

**DISEASE:**

**Heterozygous familial hypercholesterolemia (NON RARE IN EUROPE)**

<b>NAME:</b>	Heterozygous familial hypercholesterolemia (NON RARE IN EUROPE)
<b>ORPHACODE:</b>	406
<b>XREF(S):</b>	<a href="#">ORPHANET</a>
<b>CREATED:</b>	24 Sep 2020 - 17:11
<b>CHANGED:</b>	24 Sep 2020 - 17:14

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## RELATED CONTENT

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### Related Genetic Tests

- [Hypercholesterolemia, Familial \(9 genes\)](#)
- [Hypercholesterolemia, Familial \(gene panel\)](#)

### Related Laboratories

- [Centre de Génétique Humaine - CHU Sart-Tilman](#)
- [Centrum Medische Genetica - UZ Antwerpen](#)

### Related Gene Panels

- [Familial Hypercholesterolemia panel \(8 genes\) - UZA](#)
- [Familial Hypercholesterolemia panel \(9 genes\) - ULG](#)

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