DISEASE:
Tatton-Brown-Rahman syndrome

NAME:	Tatton-Brown-Rahman syndrome
DESCRIPTION:	A rare multiple congenital anomalies syndrome characterized by tall stature due to postnatal overgrowth, mild to moderate intellectual disability and subtle distinctive facial features, which often become apparent during adolescence, such as round face, low-set, thick horizontal eyebrows, narrow palpebral fissures and prominent upper-central incisors. Joint hypermobility, hypotonia and scoliosis are common.
ORPHACODE:	404443
SYNONYMS:	DNMT3A-related overgrowth syndrome Tatton-Brown-Rahman overgrowth syndrome
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	DNMT3A
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