

DISEASE:**Rolandic epilepsy-paroxysmal exercise-induced dystonia-writer's cramp syndrome**

NAME:	Rolandic epilepsy-paroxysmal exercise-induced dystonia-writer's cramp syndrome
DESCRIPTION:	A rare genetic epilepsy syndrome characterized by infantile or childhood onset of focal motor seizures remitting with age, as well as childhood onset of exercise-induced dystonia which often persists into adulthood. Additional reported features include nystagmus and postural tremor of the hands.
ORPHACODE:	163727
SYNONYMS:	Rolandic epilepsy exercise-induced dystonia
XREF(S):	<u>Orphanet</u> <u>MeSH</u> <u>OMIM</u> <u>ICD-10</u>
ANALYTE(S):	<u>TBC1D24</u>
CREATED:	04 Feb 2020 - 15:13
CHANGED:	22 Jun 2023 - 16:14

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