

Resource Summary Report

Generated by [NIF](#) on Jul 30, 2026

[epialleleR](#)

RRID:SCR_023913

Type: Tool

Proper Citation

epialleleR (RRID:SCR_023913)

Resource Information

URL: <http://bioconductor.org/packages/epialleleR/>

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Description: Software R package for calling hypermethylated variant epiallele frequencies at level of genomic regions or individual cytosines in next-generation sequencing data using binary alignment map files as input. Used for sensitive allele specific methylation analysis in next generation sequencing data. Used for sensitive detection, quantification and visualisation of mosaic epimutations in methylation sequencing data.

Resource Type: software toolkit, software resource

Defining Citation: [DOI:10.1101/2022.06.30.498213](https://doi.org/10.1101/2022.06.30.498213)

Keywords: BAM files, binary alignment map files, allele specific methylation analysis, methylation sequencing data, next generation sequencing data, hypermethylated variant epiallele frequencies calling,

Funding: K.G.Jebesen Foundation ;
Norwegian Cancer Society ;
Norwegian Research Council

Availability: Free, Available for download, Freely available

Resource Name: epialleleR

Resource ID: SCR_023913

Alternate URLs: <https://github.com/BBCG/epialleleR>

License: Artistic 2.0

Record Creation Time: 20230804T050227+0000

Record Last Update: 20260729T044558+0000

Ratings and Alerts

No rating or validation information has been found for epialleleR.

No alerts have been found for epialleleR.

Data and Source Information

Source: [SciCrunch Registry](#)

Usage and Citation Metrics

We found 1 mentions in open access literature.

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

Nikolaienko O, et al. (2022) epialleleR: an R/Bioconductor package for sensitive allele-specific methylation analysis in NGS data. GigaScience, 12.