

Package ‘dandelionR’

October 15, 2025

Title Single-cell Immune Repertoire Trajectory Analysis in R

Version 1.0.2

Description dandelionR is an R package for performing single-cell immune repertoire trajectory analysis, based on the original python implementation. It provides the necessary functions to interface with scRepertoire and a custom implementation of an absorbing Markov chain for pseudo-time inference, inspired by the Palantir Python package.

License MIT + file LICENSE

Encoding UTF-8

Roxygen list(markdown = TRUE)

RoxygenNote 7.3.2

biocViews Software, ImmunoOncology, SingleCell

Collate 'check.R' 'constructMarkovChain.R' 'dandelionR.R' 'data.R'
'determMultiscaleSpace.R' 'terminalStateFromMarkovChain.R'
'differentiationProbabilities.R' 'filterCells.R' 'getPbs.R'
'projectProbability.R' 'maxMinSampling.R' 'minMaxScale.R'
'markovProbability.R' 'miloUmap.R' 'projectPseudotimeToCell.R'
'setupVdjPseudobulk.R' 'splitCTgene.R' 'vdjPseudobulk.R'

Imports BiocGenerics, bluster, destiny, igraph, MASS, Matrix, methods,
miloR, purrr, rlang, S4Vectors, SingleCellExperiment, spam,
stats, SummarizedExperiment, uwot, RANN

Suggests BiocStyle, knitr, rmarkdown, RColorBrewer, scater,
scRepertoire, testthat

VignetteBuilder knitr

URL <https://www.github.com/tuonglab/dandelionR/>

BugReports <https://www.github.com/tuonglab/dandelionR/issues>

Depends R (>= 4.4.0)

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

git_branch RELEASE_3_21

git_last_commit 6da8e23

git_last_commit_date 2025-07-21

Repository Bioconductor 3.21

Date/Publication 2025-10-15

Author Jiawei Yu [aut] (ORCID: <<https://orcid.org/0009-0005-9170-7881>>),
 Nicholas Borcharding [aut] (ORCID:
 <<https://orcid.org/0000-0003-1427-6342>>),
 Kelvin Tuong [aut, cre] (ORCID:
 <<https://orcid.org/0000-0002-6735-6808>>)

Maintainer Kelvin Tuong <z.tuong@uq.edu.au>

Contents

.addColData	3
.allowedChain	4
.calDif	4
.classCheck	5
.collapse_nested_list	5
.constructMarkovChain	6
.determExtractColN	6
.determineMultiscaleSpace	7
.determTerminal	7
.extractVdj	8
.featureSpaceConstruct	8
.filterCells	9
.filterProductivity	10
.filterUnmapped	10
.findNewWaypoints	11
.generateExtractColumn	12
.generateExtractName	13
.getPbs	13
.getPbsCol	14
.getPbsPerCol	14
.KNNind	15
.maxMinSampling	15
.minMaxScale	16
.normalizeFeatureSpace	16
.normalizePerVDJ	17
.packFeatureSpace	18
.RANNinx	18
.removeEdge	19
.subsetSce	19
.terminalStateFromMarkovChain	20
.typeCheck	20
.waypiontsPerCol	21
chainAssign	21
dandelionR	22
demo_airr	23
demo_sce	23

`.addColData` 3

differentiationProbabilities	24
formatVdj	25
markovProbability	26
miloUmap	28
projectProbability	29
projectPseudotimeToCell	30
sce_vdj	32
setupVdjPseudobulk	33
splitCTgene	36
vdjPseudobulk	36
Index	39

<code>.addColData</code>	<i>add the calculated probability to the colData</i>
--------------------------	--

Description

add the calculated probability to the colData

Usage

```
.addColData(probabilities_proj, terminal_state, milo, verbose)
```

Arguments

probabilities_proj	the probabilities need to be stored
terminal_state	Integer. The index of the terminal state in the Markov chain, passed from markovProbability
milo	the milo object provided by user
verbose	logical, print warnings.

Value

a Milo object with probabilities and pseudotime in its colData slot

.allowedChain	<i>filtering cell without allowed chain status</i>
---------------	--

Description

filtering cell without allowed chain status

Usage

```
.allowedChain(sce, allowed_chain_status, verbose)
```

Arguments

sce	SingleCellExperiment object input
allowed_chain_status	the chain needs to be retain, passed from setupVdjPseudobulk
verbose	logical, print messages. Default is TRUE.

Value

SingleCellExperiment object with allowed chain status

.calDif	<i>function help to calculate the diffusion distance</i>
---------	--

Description

function help to calculate the diffusion distance

Usage

```
.calDif(idx, eigenvector, lambda_t, K)
```

Arguments

idx	integer the index of the calculated value
eigenvector	numeric vector, the eigenvector from diffusion map
lambda_t	eigenvalues to the power of t(diffusion time)
K	The number of the eigenvectors to be used in calculation

Value

updated diffusion distance matrix after one iteration

`.classCheck`*.classCheck*

Description

check whether the input is with the correct class

Usage

```
.classCheck(input, must)
```

Arguments

input	the input need to be check
must	the type we need

Value

whether or not the input is the correct class

`.collapse_nested_list` *Collapse a nested list*

Description

Collapse a nested list

Usage

```
.collapse_nested_list(input_list)
```

Arguments

input_list	input nested list.
------------	--------------------

Value

collapsed list

```
.constructMarkovChain .constructMarkovChain
```

Description

Markov chain construction

Usage

```
.constructMarkovChain(wp_data, knn., pseudotime, waypoints, vb, use_RANN)
```

Arguments

wp_data	Multi scale data of the waypoints
knn.	Number of nearest neighbors for graph construction
pseudotime	pseudotime ordering of cells
waypoints	integer vector, index of selected waypoint used to
vb	whether to print messages
use_RANN	parameter to make user choose whether to use RANN to construct Markov chain, or keep using bluster

Value

transition matrix of the markov chain

```
.determExtractColN determine the columns in the colData where the main VDJ information is stored
```

Description

determine the columns in the colData where the main VDJ information is stored

Usage

```
.determExtractColN(extract_cols, mode_option, milo)
```

Arguments

extract_cols	names of columns in the colData where the main VDJ information stores, passed from vdjPseudobulk
mode_option	Specifies the mode for extracting V(D)J genes
milo	Milo or SingleCellExperiment object provided by user

Value

a character vector stores the names of columns in the colData where the main VDJ information stores

```
.determineMultiscaleSpace  
    .determineMultiscaleSpace
```

Description

.determineMultiscaleSpace

Usage

```
.determineMultiscaleSpace(diffusionmap, n_eigs = NULL)
```

Arguments

diffusionmap	DiffusionMap object
n_eigs	integer, default is NULL. Number of eigen vectors to use. • If is not specified, the number of eigen vectors will be determined using the eigen gap.

