DISEASE:
Congenital muscular dystrophy, Fukuyama type

NAME:	Congenital muscular dystrophy, Fukuyama type
DESCRIPTION:	A rare congenital progressive muscular dystrophy often characterized by brain malformation (cobblestone lissencephaly), dystrophic changes in skeletal muscle, severe intellectual deficit, epilepsy and motor impairment.
ORPHACODE:	272
SYNONYMS:	FCMD FKTN-related congenital muscular dystrophy Fukuyama congenital muscular dystrophy
XREF(S):	Orphanet OMIM ICD-10
ANALYTE(S):	FKTN
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- Fukuyama congenital muscular dystrophy
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Related Laboratories

- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)
- Centrum Medische Genetica - UZ Brussel VUB
- Centrum Menselijke Erfelijkheid - KUL

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- fukutin

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