

A134T — WFS1 Molecular Atlas Card

Variant type: Missense

Substitution: Alanine (A) → Threonine (T) at position 134

Domain context: N-terminal cytoplasmic (intrinsically disordered)

ALPHAMISSENSE

- **Pathogenicity score:** 0.4113
- **Class:** ambiguous

ALPHAFOLD CONFIDENCE

- **pLDDT at residue 134:** 90.94

> **DynaMut2 $\Delta\Delta G$:** not yet computed for this variant — AlphaMissense + AlphaFold

> confidence shown above. Stability $\Delta\Delta G$ 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:** Monogenic diabetes; Hereditary ataxia; WFS1-Related Spectrum Disorders; Wolfram syndrome 1; Cataract 41; Autosomal dominant nonsyndromic hearing loss 6; Type 2 diabetes mellitus; Wolfram-like syndrome
- **cDNA change:** c.400G>A
- **ClinVar accession:** VCV000215410
- **Last evaluated:** 2025/10/14 00:00
- **Submissions:** 1

Card generated by `wolfram-atlas-batch` (missense AlphaMissense mint) on 2026-06-08T02:27:33.368978Z.

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