIM ICD-10
ANALYTE(S):	CUL7 OBSL1 CCDC8
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

RELATED CONTENT

Related Genetic Tests

- Short Stature (gene panel)
- Short stature/ Growth retardation/ (gene panel)

Related Laboratories

- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)
- Centrum Medische Genetica - UZ Antwerpen

Related Analytes

- coiled-coil domain containing 8
- cullin 7
- obscurin like cytoskeletal adaptor 1

Related Gene Panels

- Growth retardation/short stature (genepanel) - UZA
- Short Stature (46 genes) - IPG