# Belgian Genetic Tests database

## ANALYTE:
## ACTC1

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|---|---|
| **NAME:** | actin alpha cardiac muscle 1 |
| **SYMBOL:** | ACTC1 |
| **VERSION OF ORPHANET:** | 2023-06-22 14:14:43 |
| **SYNONYMS:** | CMD1R |
| **XREF(S):** | Orphanet<br>Ensembl<br>Genatlas<br>HGNC<br>OMIM<br>Reactome<br>SwissProt |
| **CREATED:** | 13 May 2019 - 01:01 |
| **CHANGED:** | 22 Jun 2023 - 16:14 |

Source URL: http://gentest.healthdata.be/index.php/index.php/analyte/808

# RELATED CONTENT

## Related Genetic Tests

- Cardiomyopathy, hereditary (gene panel)
- Cardiomyopathy, hypertrophic
- Cardiomyopathy: hypertrophic cardiomyopathy, dilated cardiomyopathy, restrictive cardiomyopathy, left ventricular non-compaction cardiomyopathy, arrhythmogenic right ventricular cardiomyopathy (gene panel)
- Cardiopathies, hereditary (gene panel)
- Congenital malformation (gene panel - 1721 genes)
- Congenital structural heart defects (gene panel)
- Dilated Cardiomyopathy (Gene panel)
- Dilated cardiomyopathy
- Heart / Cardio disorders / Cardiopathy (gene panel)
- Hypertrophic cardiomyopathy (gene panel)
- Primary ciliary dyskinesia (PCD) Heterotaxyies (gene panel)

## Related Diseases

- Atrial septal defect, ostium secundum type
- Familial isolated dilated cardiomyopathy
- Left ventricular noncompaction
- NON RARE IN EUROPE: Familial isolated hypertrophic cardiomyopathy

## Related Gene Panels

- Cardiomyopathy (genepanel) - UZA
- Cardiomyopathy, hereditary (208 genes) - VUB

- Cardiopathies, hereditary (102 genes) - KUL
- Congenital malformation (1721 genes) - ULB
- Congenital structural heart defects - UGent
- Dilated Cardiomyopathy (79 genes) - IPG
- Dilated cardiomyopathy - UGent
- Heterotaxie PCD - UGent
- Hypertrophic cardiomyopathy (75 genes) - IPG
- Hypertrophic cardiomyopathy - UGent
- cardiopathy panel - UGent

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