DISEASE:
Hypomyelination-congenital cataract syndrome

NAME:	Hypomyelination-congenital cataract syndrome
DESCRIPTION:	A rare developmental defect characterized by the onset of cataract either at birth or in the first two months of life, delayed psychomotor development by the end of the first year of life and moderate intellectual deficit.
ORPHACODE:	85163
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	HYCC1
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