

Resource Summary Report

Generated by [dkNET](#) on Apr 16, 2025

mCSM

RRID:SCR_010776

Type: Tool

Proper Citation

mCSM (RRID:SCR_010776)

Resource Information

URL: <http://bleoberis.bioc.cam.ac.uk/mcsm>

Proper Citation: mCSM (RRID:SCR_010776)

Description: Data analysis service to the study of missense mutations which relies on graph-based signatures.

Abbreviations: mCSM

Synonyms: mCSM: predicting the effect of mutations in proteins using graph-based signatures

Resource Type: analysis service resource, service resource, production service resource, data analysis service

Defining Citation: [PMID:24281696](#)

Keywords: mutation, protein, protein stability, protein-protein, protein-dna, data set

Funding:

Resource Name: mCSM

Resource ID: SCR_010776

Alternate IDs: OMICS_00133

Record Creation Time: 20220129T080300+0000

Record Last Update: 20250416T063610+0000

Ratings and Alerts

No rating or validation information has been found for mCSM.

No alerts have been found for mCSM.

Data and Source Information

Source: [SciCrunch Registry](#)

Usage and Citation Metrics

We found 74 mentions in open access literature.

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

Yuan M, et al. (2025) Prevalence of IMPG1 and IMPG2 Mutations Leading to Retinitis Pigmentosa or Vitelliform Macular Dystrophy in a Cohort of Patients with Inherited Retinal Dystrophies. *Genes*, 16(1).

Basrai A, et al. (2025) Computational analyses of drug resistance mutations in katG and emb complexes in *Mycobacterium tuberculosis*. *Proteins*, 93(1), 359.

Teng R, et al. (2024) Computer-Aided Design to Improve the Thermal Stability of *Rhizomucor miehei* Lipase. *Foods (Basel, Switzerland)*, 13(24).

Pan C, et al. (2024) Module control of network analysis in psychopathology. *iScience*, 27(7), 110302.

Colbert BM, et al. (2024) The natural history and genotype-phenotype correlations of TMPRSS3 hearing loss: an international, multi-center, cohort analysis. *Human genetics*, 143(5), 721.

Tanshee RR, et al. (2024) A comprehensive in silico investigation into the pathogenic SNPs in the RTEL1 gene and their biological consequences. *PloS one*, 19(9), e0309713.

Nunes WVB, et al. (2024) A comprehensive evolutionary scenario for the origin and neofunctionalization of the *Drosophila* speciation gene *Odysseus (OdsH)*. *G3 (Bethesda, Md.)*, 14(3).

Ginatt AA, et al. (2024) A metabolic modeling-based framework for predicting trophic dependencies in native rhizobiomes of crop plants. *eLife*, 13.

Goutam RK, et al. (2024) Impact of Frequent ARID1A Mutations on Protein Stability: Insights into Cancer Pathogenesis. *Research square*.

Fasken MB, et al. (2023) A Biallelic Variant of the RNA Exosome Gene EXOSC4 Causes

Translational Defects Associated with a Neurodevelopmental Disorder. medRxiv : the preprint server for health sciences.

Su L, et al. (2023) Impact of N221S missense mutation in human ribonucleotide reductase small subunit b on mitochondrial DNA depletion syndrome. *Scientific reports*, 13(1), 19899.

Lin W, et al. (2023) Comparison of clinical outcomes of modified laminoplasty with preservation of muscle group inserted into C2 and C7 spinous processes versus conventional C3-C7 laminoplasty: a prospective, randomized, controlled, noninferiority trial. *International journal of surgery (London, England)*, 109(4), 905.

Piniella D, et al. (2023) Experimental and Bioinformatic Insights into the Effects of Epileptogenic Variants on the Function and Trafficking of the GABA Transporter GAT-1. *International journal of molecular sciences*, 24(2).

Kahane I, et al. (2023) A mutation-level covariate model for mutational signatures. *PLoS computational biology*, 19(6), e1011195.

Wang Q, et al. (2023) Two SOX11 variants cause Coffin-Siris syndrome with a new feature of sensorineural hearing loss. *American journal of medical genetics. Part A*, 191(1), 183.

Suleman M, et al. (2023) Elucidating the binding mechanism of SARS-CoV-2 NSP6-TBK1 and structure-based designing of phytocompounds inhibitors for instigating the host immune response. *Frontiers in chemistry*, 11, 1346796.

Guo L, et al. (2023) Posterior endoscopic decompression combined with anterior cervical discectomy and fusion versus posterior laminectomy and fusion for multilevel cervical spondylotic myelopathy: a retrospective case-control study. *BMC musculoskeletal disorders*, 24(1), 578.

Jiang Q, et al. (2022) Liquid-Metal-Based Magnetic Controllable Soft Microswitch with Rapid and Reliable Response for Intelligent Soft Systems. *Micromachines*, 13(12).

Stephenson SEM, et al. (2022) Germline variants in tumor suppressor FBXW7 lead to impaired ubiquitination and a neurodevelopmental syndrome. *American journal of human genetics*, 109(4), 601.

Vieider RP, et al. (2022) The 50 most cited studies on posterior tibial slope in joint preserving knee surgery. *Journal of experimental orthopaedics*, 9(1), 119.