

Resource Summary Report

Generated by [RRID](#) on Sep 11, 2026

SNP and indel Imputability

RRID:SCR_010800

Type: Tool

Proper Citation

SNP and indel Imputability (RRID:SCR_010800)

Resource Information

URL: <http://www.unc.edu/~yunmli/1000G-imp/>

Proper Citation: SNP and indel Imputability (RRID:SCR_010800)

Description: A comprehensive SNP and indel imputability database.

Abbreviations: SNP and indel Imputability

Resource Type: data or information resource, database

Resource Name: SNP and indel Imputability

Resource ID: SCR_010800

Alternate IDs: OMICS_00280

Record Creation Time: 20220129T080300+0000

Record Last Update: 20260905T133200+0000

Ratings and Alerts

No rating or validation information has been found for SNP and indel Imputability.

No alerts have been found for SNP and indel Imputability.

Data and Source Information

Source: [SciCrunch Registry](#)

Usage and Citation Metrics

We found 1 mentions in open access literature.

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

Kanai M, et al. (2016) Empirical estimation of genome-wide significance thresholds based on the 1000 Genomes Project data set. Journal of human genetics, 61(10), 861.