DISEASE:
Postsynaptic congenital myasthenic syndromes

NAME:	Postsynaptic congenital myasthenic syndromes
ORPHACODE:	98913

XREF(S):	Orphanet OMIM OMIM OMIM OMIM OMIM OMIM OMIM OMIM OMIM OMIM ICD-10 OMIM OMIM OMIM OMIM OMIM OMIM OMIM OMIM

ANALYTE(S):	RAPSN SCN4A CHRNA1 CHRNA1 CHRNB1 CHRNB1 CHRND CHRND CHRNE CHRNE DOK7 MUSK LRP4 AGRN AK9 COL13A1
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RELATED CONTENT

Related Genetic Tests

- Neuromuscular disorders : congenital & distal myopathy, congenital muscle dystrophy / Limb-girdle muscular dystrophy / Rhabdomyolysis / Myopathy (with prominent contractures) / distal artrogryposis (gene panel)

Related Laboratories

- Centrum Medische Genetica - UZ Brussel VUB

Related Analytes

- agrin
- adenylate kinase 9
- cholinergic receptor nicotinic alpha 1 subunit
- cholinergic receptor nicotinic beta 1 subunit
- cholinergic receptor nicotinic delta subunit
- cholinergic receptor nicotinic epsilon subunit
- collagen type XIII alpha 1 chain
- docking protein 7
- LDL receptor related protein 4
- muscle associated receptor tyrosine kinase
- receptor associated protein of the synapse
- sodium voltage-gated channel alpha subunit 4

Related Gene Panels

- Neuromuscular disorders (166 genes) - VUB

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