# Belgian Genetic Tests database

## ANALYTE:
## MAN2B1

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|---|---|
| **NAME:** | mannosidase alpha class 2B member 1 |
| **SYMBOL:** | MAN2B1 |
| **VERSION OF ORPHANET:** | 2023-06-22 14:14:43 |
| **SYNONYMS:** | LAMAN |
| **XREF(S):** | Orphanet<br>Reactome<br>Ensembl<br>Genatlas<br>HGNC<br>OMIM<br>SwissProt |
| **CREATED:** | 13 May 2019 - 01:01 |
| **CHANGED:** | 22 Jun 2023 - 16:14 |

Source URL: http://gentest.healthdata.be/analyte/2807

# RELATED CONTENT

## Related Genetic Tests

- Ataxia Spasticity (gene panel)
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (gene panel)
- Intellectual disability (virtual gene panel)
- Leukodystrophy (gene panel)
- Lysosomal Storage Disease (gene panel)
- Neurodevelopmental disorders (1300 genes)
- Neurodevelopmental disorders gene panel
- Primary immune deficiencies (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)

## Related Diseases

- Alpha-mannosidosis, adult form
- Alpha-mannosidosis, infantile form

## Related Gene Panels

- Ataxia Spasticity - UGent
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Intellectual disability/Epilepsy (1091 genes) - ULG

- Leukodystrophy - UGent
- Lysomal Storage (64 genes) - VUB
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- Primary immune deficiencies - UGent
- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent

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