RARERESEARCH.AI · WFS1 MOLECULAR ATLAS · SYNONYMOUS

L149= — WFS1 Molecular Atlas Card

Variant type: Synonymous (silent)

Codon: position 149 (Leucine, L) — amino acid unchanged

Domain context: N-terminal cytoplasmic (intrinsically disordered)

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

No amino-acid change (L149 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:** Conflicting classifications of pathogenicity
- **Review status:** criteria provided, conflicting classifications
- **Associated conditions:** Autosomal dominant nonsyndromic hearing loss 6; Wolfram syndrome 1; WFS1-Related Spectrum Disorders
- **cDNA change:** c.445T>C
- **ClinVar accession:** VCV000907296
- **Last evaluated:** 2018/01/13 00:00
- **Submissions:** 1

Card generated by `wolfram-atlas-batch` (synonymous pipeline) on 2026-06-08T02:51:07.764297Z.

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