# Belgian Genetic Tests database

## ANALYTE:
## GLA

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|---|---|
| **NAME:** | galactosidase alpha |
| **SYMBOL:** | GLA |
| **VERSION OF ORPHANET:** | 2023-06-22 14:14:43 |
| **SYNONYMS:** | GALA |
| **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/203

# RELATED CONTENT

## Related Genetic Tests

- Cardiomyopathy, hereditary (gene panel)
- Cardiomyopathy: hypertrophic cardiomyopathy, dilated cardiomyopathy, restrictive cardiomyopathy, left ventricular non-compaction cardiomyopathy, arrhythmogenic right ventricular cardiomyopathy (gene panel)
- Cardiopathies, hereditary (gene panel)
- Charcot-Marie-Tooth (other than type 1A) (gene panel, IPN panel)
- Congenital malformation (gene panel - 1721 genes)
- Dermatogenetic panel, severe, rare and hereditary genodermatoses (gene panel - 394 genes)
- Dilated Cardiomyopathy (Gene panel)
- Early onset epileptic encephalopathy (gene panel - 845 genes)
- Epilepsy gene panel
- Fabry disease
- Fabry disease
- Heart / Cardio disorders / Cardiopathy (gene panel)
- Hypertrophic cardiomyopathy (gene panel)
- Inherited Kidney Diseases (Gene Panel)
- Leukodystrophy (gene panel)
- Lysosomal Storage Disease (gene panel)
- Nephrogenetics / Nephropathy (gene panel)
- Nephropathies, hereditary (gene panel)
- Nephrotic syndrome, Focal Segmental Glomerulosclerosis (FSGS) , Alport syndrome and podocytopathy (gene panel)
- Neuromuscular disorders (548 genes)
- Neuropathy (gene panel)
- Neuropathy (gene panel)
- Peripheral neuropathy (gene panel)
- Stroke (gene panel)

## Related Diseases

- Fabry disease

## Related Gene Panels

- Inherited Peripheral Neuropathies gene panel (139 genes) - KUL
- Cardiomyopathy (genepanel) - UZA
- Cardiomyopathy, hereditary (208 genes) - VUB
- Cardiopathies, hereditary (102 genes) - KUL
- Congenital malformation (1721 genes) - ULB
- Dermatogenetic / severe, rare and hereditary genodermatoses (394 genes) - ULB
- Dilated Cardiomyopathy (79 genes) - IPG
- Early onset epileptic encephalopathy (845 genes) - ULB
- Epilepsy gene panel - VUB
- Hypertrophic cardiomyopathy (75 genes) - IPG
- Leukodystrophy - UGent
- Lysomal Storage (64 genes) - VUB
- Nephropathies, hereditary (219 genes) - KUL
- Nephropathy panel - UGent
- Nephrotic syndrome, FSGS, Alport syndrome (76 genes) - IPG
- Neuromuscular disorders (548 genes) - ULB
- Neuropathy (148 genes) - IPG
- Neuropathy (genepanel) - UZA
- Neuropathy panel - UGent
- Panel Nephro-ULG-V1
- Stroke - UGent
- cardiopathy panel - UGent

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