Value

dataframe

```
.determTerminal    .determTerminal
```

Description

function in Reduce to provide waypoints

Usage

```
.determTerminal(terminal_states, i, dm_boudaries, wp_data)
```

Arguments

terminal_states	integer vector to store the generated waypoint index
i	iteration index
dm_boudaries	index of the maxium or minium value of transition matrix per row
wp_data	Multi scale data of the waypoints

Value

integer vector store the index of waypoints serve as terminal state

<code>.extractVdj</code>	<i>Specify the columns which store VDJ information, and extract the main chain from it</i>
--------------------------	--

Description

Specify the columns which store VDJ information, and extract the main chain from it

Usage

```
.extractVdj(sce, extract_cols, mode_option, verbose)
```

Arguments

<code>sce</code>	SingleCellExperiment object input
<code>extract_cols</code>	The setupVdjPseudobulk transferred parameter given by user to specify the VDJ information columns
<code>mode_option</code>	see document of setupVdjPseudobulk for detailed explanation
<code>verbose</code>	logical, print messages. Default is TRUE.

Value

SingleCellExperiment objects with column stores the information of the main VDJ information in colData slot

<code>.featureSpaceConstruct</code>	<i>Construct VDJ feature space</i>
-------------------------------------	------------------------------------

Description

Construct VDJ feature space

Usage

```
.featureSpaceConstruct(milo, extract_cols, pbs)
```

Arguments

<code>milo</code>	Milo or SingleCellExperiment object provided by user
<code>extract_cols</code>	columns of names where to extract the VDJ information
<code>pbs</code>	cell x pseudobulk adjacent matrix

Value

constructed feature space

<i>.filterCells</i>	<i>.filterCells</i>
---------------------	---------------------

Description

Helper function that identifies filter_pattern hits in determined column of sce, and then either removes the offending cells or masks the matched values with a uniform value of '(column's name)_missing'

Usage

```
.filterCells(  
    sce,  
    col_n,  
    filter_pattern = ",|None|No_contig",  
    remove_missing = TRUE  
)
```

Arguments

sce	SingleCellExperiment object, adata in python data after combineTCR, contain both vdj and seq
col_n	mode for extraction the V(D)J genes.
filter_pattern	character string, optional ', None No_contig' by default
remove_missing	bool, True by default • If TRUE, will remove cells with contigs matching the filter from the object. • If FALSE, will mask them with a uniform value dependent on the column name.

Value

filtered SingleCellExperiment object according to the parameter.

.filterProductivity	<i>filer out cell with unproductive chain</i>
---------------------	---

Description

filer out cell with unproductive chain

Usage

```
.filterProductivity(  
  sce,  
  mode_option,  
  productive_cols,  
  productive_vj,  
  productive_vdj,  
  verbose  
)
```

Arguments

- sce SingleCellExperiment input
- mode_option check setupVdjPseudobulk for detailed explanation
- productive_vj If TRUE, retains cells where the main VJ chain is productive.
- productive_vdj If TRUE, retains cells where the main VDJ chain is productive.
- verbose logical, print messages. Default is TRUE.

Value

SingleCellExperiment object after filtering on productive chain

.filterUnmapped	<i>Filter out cell with unclear mapping in VDJ information</i>
-----------------	--

Description

Filter out cell with unclear mapping in VDJ information

Usage

```
.filterUnmapped(  
  sce,  
  mode_option,  
  check_vj_mapping,  
  check_vdj_mapping,  
  main_cols,  
  check_extract_cols_mapping,  
  remove_missing,  
  verbose  
)
```

Arguments

sce	SingleCellExperiment object input
mode_option	see document of setupVdjPseudobulk for explanation
check_vj_mapping	logical vector to set whether to check V and J gene in VJ chain, passed from setupVdjPseudobulk
check_vdj_mapping	logical vector to set whether to check V, D and J gene in VDJ chain, passed from setupVdjPseudobulk
main_cols	column names in colData in which The information of main chain stores
check_extract_cols_mapping	character vector,the names of columns that needs to be checked, passed from setupVdjPseudobulk
remove_missing	option for removing the unclear mappin or just mask it, passed from setupVdjPseudobulk
verbose	logical, print messages.

Value

filtered SingleCellExperiment object

<i>.findNewWaypoints</i>	<i>function used in Reduce to find new waypoint in an iteration</i>
--------------------------	---

Description

function used in Reduce to find new waypoint in an iteration

Usage

```
.findNewWaypoints(iterdists, k, vecs, ind, datas)
```

Arguments

<code>iterdists</code>	a list containing both waypoints detected in the former iterations and the distance matrix used to find waypoints
<code>k</code>	the iteration number
<code>vecs</code>	a numeric vector used to calculate distance of waypoints to each points
<code>ind</code>	colnames

Value

a list containing updated distance matrix and new waypoints

`.generateExtractColumn`

Check whether the columns with specified names exist, if not, create them with CTgene columns

Description

Check whether the columns with specified names exist, if not, create them with CTgene columns

Usage

```
.generateExtractColumn(sce, extract_cols, verbose)
```

Arguments

<code>sce</code>	SingleCellExperiment object input
<code>extract_cols</code>	column names we aim to extract information from
<code>verbose</code>	logical, print messages. Default is TRUE.

Value

SingleCellExperiment with columns containing VDJ information in the names we've specified.

<code>.generateExtractName</code>	<i>Generate the name of columns with given parameter</i>
-----------------------------------	--

Description

Generate the name of columns with given parameter

Usage

```
.generateExtractName(sce, mode_option, verbose)
```

Arguments

<code>sce</code>	SingleCellExperiment object input
<code>mode_option</code>	see document of <code>setupVdjPseudobulk</code> for explanation
<code>verbose</code>	logical, print messages. Default is TRUE.

Value

a vecotor of colnames we need to perform main chain extraction

<code>.getPbs</code>	<i>.getPbs</i>
----------------------	----------------

Description

Helper function to ensure we have cells by pseudobulks matrix which we can use for pseudobulking.

