

**DISEASE:**  
**Rippling muscle disease**

<b>NAME:</b>	Rippling muscle disease
<b>DESCRIPTION:</b>	Rippling muscle disease is a rare, genetic, neuromuscular disorder characterized by muscle hyperirritability triggered by stretch, percussion or movement. Patients present wave-like, electrically-silent muscle contractions (rippling), muscle mounding, painful muscle stiffness and muscle hypertrophy, usually with elevated serum creatine kinase.
<b>ORPHACODE:</b>	97238
<b>XREF(S):</b>	<a href="#">Orphanet</a> <a href="#">MeSH</a> <a href="#">MedDRA</a> <a href="#">ICD-10</a> <a href="#">OMIM</a> <a href="#">OMIM</a>
<b>ANALYTE(S):</b>	CAV3
<b>CREATED:</b>	13 May 2019 - 01:02
<b>CHANGED:</b>	22 Jun 2023 - 16:14

## RELATED CONTENT

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### Related Genetic Tests

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

### Related Laboratories

- Centrum Medische Genetica - UZ Brussel VUB

### Related Analytes

- caveolin 3

### Related Gene Panels

- Neuromuscular disorders (166 genes) - VUB

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Source URL: <http://gentest.healthdata.be/disease/3432>