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Full name:	Familial hemiplegic Migraine (8 genes) - KUL
Abbreviation:	FHM
Laboratory:	Centrum Menselijke Erfelijkheid - KUL
Created:	28 Jun 2019 - 15:02
Changed:	09 Sep 2022 - 11:16

Related Diseases

- Familial or sporadic hemiplegic migraine

Related Analytes

GENE	% OF CODING SEQUENCE SUFFICIENTLY COVERED TO DETECT HETEROZYGOUS MUTATIONS	COPY NUMBER VARIATION	COMMENTS
ATP1A2	100.00	1	NM000702.3/ Core gene
ATP1A3	100.00	1	NM152296.5/ gene that may be associated with an inherited form of Familia Hemiplegic Migraine (FHM)
CACNA1A	100.00	1	NM001127221.1/ Core gene
KCNK18	100.00	1	NM181840.1/ gene that may be associated with an inherited form of Familia Hemiplegic Migraine (FHM)
PRRT2	100.00	1	NM145239.2/ gene that may be associated with an inherited form of Familia Hemiplegic Migraine (FHM)
SCN1A	100.00	1	NM_001165963.2/ Core gene
SLC1A3	100.00	1	NM004172.4/ gene that may be associated with an inherited form of Familia Hemiplegic Migraine (FHM)

GENE	% OF CODING SEQUENCE SUFFICIENTLY COVERED TO DETECT HETEROZYGOUS MUTATIONS	COPY NUMBER VARIATION	COMMENTS
SLC2A1	100.00	1	NM_006516.2/ gene that may be associated with an inherited form of Familia Hemiplegic Migraine (FHM)