Usage

```
.getPbs(pbs, col_to_bulk, milo, verbose = TRUE)
```

Arguments

<code>pbs</code>	pbs parameter provided by <code>vdjPseudobulk()</code> , cells by pseudobulks matrix or NULL
<code>col_to_bulk</code>	<code>col_to_bulk</code> parameter provided by <code>vdjPseudobulk()</code> , column's name of <code>colData</code> from <code>milo</code>
<code>milo</code>	SingleCellExperiment object
<code>verbose</code>	logical, whether to print messages

Value

a cell x pseudobulk matrix

<i>.getPbsCol</i>	<i>.getPbsCol</i>
-------------------	-------------------

Description

Helper function to create the new pseudobulk object's coldata.

Usage

`.getPbsCol(pbs, col_to_take, milo)`

Arguments

- `pbs` dgeMatrix, cell x pseudobulk binary matrix
- `col_to_take` character vector, names of colData of milo that need to be processed
- `milo` Milo or SingleCellExperiment object

Value

`pbs_col`, a DataFrame which will be passed to the new SingleCellExperiment object as colData of vdj x pseudobulk assays

<i>.getPbsPerCol</i>	<i>.getPbsPerCol</i>
----------------------	----------------------

Description

function used in Reduce to get the PbsCol

Usage

`.getPbsPerCol(pbs.col, anno_col, milo, pbs)`

Arguments

- `pbs.col` DataFrame object used to store the result of each iteration
- `anno_col` colname where to generate the metadata from
- `milo` milo or SingleCellExperiment objects provided by user
- `pbs` dgeMatrix, cell x pseudobulk binary matrix

Value

DataFrame object, serve as part of metadata of the new milo object

<code>.KNNind</code>	<i>Calculate the weighted adjacency matrix of knn graph and its index</i>
----------------------	---

Description

Calculate the weighted adjacency matrix of knn graph and its index

Usage

```
.KNNind(wp_data, knn.)
```

Arguments

<code>wp_data</code>	Multi scale data of the waypoints
<code>knn.</code>	Number of nearest neighbors for graph construction

Value

a list containing the weight adjacent matrix and index

<code>.maxMinSampling</code>	<i>.maxMinSampling</i>
------------------------------	------------------------

Description

function for max min sampling of waypoints

Usage

```
.maxMinSampling(datas, num_waypoints, verbose = TRUE)
```

Arguments

<code>datas</code>	data matrix along which to sample the waypoints, usually diffusion components
<code>num_waypoints</code>	number of waypoints to sample
<code>verbose</code>	logical, print progress

Value

Series reprenting the sampled waypoints

<code>.minMaxScale</code>	<i>minMaxScale</i>
---------------------------	--------------------

Description

scale the value to range 0 to 1

Usage

```
.minMaxScale(data)
```

Arguments

data dataframe need to be scale

Value

scaled value

<code>.normalizeFeatureSpace</code>	<i>Normalize Feature Space</i>
-------------------------------------	--------------------------------

Description

Make sure the sum of each V, D, and J gene within a pseudobulk equals to 1

Usage

```
.normalizeFeatureSpace(  
  pseudo_vdj_feature,  
  extract_cols,  
  min_count,  
  renormalize,  
  milo  
)
```

Arguments

pseudo_vdj_feature constructed feature space

extract_cols names of columns to extract the VDJ information

min_count the minim count of a V/D/J gene

renormalize Whether to renormalize the matrix

milo Milo or SingleCellExperiment object provided by user

Value

normalized VDJ feature space

<code>.normalizePerVDJ</code>	<i>function to normalize a specific kind of VDJ gene in feature space</i>
-------------------------------	---

Description

function to normalize a specific kind of VDJ gene in feature space

Usage

```
.normalizePerVDJ(  
  pseudo_vdj_feature,  
  col_n,  
  renormalize,  
  define.mask,  
  milo,  
  min_count  
)
```

Arguments

<code>pseudo_vdj_feature</code>	constructed feature space
<code>col_n</code>	name of column to extract the VDJ information
<code>renormalize</code>	Whether to renormalize the matrix
<code>define.mask</code>	logical vector determine whether the V/D/J gene should be masked when normalizing
<code>milo</code>	Milo or SingleCellExperiment object provided by user
<code>min_count</code>	the minim count of a V/D/J gene

Value

feature space normalized on specifed V/D/J gene

<code>.packFeatureSpace</code>	<i>Pack the normalized feature space into new Milo object</i>
--------------------------------	---

Description

The metadata will derived from the original milo

Usage

```
.packFeatureSpace(pbs, col_to_take, milo, pseudo_vdj_feature)
```

Arguments

<code>pbs</code>	cell x pseudobulk adjacent matrix
<code>col_to_take</code>	Optional character or vector of characters. Specifies names of colData of milo that need to identify the most common value for each pseudobulk
<code>milo</code>	Milo or SingleCellExperiment object provided by user
<code>pseudo_vdj_feature</code>	VDJ feature space

Value

Milo object with VDJ feature space stored in its assay

<code>.RANNinx</code>	<i>Calculate the weight adjacent matrixes of knn graph and its index using RANN</i>
-----------------------	---

Description

Calculate the weight adjacent matrixes of knn graph and its index using RANN

Usage

```
.RANNinx(wp_data, knn.)
```

Arguments

<code>wp_data</code>	Multi scale data of the waypoints
<code>knn.</code>	Number of nearest neighbors for graph construction

Value

a list containing the weight adjacent matrix and index

<code>.removeEdge</code>	<i>function used in Reduce to remove KNN's backward edges except for edges that are within the computed standard deviation</i>
--------------------------	--

Description

function used in Reduce to remove KNN's backward edges except for edges that are within the computed standard deviation

Usage

```
.removeEdge(Knn, i, rem_edges)
```

Arguments

Knn	weight KNN adjacent matrix
i	the iteration number
rem_edges	the edges that need to be removes

Value

an updated matrix after one round of iteration

<code>.subsetSce</code>	<i>Subset sce with given parameter</i>
-------------------------	--

Description

Subset sce with given parameter

Usage

```
.subsetSce(sce, subsetby, groups, verbose)
```

Arguments

sce	SingleCellExperiment object input
subsetby	subsetby Character. Name of a colData column for subsetting. given by setupVdjPseudobulk.
groups	Character vector. Specifies the subset condition for filtering. given by setupVdjPseudobulk.
verbose	logical, print messages. Default is TRUE.

