MOLECULAR ATLAS · VARIANT CARD

WFS1 R383H — Wolframin

Arg→His p383 loop AM=0.08 ddg=-1.35 pLDDT=83. ClinVar Conflicting evidence. Atlas mechanism: see structural analysis.

IDENTITY

Variant	R383H (p.Arginine383Histidine)
DNA change	c.1148G>A
Gene · Protein	WFS1 · Wolframin (890 aa)
UniProt	O76024 · WFS1_HUMAN
ClinVar accession	VCV001168671
Amino acid change	partial charge reduction

STRUCTURAL CONTEXT

AlphaFold model	AF-O76024-F1, v6
pLDDT at residue 383	**82.88** HIGH CONFIDENCE
Domain	Connecting loop
Position context	Connecting loop
IDR flag	No — pLDDT well above 50 threshold

> Position analysis: PHE384 (2.4 Å), LEU382 (2.5 Å — L382P!), LEU380 (3.8 Å). Substantial ΔΔG. Adjacent to L382P/E385K cluster. The Atlas's neighbor extraction surfaces this variant's contacts and connects them to the broader multi-variant target landscape.

COMPUTATIONAL PREDICTIONS

ALPHAMISSENSE

0.075

am_class: **LBen** — threshold > 0.564

DYNAMUT2 ΔΔG

-1.35 kcal/mol

Destabilising · Job 177992519991

PLDDT (ALPHAFOLD)

82.88

high confidence

CLINICAL EVIDENCE

ClinVar classification	**CONFLICTING CLASSIFICATIONS OF PATHOGENICITY**
Review status	criteria provided, conflicting classifications
Last evaluated	2025/12/15 00:00
Inheritance	Conflicting ClinVar classifications.
WFS1 variant landscape	R383H is 1 of ~326 pathogenic-spectrum variants in WFS1 (out of 2,243 in ClinVar)

- (no specific conditions catalogued)

RESEARCH PATH DECISION TREE

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ΔΔG < 2 + binding site affected → CATEGORY 3 — docking experiments ΔΔG 2–4 → CATEGORY 2 — pharmacological chaperones ΔΔG > 4 → CATEGORY 1 — gene therapy pLDDT < 50 → CATEGORY 5 — IDR, experimental only Stable fold + functional site hit → CATEGORY 4 — site-specific docking
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Cat 3/4 — see structural prose. AlphaMissense below threshold (AM under-call class) but mechanism is structurally identified. Therapeutic strategy: site-directed at contacts identified above, or wet-lab validation if pLDDT borderline/below 50.

Same 382-385 loop cluster.
