Resource Summary Report

Generated by PRECISE-TBI on Aug 2, 2026

VARiD

RRID:SCR_000241
Type: Tool

Proper Citation

VARiD (RRID:SCR_000241)

Resource Information

URL: http://compbio.cs.utoronto.ca/varid/

Proper Citation: VARiD (RRID:SCR_000241)

Description: Software using a Hidden Markov Model for SNP (single nucleotide polymorphism) and indel identification with AB-SOLiD color-space as well as regular letter-space reads.

Abbreviations: VARiD

Resource Type: software resource

Defining Citation: PMID:20529926

Keywords: c, single nucleotide polymorphism, indel, bio.tools

Availability: Free, Available for download, Freely available

Resource Name: VARiD

Resource ID: SCR_000241

Alternate IDs: OMICS_02163, biotools:varid

Alternate URLs: https://bio.tools/varid

Record Creation Time: 20220129T080200+0000

Record Last Update: 20260801T190112+0000

Ratings and Alerts

No rating or validation information has been found for VARiD.

No alerts have been found for VARiD.

Data and Source Information

Source: SciCrunch Registry

Usage and Citation Metrics

We have not found any literature mentions for this resource.