Value

subsetting SingleCellExperiment object

<code>.terminalStateFromMarkovChain</code>	<i>Determine terminal states using Markov chain if end states are not provided.</i>
--	---

Description

Determine terminal states using Markov chain if end states are not provided.

Usage

```
.terminalStateFromMarkovChain(  
  Transmat,  
  wp_data,  
  pseudotime,  
  waypoints,  
  verbose  
)
```

Arguments

Transmat	Transition matrix
wp_data	Multi scale data of the waypoints
pseudotime	numeric vector, pseudotime of each pseudobulk
waypoints	integer vector, waypoint selected to construct markov chain.
verbose	Boolean, whether to print messages/warnings.

Value

terminal_state

<code>.typeCheck</code>	<i>.typeCheck</i>
-------------------------	-------------------

Description

check whether the input has the correct type

Usage

```
.typeCheck(input, must)
```

Arguments

input	the input need to be check
must	the type we need

Value

whether or not the input is the correct type

<i>.waypiontsPerCol</i>	<i>find the waypoints according to certain columns of data</i>
-------------------------	--

Description

find the waypoints according to certain columns of data

Usage

```
.waypiontsPerCol(waypoints, ind, datas, no.iterations)
```

Arguments

waypoints	integer vector used to store waypoints
ind	columns' colnames
datas	scaled diffusionmap

Value

a numeric vector containing waypoints' index

<i>chainAssign</i>	<i>Assign the V(D)J gene to the right chain.</i>
--------------------	--

Description

Assign the V(D)J gene to the right chain.

Usage

```
chainAssign(vec, num, min_len, max_len)
```

Arguments

vec	vector of V(D)J genes to assign to the right chain.
num	number of genes to return. should be 2(vj) or 3(vdj)
min_len	minimum length of the vector to account for missing constant gene.
max_len	maximum length of the vector to account for multiple chains.

Value

list contain vector of VJ + VDJ of the cell input

dandelionR

dandelionR: Single-cell immune repertoire trajectory analysis

Description

dandelionR is an R package for performing single-cell immune repertoire trajectory analysis, based on the original python implementation. It provides the necessary functions to interface with scRepertoire and a custom implementation of an absorbing Markov chain for pseudotime inference, inspired by the Palantir Python package.

Main functions

- `setupVdjPseudobulk`: Preprocess V(D)J Data for Pseudobulk Analysis.
- `vdjPseudobulk`: Generate Pseudobulk V(D)J Feature Space.
- `markovProbability`: Markov Chain Construction and Probability Calculation.
- `projectPseudotimeToCell`: Project Pseudotime and Branch Probabilities to Single Cells.

Vignettes

See the package vignettes for detailed workflows: `vignette('dandelionR')`

Installation

To install from Bioconductor, use:

```
if (!requireNamespace('BiocManager', quietly = TRUE))
  install.packages('BiocManager')
BiocManager::install('dandelionR')
```

Author(s)

Maintainer: Kelvin Tuong <z.tuong@uq.edu.au> ([ORCID](#))

Authors:

- Jiawei Yu <jiawei.yu@uq.edu.au> ([ORCID](#))
- Nicholas Borchering <borchering@wustl.edu> ([ORCID](#))

See Also

Useful links:

- <https://www.github.com/tuonglab/dandelionR/>
- Report bugs at <https://www.github.com/tuonglab/dandelionR/issues>

demo_airr*Example AIRR Dataset for V(D)J Analysis*

Description

The demo_airr object is a list of AIRR data frames from a down-sampled demo dataset derived from Suo et al., 2024, *Nature Biotechnology*.

This dataset is used in vignettes to demonstrate workflows for V(D)J analysis.

For details, see the original publication at <https://www.nature.com/articles/s41587-023-01734-7>.

The original files are available at <https://github.com/zktuong/dandelion-demo-files>.

Usage

```
data(demo_airr)
```

Format

A SingleCellExperiment object with the following slots:

list List of DataFrames containing the standardised AIRR data for each sample.

For information of AIRR rearrangements, see the AIRR Community standards at <https://docs.airr-community.org/>.

Source

Suo et al., 2024, *Nature Biotechnology*.

<https://www.nature.com/articles/s41587-023-01734-7>.

Examples

```
data(demo_airr)
```

demo_sce*Example SCE Dataset that does not contain V(D)J information*

Description

The demo_sce object is a down-sampled demo dataset derived from Suo et al., 2024, *Nature Biotechnology*.

This dataset is used in vignettes to demonstrate workflows for V(D)J analysis.

For details, see the original publication at <https://www.nature.com/articles/s41587-023-01734-7>.

The original Lymphoid cells data in h5ad format is available at <https://developmental.cellatlas.io/fetal-immune>.

Usage

```
data(demo_sce)
```

Format

A SingleCellExperiment object with the following slots:

`colData` A minimall DataFrame containing metadata about each sample, corresponding to obs in AnnData (Python). The following columns are relevant for vignette usage:

`anno_lvl2_final_clean` Cell type annotations.

`int_colData` A DataFrame containing additional assay metadata important for further analysis. Includes:

- `X_scvi`: A dimensionality reduction matrix from the scVI model.
- `UMAP`: A UMAP reduction matrix.

Source

Suo et al., 2024, *Nature Biotechnology*.

<https://www.nature.com/articles/s41587-023-01734-7>.

Examples

```
data(demo_sce)
```

```
differentiationProbabilities
```

Compute Branch Probabilities Using Markov Chain

Description

This function calculates branch probabilities for differentiation trajectories based on a Markov chain constructed from waypoint data and pseudotime ordering.

Usage

```
differentiationProbabilities(
  wp_data,
  terminal_states = NULL,
  knn = 30L,
  pseudotime,
  waypoints,
  verbose = TRUE,
  use_RANN = TRUE
)
```

Arguments

wp_data	A multi-scale data matrix or data frame representing the waypoints.
terminal_states	Integer vector. Indices of the terminal states. Default is NULL.
knn	Integer. Number of nearest neighbors for graph construction. Default is 30L.
pseudotime	Numeric vector. Pseudotime ordering of cells.
waypoints	Integer vector. Indices of selected waypoints used to construct the Markov chain.
verbose	Boolean, whether to print messages/warnings.
use_RANN	parameter to make user choose whether to use RANN to construct Markov chain, or keep using bluster

Value

A numeric matrix or data frame containing branch probabilities for each waypoint.

formatVdj	<i>Change the format of splitCTgene output.</i>
-----------	---

Description

Change the format of splitCTgene output.

