

**DISEASE:**

Atrial septal defect, ostium secundum type

<b>NAME:</b>	Atrial septal defect, ostium secundum type
<b>ORPHACODE:</b>	99103
<b>SYNOMYS:</b>	ASD, ostium secundum type
<b>XREF(S):</b>	<a href="#">Orphanet</a> <a href="#">OMIM</a> <a href="#">OMIM</a> <a href="#">MedDRA</a> <a href="#">MedDRA</a> <a href="#">ICD-10</a> <a href="#">OMIM</a>

<b>ANALYTE(S):</b>	ACTC1 <u>GATA4</u> MYH6 <u>NKX2-5</u> TBX20 <u>TBX20</u> TLL1 <u>CITED2</u> GATA6
<b>CREATED:</b>	13 May 2019 - 01:02
<b>CHANGED:</b>	22 Jun 2023 - 16:14

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## RELATED CONTENT

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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

- [actin alpha cardiac muscle 1](#)
- [Cbp/p300 interacting transactivator with Glu/Asp rich carboxy-terminal domain 2](#)
- [GATA binding protein 4](#)
- [GATA binding protein 6](#)
- [myosin heavy chain 6](#)
- [NK2 homeobox 5](#)
- [T-box transcription factor 20](#)
- [tolloid like 1](#)

### Related Gene Panels

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

- Cardiopathies, hereditary (102 genes) - KUL
- Congenital heart disease (29 genes) - VUB

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