

# A134= — WFS1 Molecular Atlas Card

**Variant type:** Synonymous (silent)

**Codon:** position 134 (Alanine, A) — amino acid unchanged

**Domain context:** N-terminal cytoplasmic (intrinsically disordered)

---

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

---

No amino-acid change (A134 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/Likely 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.402G>A
- **ClinVar accession:** VCV000045458
- **Last evaluated:** 2025/12/15 00:00
- **Submissions:** 1

---

*Card generated by wolfram-atlas-batch (synonymous pipeline) on 2026-06-08T02:51:03.665836Z.*

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