Usage

```
formatVdj(gene_list)
```

Arguments

gene_list	list containing the output from splitCTgene.
-----------	--

Value

list contain vector of VJ + VDJ information of the cell input

markovProbability *Markov Chain Construction and Probability Calculation*

Description

This function preprocesses data, constructs a Markov chain, and calculates transition probabilities based on pseudotime information.

Usage

```
markovProbability(
  milo,
  diffusionmap,
  terminal_state = NULL,
  root_cell,
  knn = 30L,
  diffusiontime = NULL,
  pseudotime_key = "pseudotime",
  scale_components = TRUE,
  num_waypoints = 500,
  n_eigs = NULL,
  verbose = TRUE,
  use_RANN = TRUE
)
```

Arguments

milo	A Milo or SingleCellExperiment object. This object should have pseudotime stored in colData, which will be used to calculate probabilities. If pseudotime is available in milo, it takes precedence over the value provided through the diffusiontime parameter.
diffusionmap	A DiffusionMap object corresponding to the milo object. Used for Markov chain construction.
terminal_state	Integer. The index of the terminal state in the Markov chain.
root_cell	Integer. The index of the root state in the Markov chain.
knn	Integer. The number of nearest neighbors for graph construction. Default is 30L.
diffusiontime	Numeric vector. If pseudotime is not stored in milo, this parameter can be used to provide pseudotime values to the function.
pseudotime_key	Character. The name of the column in colData that contains the inferred pseudotime.
scale_components	Logical. If TRUE, the components will be scaled before constructing the Markov chain. Default is FALSE.
num_waypoints	Integer. The number of waypoints to sample when constructing the Markov chain. Default is 500L.

n_eigs	integer, default is NULL. Number of eigen vectors to use. • If is not specified, the number of eigen vectors will be determined using the eigen gap.
verbose	Logical. If TRUE, print progress. Default is TRUE.
use_RANN	parameter to make user choose whether to use RANN to construct Markov chain, or keep using bluster

Value

milo or SingleCellExperiment object with pseudotime, probabilities in its colData

Examples

```
data(sce_vdj)
# downsample to first 2000 cells
sce_vdj <- sce_vdj[, 1:2000]
sce_vdj <- setupVdjPseudobulk(sce_vdj,
  already.productive = FALSE,
  allowed_chain_status = c("Single pair", "Extra pair")
)
# Build Milo Object
set.seed(100)
milo_object <- miloR::Milo(sce_vdj)
milo_object <- miloR::buildGraph(milo_object,
  k = 50, d = 20,
  reduced.dim = "X_scvi"
)
milo_object <- miloR::makeNhoods(milo_object,
  reduced_dims = "X_scvi",
  d = 20
)

# Construct Pseudobulked VDJ Feature Space
pb.milo <- vdjPseudobulk(milo_object, col_to_take = "anno_lvl_2_final_clean")
pb.milo <- scater::runPCA(pb.milo, assay.type = "Feature_space")

# Define root and branch tips
pca <- t(as.matrix(SingleCellExperiment::reducedDim(pb.milo, type = "PCA")))
branch.tips <- c(which.min(pca[, 2]), which.max(pca[, 2]))
names(branch.tips) <- c("CD8+T", "CD4+T")
root <- which.min(pca[, 1])

# Construct Diffusion Map
dm <- destiny::DiffusionMap(t(pca), n_pcs = 10, n_eigs = 5)
dif.pse <- destiny::DPT(dm, tips = c(root, branch.tips), w_width = 0.1)

# Markov Chain Construction
pb.milo <- markovProbability(
  milo = pb.milo,
  diffusionmap = dm,
  diffusiontime = dif.pse[[paste0("DPT", root)]],
```

```

    terminal_state = branch.tips,
    root_cell = root,
    pseudotime_key = "pseudotime"
)

```

miloUmap

Perform UMAP on the Adjacency Matrix of a Milo Object

Description

This function uses `uwot::umap` to perform UMAP dimensionality reduction on the adjacency matrix of the KNN graph in a Milo object.

Usage

```

miloUmap(
  milo,
  slot_name = "UMAP_knngraph",
  n_neighbors = 50L,
  metric = "euclidean",
  min_dist = 0.3,
  use_graph = TRUE,
  ...
)

```

Arguments

<code>milo</code>	the milo object with knn graph that needed to conduct umap on.
<code>slot_name</code>	character, with default 'UMAP_knngraph'. The slot name in <code>reduceDim</code> where the result store
<code>n_neighbors</code>	integer, with default 50L. - the size of local neighborhood (in terms of number of neighboring sample points) used for manifold approximation. - Here, the goal is to create large enough neighborhoods to capture the local manifold structure to allow for hypersampling.
<code>metric</code>	character, with default 'euclidean' the choice of metric used to measure distance to find nearest neighbors. Default is 'euclidean'.
<code>min_dist</code>	numeric, with default 0.3 the minimum distance between points in the low dimensional space
<code>use_graph</code>	Logical, default TRUE. Whether to run UMAP on the graph adjacency matrix (TRUE) as in Dandelion, or directly on the latent space (FALSE) for faster performance.
<code>...</code>	other parameters passed to <code>uwot::umap</code>

Value

milo object with umap reduction

Examples

```
data(sce_vdj)
# downsample to just 1000 cells
sce_vdj <- sce_vdj[, 1:1000]
sce_vdj <- setupVdjPseudobulk(sce_vdj,
  already.productive = FALSE,
  allowed_chain_status = c("Single pair", "Extra pair")
)
# Build Milo Object
milo_object <- miloR::Milo(sce_vdj)
milo_object <- miloR::buildGraph(milo_object,
  k = 50, d = 20,
  reduced.dim = "X_scvi"
)
milo_object <- miloR::makeNhoods(milo_object,
  reduced_dims = "X_scvi", d = 20
)

# Construct UMAP on Milo Neighbor Graph
milo_object <- miloUmap(milo_object)
```

projectProbability	<i>Project Probabilities from Markov Chain to Pseudobulks</i>
--------------------	---

Description

This function projects probabilities calculated from a Markov chain onto each pseudobulk based on a diffusion distance matrix.

