

**DISEASE:**  
**Rolandic epilepsy**

|                     |                                                                                                                                                                                                                                                                                      |
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| <b>NAME:</b>        | Rolandic epilepsy                                                                                                                                                                                                                                                                    |
| <b>DESCRIPTION:</b> | Rolandic epilepsy (RE) is a focal childhood epilepsy characterized by seizures consisting of unilateral facial sensory-motor symptoms, with electroencephalogram (EEG) showing sharp biphasic waves over the rolandic region. It is an age-related epilepsy, with excellent outcome. |
| <b>ORPHACODE:</b>   | 1945                                                                                                                                                                                                                                                                                 |
| <b>SYNONYMS:</b>    | <p>BECRS<br/> BECTS<br/> BRE<br/> Benign epilepsy of childhood with centrotemporal spikes<br/> Benign familial epilepsy of childhood with rolandic spikes<br/> Benign rolandic epilepsy<br/> Centrotemporal epilepsy</p>                                                             |
| <b>XREF(S):</b>     | <p><u><a href="#">Orphanet</a></u><br/> <u><a href="#">ICD-10</a></u><br/> <u><a href="#">OMIM</a></u><br/> <u><a href="#">OMIM</a></u></p>                                                                                                                                          |

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|--------------------|------------------------------------------------|
| <b>ANALYTE(S):</b> | <u>GABRG2</u><br><u>GRIN2A</u><br><u>SRPX2</u> |
| <b>CREATED:</b>    | 13 May 2019 - 01:02                            |
| <b>CHANGED:</b>    | 22 Jun 2023 - 16:14                            |

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- [Epilepsy \(gene panel\)](#)

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- [Centrum Medische Genetica - UZ Antwerpen](#)

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- [glutamate ionotropic receptor NMDA type subunit 2A](#)
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