

**DISEASE:**

**Isolated focal non-epidermolytic palmoplantar keratoderma**

<b>NAME:</b>	Isolated focal non-epidermolytic palmoplantar keratoderma
<b>DESCRIPTION:</b>	A rare hereditary palmoplantar keratoderma characterized by focal hyperkeratotic lesions on the palms and soles. Histopathologic examination reveals prominent hyperkeratosis, thickened stratum spinosum with reduced stratum granulosum, disadhesion of cells in the suprabasal layers, elongation of rete ridges, and sparse lymphocyte infiltration in the dermis.
<b>ORPHACODE:</b>	448264
<b>XREF(S):</b>	<a href="#">Orphanet</a> <a href="#">ICD-10</a> <a href="#">OMIM</a> <a href="#">OMIM</a>
<b>ANALYTE(S):</b>	<a href="#">TRPV3</a> <a href="#">KRT16</a>
<b>CREATED:</b>	13 May 2019 - 01:02
<b>CHANGED:</b>	22 Jun 2023 - 16:14

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Source URL: <http://gentest.healthdata.be/disease/2173>

## RELATED CONTENT

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### Related Genetic Tests

- Ichthyosis (gene panel)

### Related Laboratories

- Centrum Menselijke Erfelijkheid - KUL

### Related Analytes

- keratin 16
- transient receptor potential cation channel subfamily V member 3

### Related Gene Panels

- Ichthyosis and erythroderma (98 genes) - KUL

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