Usage

```
projectProbability(
  diffusionmap,
  waypoints,
  probabilities,
  t = 1,
  verbose = TRUE
)
```

Arguments

diffusionmap	diffusion map, used to reconstruct diffusion distance matrix
waypoints	Integer vector. Indices of the waypoints used in the Markov chain.

probabilities	Numeric vector. Probabilities associated with the waypoints, calculated from the Markov chain.
t	Numeric. The diffusion time to be used in the projection.
verbose	Boolean, whether to print messages/warnings.

Value

each pseudobulk's probabilities

projectPseudotimeToCell

Project Pseudotime and Branch Probabilities to Single Cells

Description

This function projects pseudotime and branch probabilities from pseudobulk data to single-cell resolution (milo). The results are stored in the colData of the milo object.

Usage

```
projectPseudotimeToCell(
  milo,
  pb_milo,
  term_states = NULL,
  pseudotime_key = "pseudotime",
  suffix = "",
  verbose = TRUE
)
```

Arguments

milo	A SingleCellExperiment or Milo object. Represents single-cell data where pseudotime and branch probabilities will be projected.
pb_milo	A pseudobulk Milo object. Contains aggregated branch probabilities and pseudotime information to be transferred to single cells.
term_states	Named vector of terminal states, with branch probabilities to be transferred. The names should correspond to branches of interest.
pseudotime_key	Character. The column name in colData of pb_milo that contains the pseudotime information which was used in the markovProbability function. Default is "pseudotime".
suffix	Character. A suffix to be added to the new column names in colData. Default is an empty string ('').
verbose	Boolean, whether to print messages/warnings.

Value

subset of milo or SingleCellExperiment object where cell that do not belong to any neighbourhood are removed and projected pseudotime information stored colData

Examples

```
data(sce_vdj)
# downsample to first 2000 cells
sce_vdj <- sce_vdj[, 1:2000]
sce_vdj <- setupVdjPseudobulk(sce_vdj,
  already.productive = FALSE,
  allowed_chain_status = c("Single pair", "Extra pair")
)
# Build Milo Object
set.seed(100)
milo_object <- miloR::Milo(sce_vdj)
milo_object <- miloR::buildGraph(milo_object,
  k = 50, d = 20,
  reduced.dim = "X_scvi"
)
milo_object <- miloR::makeNhoods(milo_object,
  reduced_dims = "X_scvi",
  d = 20
)

# Construct Pseudobulked VDJ Feature Space
pb.milo <- vdjPseudobulk(milo_object, col_to_take = "anno_lvl_2_final_clean")
pb.milo <- scater::runPCA(pb.milo, assay.type = "Feature_space")

# Define root and branch tips
pca <- t(as.matrix(SingleCellExperiment::reducedDim(pb.milo, type = "PCA")))
branch.tips <- c(which.min(pca[, 2]), which.max(pca[, 2]))
names(branch.tips) <- c("CD8+T", "CD4+T")
root <- which.min(pca[, 1])

# Construct Diffusion Map
dm <- destiny::DiffusionMap(t(pca), n_pcs = 10, n_eigs = 5)
dif.pse <- destiny::DPT(dm, tips = c(root, branch.tips), w_width = 0.1)

# Markov Chain Construction
pb.milo <- markovProbability(
  milo = pb.milo,
  diffusionmap = dm,
  diffusiontime = dif.pse[[paste0("DPT", root)]],
  terminal_state = branch.tips,
  root_cell = root,
  pseudotime_key = "pseudotime"
)
# Project Pseudobulk Data
projected_milo <- projectPseudotimeToCell(
  milo_object,
  pb.milo,
```

```
branch.tips,
pseudotime_key = "pseudotime"
)
```

sce_vdj	Example Dataset for V(D)J Analysis
---------	------------------------------------

Description

The sce_vdj object is a down-sampled demo dataset derived from Suo et al., 2024, *Nature Biotechnology*.
This dataset is used in vignettes to demonstrate workflows for V(D)J analysis.
For details, see the original publication at <https://www.nature.com/articles/s41587-023-01734-7>.

Usage

```
data(sce_vdj)
```

Format

A SingleCellExperiment object with the following slots:

colData A DataFrame containing metadata about each sample, corresponding to obs in AnnData (Python). The following columns are relevant for vignette usage:

productive_(mode)_VDJ, productive_(mode)_VJ Factors indicating whether the heavy or light chain is productive. mode refers to the extraction mode for V(D)J genes and can be one of:

- 'abT': TCR alpha-beta
- 'gdT': TCR gamma-delta
- 'B': BCR

Gene segment fields Gene segment annotations with column names in the format (v/d/j)_call_(mode)_(VDJ/VJ). Examples include:

- v_call_abT_VDJ: V gene for TCR alpha-beta VDJ recombination
- d_call_abT_VJ: D gene for TCR alpha-beta VJ recombination

chain_status A factor describing the receptor chain's status.

anno_lvl1_2_final_clean Cell type annotations.

int_colData A DataFrame containing additional assay metadata important for further analysis. Includes:

- X_scvi: A dimensionality reduction matrix from the scVI model.
- UMAP: A UMAP reduction matrix.

Source

Suo et al., 2024, *Nature Biotechnology*.
<https://www.nature.com/articles/s41587-023-01734-7>.

Examples

```
data(sce_vdj)
```

```
setupVdjPseudobulk      Preprocess V(D)J Data for Pseudobulk Analysis
```

Description

This function preprocesses single-cell V(D)J sequencing data for pseudobulk analysis. It filters data based on productivity and chain status, subsets data, extracts main V(D)J genes, and removes unmapped entries.

