DISEASE:
KDM5C-related syndromic X-linked intellectual disability

NAME:	KDM5C-related syndromic X-linked intellectual disability
DESCRIPTION:	A rare multiple congenital anomalies/dysmorphic syndrome characterized by mild to severe intellectual deficit associated with variable clinical manifestations including spasticity, cryptorchidism, maxillary hypoplasia, alopecia areata, epilepsy, short stature, impaired speech, and behavioral problems.
ORPHACODE:	85279
XREF(S):	Orphanet OMIM ICD-10
ANALYTE(S):	KDM5C
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

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