

Package ‘eds’

August 20, 2025

Title eds: Low-level reader for Alevin EDS format

Version 1.11.0

Description This packages provides a single function, readEDS.

This is a low-level utility for reading in Alevin EDS format into R.

This function is not designed for end-users but instead the package is predominantly for simplifying package dependency graph for other Bioconductor packages.

Depends Matrix

Imports Rcpp

Suggests knitr, tximportData, testthat (>= 3.0.0)

LinkingTo Rcpp

SystemRequirements C++11

License GPL-2

Encoding UTF-8

URL <https://github.com/mikelove/eds>

biocViews Sequencing, RNASeq, GeneExpression, SingleCell

VignetteBuilder knitr

LazyData true

RoxygenNote 7.1.2

Config/testthat/edition 3

git_url <https://git.bioconductor.org/packages/eds>

git_branch devel

git_last_commit e1f3039

git_last_commit_date 2025-04-15

Repository Bioconductor 3.22

Date/Publication 2025-08-20

Author Avi Srivastava [aut, cre],
Michael Love [aut, ctb]

Maintainer Avi Srivastava <asrivastava@cs.stonybrook.edu>

Contents

readEDS	2
-------------------	---

Index**3**

readEDS

*A low-level utility function for quickly reading in Alevin EDS format***Description**

This provides a simple utility for reading in EDS format. Note that most users will prefer to use tximport or tximeta. This function and package exist in order to simplify the dependency graph for other packages.

Usage

```
readEDS(numOfGenes, numOfOriginalCells, countMatFilename, tierImport = FALSE)
```

Arguments

```
numOfGenes      number of genes
numOfOriginalCells
                number of cells
countMatFilename
                pointer to the EDS file, quants_mat.gz
tierImport      whether the countMatFilename refers to a quants tier file
```

Value

a genes x cells sparse matrix, of the class dgCMatrx

Examples

```
# point to files
dir0 <- system.file("extdata", package="tximportData")
samps <- list.files(file.path(dir0, "alevin"))
dir <- file.path(dir0, "alevin", samps[3], "alevin")
quant.mat.file <- file.path(dir, "quants_mat.gz")
barcode.file <- file.path(dir, "quants_mat_rows.txt")
gene.file <- file.path(dir, "quants_mat_cols.txt")

# readEDS() requires knowing the number of cells and genes
cell.names <- readLines(barcode.file)
gene.names <- readLines(gene.file)
num.cells <- length(cell.names)
num.genes <- length(gene.names)

# reading in the sparse matrix
mat <- readEDS(numOfGenes=num.genes,
               numOfOriginalCells=num.cells,
               countMatFilename=quant.mat.file)
```

Index

readEDS, [2](#)