DISEASE:
Dravet syndrome

NAME:	Dravet syndrome
DESCRIPTION:	A rare, genetic, developmental and epileptic encephalopathy characterized by infantile onset of intractable seizures that are often febrile, and associated with cognitive and motor impairment.
ORPHACODE:	33069
SYNONYMS:	SMEI Severe myoclonic epilepsy of infancy Severe myoclonus epilepsy of infancy
XREF(S):	Orphanet OMIM OMIM OMIM ICD-10

ANALYTE(S):	SCN1A SCN1B SCN9A GABRA1 GABRG2 PCDH19 SCN2A
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RELATED CONTENT

Related Genetic Tests

- Epilepsy (gene panel)
- Neuropathy (gene panel)

Related Laboratories

- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)
- Centrum Medische Genetica - UZ Antwerpen

Related Analytes

- gamma-aminobutyric acid type A receptor subunit alpha1
- gamma-aminobutyric acid type A receptor subunit gamma2
- protocadherin 19
- sodium voltage-gated channel alpha subunit 1
- sodium voltage-gated channel beta subunit 1
- sodium voltage-gated channel alpha subunit 2
- sodium voltage-gated channel alpha subunit 9

Related Gene Panels

- Neuropathy (148 genes) - IPG
- Rare epilepsy with developmental delay (> 240 genes) - UZA
