

**DISEASE:**  
**Naxos disease**

<b>NAME:</b>	Naxos disease
<b>DESCRIPTION:</b>	A recessively inherited condition with arrhythmogenic right ventricular dysplasia/cardiomyopathy (ARVD/C) and a cutaneous phenotype, characterised by peculiar woolly hair and palmoplantar keratoderma.
<b>ORPHACODE:</b>	34217
<b>SYNONYMS:</b>	KWWH type I Keratoderma with woolly hair type I Keratosis palmoplantaris with arrythmogenic cardiomyopathy Palmoplantar hyperkeratosis with arrythmogenic cardiomyopathy Palmoplantar keratoderma with arrythmogenic cardiomyopathy
<b>XREF(S):</b>	<a href="#">Orphanet</a> <a href="#">MeSH</a> <a href="#">OMIM</a> <a href="#">ICD-10</a>
<b>ANALYTE(S):</b>	<a href="#">JUP</a>
<b>CREATED:</b>	13 May 2019 - 01:02
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### Related Genetic Tests

- [Cardiomyopathy, hereditary \(gene panel\)](#)
- [Cardiopathies, hereditary \(gene panel\)](#)

### Related Laboratories

- [Centrum Medische Genetica - UZ Brussel VUB](#)
- [Centrum Menselijke Erfelijkheid - KUL](#)

### Related Analytes

- [junction plakoglobin](#)

### Related Gene Panels

- [Cardiomyopathy, hereditary \(208 genes\) - VUB](#)
- [Cardiopathies, hereditary \(102 genes\) - KUL](#)

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