# DISEASE:
# GM1 gangliosidosis type 3

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| **NAME:** | GM1 gangliosidosis type 3 |
| **DESCRIPTION:** | GM1 gangliosidosis type 3 is a mild, chronic, adult form of GM1 gangliosidosis (see this term) characterized by onset generally during childhood or adolescence and by cerebellar dysfunction. |
| **ORPHACODE:** | 79257 |
| **SYNONYMS:** | Adult-onset GM1 gangliosidosis |
| **XREF(S):** | Orphanet<br>OMIM<br>ICD-10 |
| **ANALYTE(S):** | GLB1 |
| **CREATED:** | 13 May 2019 - 01:02 |
| **CHANGED:** | 22 Jun 2023 - 16:14 |

# RELATED CONTENT

## Related Genetic Tests

- Lysosomal Storage Disease (gene panel)
- Mucopolysaccharidosis (MPS) type IVB, Morquio B syndrome / GM1 gangliosidosis

## Related Laboratories

- Centrum Medische Genetica - UZ Brussel VUB

## Related Analytes

- galactosidase beta 1

## Related Gene Panels

- Lysomal Storage (64 genes) - VUB