

Resource Summary Report

Generated by [PRECISE-TBI](#) on Sep 7, 2026

SIMLINK

RRID:SCR_009387

Type: Tool

Proper Citation

SIMLINK (RRID:SCR_009387)

Resource Information

URL: <http://csg.sph.umich.edu/boehnke/simlink.php>

Proper Citation: SIMLINK (RRID:SCR_009387)

Description: Software program to estimate the probability (power) of detecting linkage given family history information on a set of identified pedigrees. (entry from Genetic Analysis Software)

Resource Type: software application, software resource

Keywords: gene, genetic, genomic, fortran, ms-dos, unix, sunos, vms

Resource Name: SIMLINK

Resource ID: SCR_009387

Alternate IDs: nlx_154625

Record Creation Time: 20220129T080252+0000

Record Last Update: 20260905T133251+0000

Ratings and Alerts

No rating or validation information has been found for SIMLINK.

No alerts have been found for SIMLINK.

Data and Source Information

Source: [SciCrunch Registry](#)

Usage and Citation Metrics

We found 3 mentions in open access literature.

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

Tomberg K, et al. (2018) Whole exome sequencing of ENU-induced thrombosis modifier mutations in the mouse. PLoS genetics, 14(9), e1007658.

Dong Y, et al. (2013) Genome-wide scan for hypertension linkage to chromosome 12q23.1 - q23.3 in a Chinese family. The Indian journal of medical research, 137(5), 935.

Kuchtey J, et al. (2011) Mapping of the disease locus and identification of ADAMTS10 as a candidate gene in a canine model of primary open angle glaucoma. PLoS genetics, 7(2), e1001306.