BanglaVariant-Explain | Evidence snapshot
Research prototype | Nafisa Tasneem Esha

Exported 2026-10-17T11:09:49.970Z
Snapshot 2026-10-17
Gene: BRCA1
Variant: NM_007294.4(BRCA1):c.5266dup (p.Gln1756fs)
ClinVar accession: VCV000017677.176
Source classification: Pathogenic
Source review status: reviewed by expert panel
Source record updated: 2026-10-10
Original source: https://www.ncbi.nlm.nih.gov/clinvar/variation/17677/
South Asian and global allele frequency: unavailable in this release.
ML pathogenicity prediction / confidence / SHAP: not implemented.
Literature retrieval: TF-IDF keyword matching, not variant-level evidence verification.
This report contains no uploaded VCF data or personal genotype.
For research and education only. Not medical advice or a diagnosis.
A source classification is not a personal disease-risk probability. Check the original record and relevant professional guidance.