# DISEASE:
# Microcephaly-lymphedema-chorioretinopathy syndrome

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| **NAME:** | Microcephaly-lymphedema-chorioretinopathy syndrome |
| **DESCRIPTION:** | Microcephaly with or without chorioretinopathy, lymphedema or intellectual disability (MCLID) is a rare autosomal dominant condition characterized by variable expression of microcephaly, ocular disorders including chorioretinopathy, congenital lymphedema of the lower limbs, and mild to moderate intellectual disability. |
| **ORPHACODE:** | 2526 |
| **SYNONYMS:** | MLCRD |
| **XREF(S):** | Orphanet<br>OMIM<br>ICD-10<br>MeSH |
| **ANALYTE(S):** | KIF11 |
| **CREATED:** | 13 May 2019 - 01:02 |
| **CHANGED:** | 22 Jun 2023 - 16:14 |

# RELATED CONTENT

## Related Genetic Tests

- Primary lymphedema / fetal hydrops (gene panel)

## Related Laboratories

- Centre de Génétique Médicale UCL

## Related Analytes

- kinesin family member 11