# DISEASE:
# X-linked complicated corpus callosum dysgenesis

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|---|---|
| **NAME:** | X-linked complicated corpus callosum dysgenesis |
| **DESCRIPTION:** | A congenital, X-linked, clinical subtype of L1 syndrome, characterized by variable spastic paraplegia, mild to moderate intellectual disability, and dysplasia, hypoplasia or aplasia of the corpus callosum. In this subtype hydrocephalus, adducted thumbs, or absent speech are not observed. |
| **ORPHACODE:** | 1497 |
| **XREF(S):** | Orphanet<br>OMIM<br>ICD-10 |
| **ANALYTE(S):** | L1CAM |
| **CREATED:** | 13 May 2019 - 01:02 |
| **CHANGED:** | 22 Jun 2023 - 16:14 |

Source URL: http://gentest.healthdata.be/disease/748

# RELATED CONTENT

## Related Genetic Tests

- Brain malformations (gene panel)

## Related Laboratories

- Centre de Génétique Humaine - Erasme ULB

## Related Analytes

- L1 cell adhesion molecule

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