RARERESEARCH.AI · WFS1 MOLECULAR ATLAS · MISSENSE

A214V — WFS1 Molecular Atlas Card

Variant type: Missense

Substitution: Alanine (A) → Valine (V) at position 214

Domain context: N-terminal cytoplasmic (intrinsically disordered)

ALPHAMISSENSE

- **Pathogenicity score:** 0.063
- **Class:** likely benign

ALPHAFOLD CONFIDENCE

- **pLDDT at residue 214:** 40.25

> **DynaMut2 ΔΔG:** not yet computed for this variant — AlphaMissense + AlphaFold

> confidence shown above. Stability ΔΔG and the wild-type/mutant structural

> comparison backfill behind this note.

CLINICAL EVIDENCE

- **Classification:** Uncertain significance
- **Review status:** criteria provided, multiple submitters, no conflicts
- **Associated conditions:** Inborn genetic diseases; Wolfram syndrome 1; Autosomal dominant nonsyndromic hearing loss 6; Type 2 diabetes mellitus; Wolfram-like syndrome; Cataract 41
- **cDNA change:** c.641C>T
- **ClinVar accession:** VCV001496004
- **Last evaluated:** 2025/10/29 00:00
- **Submissions:** 1

Card generated by `wolfram-atlas-batch` (missense AlphaMissense mint) on 2026-06-08T02:27:33.402499Z.

AlphaMissense (Cheng et al. 2023) · AlphaFold model v6 · UniProt O76024.