

# Q687\* — WFS1 Molecular Atlas Card

**Variant type:** Nonsense (premature stop codon)

**Position:** 687

**Wild-type residue:** Glutamine (Q)

**Domain context (where the stop falls):** C-terminal ER-lumenal (calcium binding, calmodulin, chaperone)

---

## SCHEMA CATEGORY: N3 — NMD-ESCAPE, MODERATE TRUNCATION — CHAPERONE EXPLORATION

---

Truncated protein retains substantial structure but loses C-terminal domains. Worth screening generic ER chaperones (4-PBA, TUDCA) and sigma-1 receptor agonists. Lower confidence than for missense variants, but a candidate for the high-content drug screen (Initiative 8).

---

## NMD PREDICTION

---

- **Status:** NMD-escape
- **Confidence:** high
- **Reasoning:** Stop codon at position 687 is in the last exon (exon 8, starts ~aa 413). NMD does not target stop codons in the last exon — a truncated protein is produced.

---

## TRUNCATION ANALYSIS

---

- **Residues retained:** 1 – 686 (77.1% of full-length protein)
- **Residues lost:** 687 – 890 (22.9% of full-length protein)

### Retained domains

- N-terminal cytoplasmic (intrinsically disordered) (aa 1–310)

- Transmembrane helix 1 (aa 311–331)
- Cytoplasmic loop 1 (aa 332–340)
- Transmembrane helix 2 (aa 341–361)
- Luminal loop 1 (aa 362–370)
- Transmembrane helix 3 (aa 371–391)
- Cytoplasmic loop 2 (aa 392–400)
- Transmembrane helix 4 (aa 401–421)
- Luminal loop 2 (aa 422–431)
- Transmembrane helix 5 (aa 432–452)
- Cytoplasmic loop 3 (aa 453–461)
- Transmembrane helix 6 (aa 462–482)
- Luminal loop 3 (aa 483–496)
- Transmembrane helix 7 (aa 497–517)
- Cytoplasmic loop 4 (aa 518–532)
- Transmembrane helix 8 (aa 533–553)
- Luminal loop 4 (aa 554–573)
- Transmembrane helix 9 (aa 574–594)
- Cytoplasmic loop 5 / pre-luminal (aa 595–599)

### Partially retained at truncation point

- **C-terminal ER-luminal (calcium binding, calmodulin, chaperone)** — partial: aa 600–686 retained, aa 687–890 lost

### Lost domains

*(no full domains lost — only distal C-terminus)*

---

## CLINICAL EVIDENCE

---

- **Classification:** Pathogenic/Likely pathogenic
  - **Review status:** criteria provided, multiple submitters, no conflicts
  - **cDNA change:** c.2059C>T
  - **ClinVar accession:** VCV001298956
  - **Last evaluated:** 2024/04/13 00:00
  - **Submissions:** 1
-

## WHY THIS VARIANT MATTERS

---

Moderate truncation leaves some of the protein intact, including portions of the transmembrane bundle. Whether the partial protein can be coaxed into function with chaperones is an open question — the atlas surfaces it as a candidate for the Initiative 8 drug screen, with the explicit structural data needed to design that screen.

---

*Card generated by `wolfram-atlas-batch` skill (v1) on 2026-06-08T02:18:47.450239Z.*

*NMD rule and schema definitions: `reference/nmd` `rules.md`, `reference/cardschemaextension.md` .*

*WFS1 reference: UniProt O76024, AlphaFold model v6.*