# DISEASE:
# 15q24 microdeletion syndrome

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| **NAME:** | 15q24 microdeletion syndrome |
| **DESCRIPTION:** | 15q24 microdeletion syndrome is a rare chromosomal anomaly characterized cytogenetically by a 1.7-6.1 Mb deletion in chromosome 15q24 and clinically by pre- and post-natal growth retardation, intellectual disability, distinct facial features, and genital, skeletal, and digital anomalies. |
| **ORPHACODE:** | 94065 |
| **SYNONYMS:** | Del(15)(q24)<br>Monosomy 15q24 |
| **XREF(S):** | Orphanet<br>OMIM<br>ICD-10 |
| **ANALYTE(S):** | SIN3A |
| **CREATED:** | 13 May 2019 - 01:02 |
| **CHANGED:** | 22 Jun 2023 - 16:14 |

## RELATED CONTENT

### Related Analytes

- SIN3 transcription regulator family member A

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