

Resource Summary Report

Generated by [RRID](#) on Sep 10, 2026

FASTLINK

RRID:SCR_009177

Type: Tool

Proper Citation

FASTLINK (RRID:SCR_009177)

Resource Information

URL: <http://www.ncbi.nlm.nih.gov/CBBresearch/Schaffer/fastlink.html>

Proper Citation: FASTLINK (RRID:SCR_009177)

Description: Software application (entry from Genetic Analysis Software)

Abbreviations: FASTLINK

Synonyms: faster version of LINKAGE LINKAGE

Resource Type: software application, software resource

Defining Citation: [PMID:8807326](#)

Keywords: gene, genetic, genomic, c, unix, vms, ms-dos, .. and can also run in parallel on shared memory unix machines

Resource Name: FASTLINK

Resource ID: SCR_009177

Alternate IDs: OMICS_28405, nlx_154309

Alternate URLs: <https://sources.debian.org/src/fastlink/>

Record Creation Time: 20220129T080251+0000

Record Last Update: 20260905T133245+0000

Ratings and Alerts

No rating or validation information has been found for FASTLINK.

No alerts have been found for FASTLINK.

Data and Source Information

Source: [SciCrunch Registry](#)

Usage and Citation Metrics

We found 59 mentions in open access literature.

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

Sun W, et al. (2024) Altered chromatin topologies caused by balanced chromosomal translocation lead to central iris hypoplasia. Nature communications, 15(1), 5048.

Posadas-Cantera S, et al. (2024) The effect of HLA genotype on disease onset and severity in CTLA-4 insufficiency. Frontiers in immunology, 15, 1447995.

Dokshin FA, et al. (2024) Party affiliation predicts homeowners' decisions to install solar PV, but partisan gap wanes with improved economics of solar. Proceedings of the National Academy of Sciences of the United States of America, 121(29), e2303519121.

Knight DJ, et al. (2024) Residential mobility and persistently depressed voting among disadvantaged adults in a large housing experiment. Proceedings of the National Academy of Sciences of the United States of America, 121(20), e2306287121.

Postans M, et al. (2024) Evaluating the risk of SARS-CoV-2 reinfection with the Omicron or Delta variant in Wales, UK. PloS one, 19(9), e0309645.

Hendricks MA, et al. (2023) Association of Household Opioid Availability With Opioid Overdose. JAMA network open, 6(3), e233385.

Tahamont S, et al. (2023) No ground truth? No problem: Improving administrative data linking using active learning and a little bit of guile. PloS one, 18(4), e0283811.

Schnell R, et al. (2023) Microsimulation of an educational attainment register to predict future record linkage quality. International journal of population data science, 8(1), 2122.

Mikl M, et al. (2022) A massively parallel reporter assay reveals focused and broadly encoded RNA localization signals in neurons. Nucleic acids research, 50(18), 10643.

Weiner SG, et al. (2022) Factors Associated With Opioid Overdose After an Initial Opioid Prescription. JAMA network open, 5(1), e2145691.

Mo CH, et al. (2022) Civilian national service programs can powerfully increase youth voter turnout. Proceedings of the National Academy of Sciences of the United States of America, 119(29), e2122996119.

Andrianou XD, et al. (2021) Evaluation of the national surveillance system for invasive meningococcal disease, Italy, 2015-2018. PloS one, 16(1), e0244889.

Leache L, et al. (2021) Incidence of Attention Deficit Hyperactivity Disorder (ADHD) Diagnoses in Navarre (Spain) from 2003 to 2019. International journal of environmental

research and public health, 18(17).

Rane MS, et al. (2021) Association of Diphtheria-Tetanus-Acellular Pertussis Vaccine Timeliness and Number of Doses With Age-Specific Pertussis Risk in Infants and Young Children. JAMA network open, 4(8), e2119118.

Iqbal H, et al. (2020) Mutations in FYCO1 identified in families with congenital cataracts. Molecular vision, 26, 334.

Andrasfay T, et al. (2020) Intergenerational Change in Birthweight: Effects of Foreign-born Status and Race/Ethnicity. Epidemiology (Cambridge, Mass.), 31(5), 649.

Avoundjian T, et al. (2020) Comparing Methods for Record Linkage for Public Health Action: Matching Algorithm Validation Study. JMIR public health and surveillance, 6(2), e15917.

Rauf B, et al. (2020) Novel mutations in LTBP2 identified in familial cases of primary congenital glaucoma. Molecular vision, 26, 14.

Farooq M, et al. (2020) RRP7A links primary microcephaly to dysfunction of ribosome biogenesis, resorption of primary cilia, and neurogenesis. Nature communications, 11(1), 5816.

Jiao X, et al. (2019) Autosomal recessive congenital cataracts linked to HSF4 in a consanguineous Pakistani family. PloS one, 14(12), e0225010.