

# Resource Summary Report

Generated by [NIF](#) on Aug 26, 2026

## SAPRED

RRID:SCR\_010785

Type: Tool

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### Proper Citation

SAPRED (RRID:SCR\_010785)

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### Resource Information

**URL:** <http://sapred.cbi.pku.edu.cn/>

**Proper Citation:** SAPRED (RRID:SCR\_010785)

**Description:** Offers the researchers an automatic pipeline to predict the disease-association of SAPs.

**Abbreviations:** SAPRED

**Synonyms:** SAP Disease-Association Predictor

**Resource Type:** software resource

**Defining Citation:** [PMID:17384424](#)

**Resource Name:** SAPRED

**Resource ID:** SCR\_010785

**Alternate IDs:** OMICS\_00161

**Record Creation Time:** 20220129T080300+0000

**Record Last Update:** 20260815T182412+0000

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### Ratings and Alerts

No rating or validation information has been found for SAPRED.

No alerts have been found for SAPRED.

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### Data and Source Information

**Source:** [SciCrunch Registry](#)

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## Usage and Citation Metrics

We found 4 mentions in open access literature.

**Listed below are recent publications.** The full list is available at [NIF](#) .

Wang Q, et al. (2015) A review of study designs and statistical methods for genomic epidemiology studies using next generation sequencing. *Frontiers in genetics*, 6, 149.

Katsonis P, et al. (2014) Single nucleotide variations: biological impact and theoretical interpretation. *Protein science : a publication of the Protein Society*, 23(12), 1650.

Yang X, et al. (2014) ATP1A3 mutations and genotype-phenotype correlation of alternating hemiplegia of childhood in Chinese patients. *PloS one*, 9(5), e97274.

Zelinger L, et al. (2011) A missense mutation in DHDDS, encoding dehydrodolichyl diphosphate synthase, is associated with autosomal-recessive retinitis pigmentosa in Ashkenazi Jews. *American journal of human genetics*, 88(2), 207.