# GENETIC TEST:
# Deafness, autosomal dominant 6/14 / Wolfram syndrome

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|---|---|
| **FULL NAME:** | Deafness, autosomal dominant 6/14 / Wolfram syndrome |
| **TEST TYPE:** | Clinical |
| **TEST SPECIALTY:** | Molecular Genetics |
| **TEST PURPOSE:** | Post-natal Diagnosis |
| **SPECIMEN:** | Peripheral (whole) blood on EDTA |
| **METHOD CATEGORY:** | Sequence analysis: entire coding region |
| **METHOD TECHNIQUE:** | Next Generation Sequencing (NGS) |
| **RIZIV CODE:** | 565471-565482 |
| **ACCREDITATION (ISO 15189):** | 2023-11-09 / 2024-05-08 |
| **TURNAROUND TIME (MAXIMUM):** | Unknown |
| **CREATED:** | 21 Aug 2019 - 08:57 |
| **CHANGED:** | 22 Jan 2024 - 13:42 |

# RELATED CONTENT

## Related Diseases

- Rare autosomal dominant non-syndromic sensorineural deafness type DFNA
- Wolfram syndrome

## Related Laboratories

- Centrum Medische Genetica - UZ Antwerpen

## Related Analytes

- wolframin ER transmembrane glycoprotein