Resource Summary Report

Generated by FDI Lab - SciCrunch.org on Apr 20, 2025

VarFish

RRID:SCR_023710
Type: Tool

Proper Citation

VarFish (RRID:SCR_023710)

Resource Information

URL: https://www.cubi.bihealth.org/software/varfish/

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Description: Web application for quality control, filtering, prioritization, analysis, and user based annotation of DNA variant data with focus on rare disease genetics. Comprehensive DNA variant analysis for diagnostics and research.

Resource Type: data processing software, data analysis software, software application, software resource

Defining Citation: DOI:10.1093/nar/gkaa241

Keywords: Genetics, Variant Call Format, Filtering, Quality control,

Funding: Berlin Institute of Health Charité funds ;
Stiftung Charité

Availability: Free, Available for download, Freely available

Resource Name: VarFish

Resource ID: SCR_023710

Alternate URLs: https://github.com/bihealth/varfish-server

License: MIT license

Record Creation Time: 20230622T050213+0000

Record Last Update: 20250420T015252+0000

Ratings and Alerts

No rating or validation information has been found for VarFish.

No alerts have been found for VarFish.

Data and Source Information

Source: SciCrunch Registry

Usage and Citation Metrics

We found 2 mentions in open access literature.

Listed below are recent publications. The full list is available at FDI Lab - SciCrunch.org.

Kakar N, et al. (2025) Further evidence of biallelic NAV3 variants associated with recessive neurodevelopmental disorder with dysmorphism, developmental delay, intellectual disability, and behavioral abnormalities. Human genetics, 144(1), 55.

Nieminen M, et al. (2022) SODAR: managing multiomics study data and metadata. GigaScience, 12.