Usage

```
setupVdjPseudobulk(
  sce,
  mode_option = c("abT", "gdT", "B"),
  already.productive = TRUE,
  productive_cols = NULL,
  productive_vj = TRUE,
  productive_vdj = TRUE,
  allowed_chain_status = NULL,
  subsetby = NULL,
  groups = NULL,
  extract_cols = NULL,
  filter_unmapped = TRUE,
  check_vj_mapping = c(TRUE, TRUE),
  check_vdj_mapping = c(TRUE, FALSE, TRUE),
  check_extract_cols_mapping = NULL,
  remove_missing = TRUE,
  verbose = TRUE
)
```

Arguments

sce	A SingleCellExperiment object. V(D)J data should be contained in colData for filtering.
mode_option	Optional character. Specifies the mode for extracting V(D)J genes. If NULL, extract_cols must be specified. Default is NULL.
already.productive	Logical. Whether the data has already been filtered for productivity. If TRUE, skips productivity filtering. Default is FALSE.
productive_cols	Character vector. Names of colData columns used for productivity filtering. Default is NULL.

productive_vj	Logical. If TRUE, retains cells where the main VJ chain is productive. Default is TRUE.
productive_vdj	Logical. If TRUE, retains cells where the main VDJ chain is productive. Default is TRUE.
allowed_chain_status	Character vector. Specifies chain statuses to retain. Valid options include `c('single pair', 'Extra pair', 'Extra pair-exception', 'Orphan VDJ', 'Orphan VDJ-exception')`. Default is NULL.
subsetby	Character. Name of a colData column for subsetting. Default is NULL.
groups	Character vector. Specifies the subset condition for filtering. Default is NULL.
extract_cols	Character vector. Names of colData columns where V(D)J information is stored, used instead of the standard columns. Default is NULL.
filter_unmapped	Logic. Whether to filter unmapped data. Default is TRUE.
check_vj_mapping	Logic vector. Whether to check for VJ mapping. Default is c(TRUE, TRUE). • If the first element is TRUE, function will filter the unmapped data in V gene of the VJ chain • If the second element is TRUE, function will filter the unmapped data in J gene of the VJ chain
check_vdj_mapping	Logic vector. Specifies columns to check for VDJ mapping. Default is c(TRUE, FALSE, 'TRUE'). • If the first element is TRUE, function will filter the unmapped data in V gene of the VDJ chain • If the second element is TRUE, function will filter the unmapped data in D gene of the VDJ chain • If the third element is TRUE, function will filter the unmapped data in J gene of the VDJ chain
check_extract_cols_mapping	Character vector. Specifies columns related to extract_cols for mapping checks. Default is NULL.
remove_missing	Logical. If TRUE, removes cells with contigs matching the filter. If FALSE, masks them with uniform values. Default is TRUE.
verbose	Logical. Whether to print messages. Default is TRUE.

Details

The function performs the following preprocessing steps:

- **Productivity Filtering:**
 - Skipped if already.productive = TRUE.
 - Filters cells based on productivity using productive_cols or standard colData columns named productive_{mode_option}_{type} (where type is 'VDJ' or 'VJ').
 - *mode_option*

- * function will check colData(s) named `productive_{mode_option}_{type}`, where type should be 'VDJ' or 'VJ' or both, depending on values of `productive_vj` and `productive_vdj`.
- * If set as NULL, the function needs the option 'extract_cols' to be specified
- *productive_cols*
 - * must be specified when productivity filtering is need to conduct and mode_option is NULL.
 - * where VDJ/VJ information is stored so that this will be used instead of the standard columns.
- *productive_vj, productive_vdj*
 - * If TRUE, cell will only be kept if the main V(D)J chain is productive
- **Chain Status Filtering:**
 - Retains cells with chain statuses specified by `allowed_chain_status`.
- **Subsetting:**
 - Conducted only if both `subsetby` and `groups` are provided.
 - Retains cells matching the `groups` condition in the `subsetby` column.
- **Main V(D)J Extraction:**
 - Uses `extract_cols` to specify custom columns for extracting V(D)J information.
- **Unmapped Data Filtering:**
 - decided to removes or masks cells based on `filter_unmapped`.
 - Checks specific columns for unclear mappings using `check_vj_mapping`, `check_vdj_mapping`, or `check_extract_cols_mapping`.
 - *filter_unmapped*
 - * pattern to be filtered from object.
 - * If is set to be NULL, the filtering process will not start
 - *check_vj_mapping, check_vdj_mapping*
 - * only colData specified by these arguments (`check_vj_mapping` and `check_vdj_mapping`) will be checked for unclear mappings
 - *check_extract_cols_mapping, related to extract_cols*
 - * Only colData specified by the argument will be checked for unclear mapping, the colData should first specified by `extract_cols`
 - *remove_missing*
 - * If TRUE, will remove cells with contigs matching the filter from the object.
 - * If FALSE, will mask them with a uniform value dependent on the column name.

Value

filtered SingleCellExperiment object

Examples

```
# load data
data(sce_vdj)
# check the dimension
```

```

dim(sce_vdj)
# filtered the data
sce_vdj <- setupVdjPseudobulk(
  sce = sce_vdj,
  mode_option = "abT", # set the mode to alpha-beta TCR
  allowed_chain_status = c("Single pair", "Extra pair"),
  already.productive = FALSE
) # need to filter the unproductive cells
# check the remaining dim
dim(sce_vdj)

```

splitCTgene	<i>Split the V(D)J genes from CTgene column and store them separately.</i>
-------------	--

Description

Split the V(D)J genes from CTgene column and store them separately.

Usage

```
splitCTgene(sce)
```

Arguments

sce SingleCellExperiment object after conducting scRepertoire::combineTCR()

Value

list contain vector of VJ & VDJ genes from each cell

vdjPseudobulk	<i>Generate Pseudobulk V(D)J Feature Space</i>
---------------	--

Description

This function creates a pseudobulk V(D)J feature space from single-cell data, aggregating V(D)J information into pseudobulk groups. It supports input as either a Milo object or a SingleCellExperiment object.

Usage

```
vdjPseudobulk(
  milo,
  pbs = NULL,
  col_to_bulk = NULL,
  extract_cols = c("v_call_abT_VDJ_main", "j_call_abT_VDJ_main", "v_call_abT_VJ_main",
    "j_call_abT_VJ_main"),
  mode_option = c("abT", "gdT", "B"),
  col_to_take = NULL,
  normalise = TRUE,
  renormalize = FALSE,
  min_count = 1L,
  verbose = TRUE
)
```

