

T436= — WFS1 Molecular Atlas Card

Variant type: Synonymous (silent)

Codon: position 436 (Threonine, T) — amino acid unchanged

Domain context: Transmembrane helix 5

SCHEMA CATEGORY: SILENT — SILENT — NO AMINO-ACID CHANGE

No amino-acid change (T436 is unchanged): the codon is altered but the protein sequence is identical to wild-type. No structural, stability or AlphaMissense effect applies. Synonymous variants are typically benign unless they affect splicing or regulatory elements; this one is not adjacent to an exon boundary.

CLINICAL EVIDENCE

- **Classification:** Benign
- **Review status:** criteria provided, multiple submitters, no conflicts
- **Associated conditions:** WFS1-Related Spectrum Disorders; Wolfram syndrome 1; Autosomal dominant nonsyndromic hearing loss 6
- **cDNA change:** c.1308C>T
- **ClinVar accession:** VCV000045433
- **Last evaluated:** 2026/03/01 00:00
- **Submissions:** 1

Card generated by wolfram-atlas-batch (synonymous pipeline) on 2026-06-08T02:53:17.850840Z.

WFS1: UniProt O76024, AlphaFold v6. Synonymous variants carry no protein-structural effect.