

**DISEASE:**  
**Paroxysmal nocturnal hemoglobinuria**

<b>NAME:</b>	Paroxysmal nocturnal hemoglobinuria
<b>DESCRIPTION:</b>	Paroxysmal nocturnal hemoglobinuria (PNH) is an acquired clonal hematopoietic stem cell disorder characterized by corpuscular hemolytic anemia, bone marrow failure and frequent thrombotic events.
<b>ORPHACODE:</b>	447
<b>SYNONYMS:</b>	Marchiafava-Micheli disease PNH
<b>XREF(S):</b>	<a href="#">Orphanet</a> <a href="#">OMIM</a> <a href="#">OMIM</a> <a href="#">MeSH</a> <a href="#">MedDRA</a> <a href="#">ICD-10</a>
<b>ANALYTE(S):</b>	<a href="#">PIGA</a>
<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

- Trombosis - Hemostasis (gene panel)

### Related Laboratories

- Centrum Menselijke Erfelijheid - KUL

### Related Analytes

- phosphatidylinositol glycan anchor biosynthesis class A

### Related Gene Panels

- Trombosis - Hemostasis (107 genes) - KUL

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Source URL: <http://gentest.healthdata.be/index.php/index.php/disease/63>