DISEASE:
Familial or sporadic hemiplegic migraine

NAME:	Familial or sporadic hemiplegic migraine
DESCRIPTION:	A rare variety of migraine with aura characterized by the presence of a motor weakness during the aura. There are two main forms depending on the familial history: patients with at least one first- or second-degree relative who has aura including motor weakness have familial hemiplegic migraine (FHM); patients without such familial history have sporadic hemiplegic migraine (SHM).
ORPHACODE:	569
XREF(S):	Orphanet ICD-10 OMIM OMIM OMIM OMIM
ANALYTE(S):	SCN1A CACNA1A ATP1A2 PRRT2
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