# DISEASE:
# 6q25 microdeletion syndrome

| | |
|---|---|
| **NAME:** | 6q25 microdeletion syndrome |
| **DESCRIPTION:** | 6q25 microdeletion syndrome is a recently described syndrome characterized by developmental delay, facial dysmorphism and hearing loss. |
| **ORPHACODE:** | 251056 |
| **SYNONYMS:** | Del(6)(q25)<br>Monosomy 6q25 |
| **XREF(S):** | Orphanet<br>OMIM<br>ICD-10 |
| **ANALYTE(S):** | ARID1B |
| **CREATED:** | 13 May 2019 - 01:02 |
| **CHANGED:** | 22 Jun 2023 - 16:14 |

## RELATED CONTENT

### Related Analytes

- AT-rich interaction domain 1B

Source URL: http://gentest.healthdata.be/disease/930