Arguments

<code>milo</code>	A Milo or SingleCellExperiment object containing V(D)J data.
<code>pbs</code>	Optional. A binary matrix with cells as rows and pseudobulk groups as columns. • If <code>milo</code> is a Milo object, this parameter is not required. • If <code>milo</code> is a SingleCellExperiment object, either <code>pbs</code> or <code>col_to_bulk</code> must be provided.
<code>col_to_bulk</code>	Optional character or character vector. Specifies colData column(s) to generate pbs. If multiple columns are provided, they will be combined. Default is NULL. • If <code>milo</code> is a Milo object, this parameter is not required. • If <code>milo</code> is a SingleCellExperiment object, either <code>pbs</code> or <code>col_to_bulk</code> must be provided.
<code>extract_cols</code>	Character vector. Specifies column names where V(D)J information is stored. Default is <code>c('v_call_abT_VDJ_main', 'j_call_abT_VDJ_main', 'v_call_abT_VJ_main', 'j_ca</code>
<code>mode_option</code>	Character. Specifies the mode for extracting V(D)J genes. Must be one of <code>c('B', 'abT', 'gdT')</code> . Default is <code>'abT'</code> . • Note: This parameter is considered only when <code>extract_cols = NULL</code>. • If NULL, uses column names such as <code>v_call_VDJ</code> instead of <code>v_call_abT_VDJ</code>.
<code>col_to_take</code>	Optional character or vector of characters. Specifies names of colData of <code>milo</code> that need to identify the most common value for each pseudobulk. Default is NULL.
<code>normalise</code>	Logical. If TRUE, scales the counts of each V(D)J gene group to 1 for each pseudobulk. Default is TRUE.
<code>renormalize</code>	Logical. If TRUE, rescales the counts of each V(D)J gene group to 1 for each pseudobulk after removing 'missing' calls. Useful when <code>setupVdjPseudobulk()</code> was run with <code>remove_missing = FALSE</code> . Default is FALSE.
<code>min_count</code>	Integer. Sets pseudobulk counts in V(D)J gene groups with fewer than this many non-missing calls to 0. Default is 1.
<code>verbose</code>	Logical. If TRUE, prints messages and warnings. Default is TRUE.

Details

This function aggregates V(D)J data into pseudobulk groups based on the following logic:

- **Input Requirements:**
 - If `milo` is a `Milo` object, neither `pbs` nor `col_to_bulk` is required.
 - If `milo` is a `SingleCellExperiment` object, the user must provide either `pbs` or `col_to_bulk`.
- **Normalization:**
 - When `normalise = TRUE`, scales V(D)J counts to 1 for each pseudobulk group.
 - When `renormalize = TRUE`, rescales the counts after removing 'missing' calls.
- **Mode Selection:**
 - If `extract_cols = NULL`, the function relies on `mode_option` to determine which V(D)J columns to extract.
- **Filtering:**
 - Uses `min_count` to filter pseudobulks with insufficient counts for V(D)J groups.

Value

`SingleCellExperiment` object

Examples

```
data(sce_vdj)
sce_vdj <- setupVdjPseudobulk(sce_vdj,
  already.productive = FALSE,
  allowed_chain_status = c("Single pair", "Extra pair")
)
# Build Milo Object
milo_object <- miloR::Milo(sce_vdj)
milo_object <- miloR::buildGraph(milo_object,
  k = 50, d = 20,
  reduced.dim = "X_scvi"
)
milo_object <- miloR::makeNhoods(milo_object,
  reduced_dims = "X_scvi",
  d = 20
)

# Construct pseudobulked VDJ feature space
pb.milo <- vdjPseudobulk(milo_object, col_to_take = "anno_lv1_2_final_clean")
```

Index

* datasets

demo_airr, [23](#)
demo_sce, [23](#)
sce_vdj, [32](#)

* internal

.KNNind, [15](#)
.RANNinx, [18](#)
.addColData, [3](#)
.allowedChain, [4](#)
.calDif, [4](#)
.classCheck, [5](#)
.collapse_nested_list, [5](#)
.constructMarkovChain, [6](#)
.determExtractColN, [6](#)
.determTerminal, [7](#)
.determineMultiscaleSpace, [7](#)
.extractVdj, [8](#)
.featureSpaceConstruct, [8](#)
.filterCells, [9](#)
.filterProductivity, [10](#)
.filterUnmapped, [10](#)
.findNewWaypoints, [11](#)
.generateExtractColumn, [12](#)
.generateExtractName, [13](#)
.getPbs, [13](#)
.getPbsCol, [14](#)
.getPbsPerCol, [14](#)
.maxMinSampling, [15](#)
.minMaxScale, [16](#)
.normalizeFeatureSpace, [16](#)
.normalizePerVDJ, [17](#)
.packFeatureSpace, [18](#)
.removeEdge, [19](#)
.subsetSce, [19](#)
.terminalStateFromMarkovChain, [20](#)
.typeCheck, [20](#)
.waypiontsPerCol, [21](#)
chainAssign, [21](#)
dandelionR, [22](#)

formatVdj, [25](#)
splitCTgene, [36](#)
.KNNind, [15](#)
.RANNinx, [18](#)
.addColData, [3](#)
.allowedChain, [4](#)
.calDif, [4](#)
.classCheck, [5](#)
.collapse_nested_list, [5](#)
.constructMarkovChain, [6](#)
.determExtractColN, [6](#)
.determTerminal, [7](#)
.determineMultiscaleSpace, [7](#)
.extractVdj, [8](#)
.featureSpaceConstruct, [8](#)
.filterCells, [9](#)
.filterProductivity, [10](#)
.filterUnmapped, [10](#)
.findNewWaypoints, [11](#)
.generateExtractColumn, [12](#)
.generateExtractName, [13](#)
.getPbs, [13](#)
.getPbsCol, [14](#)
.getPbsPerCol, [14](#)
.maxMinSampling, [15](#)
.minMaxScale, [16](#)
.normalizeFeatureSpace, [16](#)
.normalizePerVDJ, [17](#)
.packFeatureSpace, [18](#)
.removeEdge, [19](#)
.subsetSce, [19](#)
.terminalStateFromMarkovChain, [20](#)
.typeCheck, [20](#)
.waypiontsPerCol, [21](#)
chainAssign, [21](#)
dandelionR, [22](#)
dandelionR-package (dandelionR), [22](#)
demo_airr, [23](#)

demo_sce, [23](#)
differentiationProbabilities, [24](#)

formatVdj, [25](#)

markovProbability, [22](#), [26](#)
miloUmap, [28](#)

projectProbability, [29](#)
projectPseudotimeToCell, [22](#), [30](#)

sce_vdj, [32](#)
setupVdjPseudobulk, [22](#), [33](#)
splitCTgene, [36](#)

vdjPseudobulk, [22](#), [36](#)