DISEASE:
Rolandic epilepsy-speech dyspraxia syndrome

NAME:	Rolandic epilepsy-speech dyspraxia syndrome
DESCRIPTION:	Rolandic epilepsy-speech dyspraxia syndrome is a rare, genetic epilepsy characterized by speech disorder (including a range of symptoms from dysarthria, speech dyspraxia, receptive and expressive language delay/regression and acquired aphasia to subtle impairments of conversational speech) and epilepsy (mostly focal and secondary generalized childhood-onset seizures, sometimes with aura). Mild to severe intellectual disability may also be observed.
ORPHACODE:	163721
XREF(S):	Orphanet ICD-10 OMIM OMIM
ANALYTE(S):	SRPX2 GRIN2A
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RELATED CONTENT

Related Genetic Tests

- Epilepsy (gene panel)
- Epilepsy gene panel

Related Laboratories

- Centrum Medische Genetica - UZ Antwerpen
- Centrum Medische Genetica - UZ Brussel VUB

Related Analytes

- glutamate ionotropic receptor NMDA type subunit 2A
- sushi repeat containing protein X-linked 2

Related Gene Panels

- Epilepsy gene panel - VUB
- Rare epilepsy with developmental delay (> 240 genes) - UZA