

Package ‘spillR’

September 23, 2024

Type Package

Title Spillover Compensation in Mass Cytometry Data

Version 1.0.1

Description Channel interference in mass cytometry can cause spillover and may result in miscounting of protein markers. We develop a nonparametric finite mixture model and use the mixture components to estimate the probability of spillover. We implement our method using expectation-maximization to fit the mixture model.

biocViews FlowCytometry, ImmunoOncology, MassSpectrometry, Preprocessing, SingleCell, Software, StatisticalMethod, Visualization, Regression

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Encoding UTF-8

LazyData false

Config/testthat/edition 3

RoxygenNote 7.2.3

Imports dplyr, tibble, tidyselect, stats, ggplot2, tidyr, spatstat.univar, S4Vectors, parallel

Depends R (>= 4.3.0), SummarizedExperiment, CATALYST

Suggests knitr, rmarkdown, cowplot, testthat (>= 3.0.0), BiocStyle, hexbin

VignetteBuilder knitr

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compCytof	<i>Compute spillover probability and correct for spillover</i>
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Description

Compute spillover probability and correct for spillover

Usage

```
compCytof(
  sce,
  sce_bead,
  marker_to_barcode,
  impute_value,
  overwrite = FALSE,
  n_cores = 1,
  naive = FALSE
)
```

Arguments

sce	SingleCellExperiment for the real cells
sce_bead	SingleCellExperiment for the bead experiment
marker_to_barcode	Table that maps the marker to the barcode in the beads experiment
impute_value	Imputed value for counts that are declared as spillover
overwrite	logical; if TRUE data are overwritten if FALSE data are saved in new columns
n_cores	Number of computing cores
naive	logical; if TRUE use the naive version

Value

A `SingleCellExperiment` object

Examples

```
library(CATALYST)
library(dplyr)
bc_key <- c(139, 141:156, 158:176)
sce_bead <- prepData(ss_exp)
sce_bead <- assignPrelim(sce_bead, bc_key, verbose = FALSE)
sce_bead <- applyCutoffs(estCutoffs(sce_bead))
sce_bead <- computeSpillmat(sce_bead)
data(mp_cells, package = "CATALYST")
sce <- prepData(mp_cells)
marker_to_barcode <- rowData(sce_bead)[, c("channel_name", "is_bc")] |>
  as_tibble() |>
  filter(is_bc == TRUE) |>
  mutate(barcode = bc_key) |>
  select(marker = channel_name, barcode)
spillR::compCytotf(sce, sce_bead, marker_to_barcode, impute_value = NA)
```

 compensate

Compute spillover probability and correct for spillover

Description

Compute spillover probability and correct for spillover

Usage

```
compensate(
  tb_real,
  tb_bead,
  target_marker,
  spillover_markers,
  impute_value = NA,
  n_iter = 1000
)
```

Arguments

<code>tb_real</code>	Data frame or tibble with proteins counts of real experiment
<code>tb_bead</code>	Data frame or tibble with proteins counts of bead experiment
<code>target_marker</code>	Marker name in real experiment
<code>spillover_markers</code>	Marker names in bead experiment
<code>impute_value</code>	Value for counts that are declared as spillover
<code>n_iter</code>	Maximum number of EM steps

Value

A list of class `spillr` containing

<code>tb_compensate</code>	corrected real cells
<code>tb_spill_prob</code>	probability curve
<code>convergence</code>	covergence table of EM algorithm
<code>tb_real</code>	input real cells
<code>tb_bead</code>	input bead cells
<code>target_marker</code>	input marker in real experiment
<code>spillover_markers</code>	input markers in bead experiment

<code>compensate_naive</code>	<i>Compute spillover probability and correct for spillover from beads only</i>
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Description

Compute spillover probability and correct for spillover from beads only

Usage

```
compensate_naive(
  tb_real,
  tb_bead,
  target_marker,
  spillover_markers,
  impute_value = NA
)
```

Arguments

<code>tb_real</code>	Data frame or tibble with proteins counts of real experiment
<code>tb_bead</code>	Data frame or tibble with proteins counts of bead experiment
<code>target_marker</code>	Marker name in real experiment
<code>spillover_markers</code>	Marker names in bead experiment
<code>impute_value</code>	Value for counts that are declared as spillover

Value

A list of class `spillr` containing

<code>tb_compensate</code>	corrected real cells
<code>tb_spill_prob</code>	probability curve
<code>convergence</code>	covergence table of EM algorithm
<code>tb_real</code>	input real cells
<code>tb_bead</code>	input bead cells
<code>target_marker</code>	input marker in real experiment
<code>spillover_markers</code>	input markers in bead experiment

<code>generate_bead</code>	<i>Generate dataset for vignettes and simulation studies</i>
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Description

Generate dataset for vignettes and simulation studies

Usage

```
generate_bead()
```

Value

`tibble` data frame

Examples

```
set.seed(23)
generate_bead()
```

<code>generate_real</code>	<i>Generate dataset for vignettes and simulation studies</i>
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Description

Generate dataset for vignettes and simulation studies

Usage

```
generate_real()
```

Value

`tibble` data frame

Examples

```
set.seed(23)
generate_real()
```

<code>plotDiagnostics</code>	<i>Compute spillover probability and correct for spillover</i>
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Description

Compute spillover probability and correct for spillover

Usage

```
plotDiagnostics(sce, ch)
```

Arguments

<code>sce</code>	A <code>SingleCellExperiment</code> object
<code>ch</code>	Character string specifying the channel to plot

Value

A list of `ggplot2` plots

Examples

```
library(CATALYST)
library(dplyr)
bc_key <- c(139, 141:156, 158:176)
sce_bead <- prepData(ss_exp)
sce_bead <- assignPrelim(sce_bead, bc_key, verbose = FALSE)
sce_bead <- applyCutoffs(estCutoffs(sce_bead))
sce_bead <- computeSpillmat(sce_bead)
data(mp_cells, package = "CATALYST")
sce <- prepData(mp_cells)
marker_to_barcode <- rowData(sce_bead)[, c("channel_name", "is_bc")] |>
  as_tibble() |>
  filter(is_bc == TRUE) |>
  mutate(barcode = bc_key) |>
  select(marker = channel_name, barcode)
sce <- spillR::compCytOf(sce, sce_bead, marker_to_barcode, impute_value = NA)
plotDiagnostics(sce, "Yb173Di")
```

tfm	<i>Variance stabilizing transform of counts</i>
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Description

Variance stabilizing transform of counts

Usage

tfm(x)

Arguments

x Raw count

Value

A transformed count

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