

**DISEASE:**  
**Ichthyosis hystrix of Curth-Macklin**

<b>NAME:</b>	Ichthyosis hystrix of Curth-Macklin
<b>DESCRIPTION:</b>	Ichthyosis hystrix of Curth-Macklin (IHCM) is a rare type of keratinopathic ichthyosis (see this term) that is characterized by the presence of severe hyperkeratotic lesions and palmoplantar keratoderma (PPK, see this term).
<b>ORPHACODE:</b>	79503
<b>SYNOMYS:</b>	Ichthyosis hystrix, Curth-Macklin type
<b>XREF(S):</b>	<a href="#">Orphanet</a> <a href="#">ICD-10</a> <a href="#">OMIM</a> <a href="#">MeSH</a>
<b>ANALYTE(S):</b>	<a href="#">KRT1</a>
<b>CREATED:</b>	13 May 2019 - 01:02
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## RELATED CONTENT

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### Related Genetic Tests

- Ichthyosis (gene panel)

### Related Laboratories

- Centrum Menselijke Erfelijkheid - KUL

### Related Analytes

- keratin 1

### Related Gene Panels

- Ichthyosis and erythroderma (98 genes) - KUL

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Source URL: <http://gentest.healthdata.be/index.php/index.php/disease/3365>