Package 'ppcseq'

April 10, 2022

Title Probabilistic Outlier Identification for RNA Sequencing
Generalized Linear Models

Version 1.2.0

Description Relative transcript abundance has proven to be a valuable tool for understanding
the function of genes in biological systems. For the differential analysis of
transcript abundance using RNA sequencing data, the negative binomial model is
by far the most frequently adopted. However, common methods that are based on a
negative binomial model are not robust to extreme outliers, which we found to be
abundant in public datasets. So far, no rigorous and probabilistic methods for
detection of outliers have been developed for RNA sequencing data, leaving the
identification mostly to visual inspection. Recent advances in Bayesian computation
allow large-scale comparison of observed data against its theoretical distribution
given in a statistical model. Here we propose ppcseq, a key quality-control tool
for identifying transcripts that include outlier data points in differential expression
analysis, which do not follow a negative binomial distribution. Applying ppcseq to
analyse several publicly available datasets using popular tools, we show that from 3
to 10 percent of differentially abundant transcripts across algorithms and datasets
had statistics inflated by the presence of outliers.

License GPL-3

Encoding UTF-8

LazyData true

Biarch true

Depends R (>= 4.1.0)

Imports methods, Rcpp (>= 0.12.0), rstan (>= 2.18.1), rstantools (>=
2.0.0), tibble, dplyr, magrittr, purrr, future, furrr, tidyr
(>= 0.8.3.9000), lifecycle, ggplot2, foreach, tidybayes, edgeR,
benchmarkme, parallel, rlang, stats, utils, graphics

LinkingTo BH (>= 1.66.0), Rcpp (>= 0.12.0), RcppEigen (>= 0.3.3.3.0),
rstan (>= 2.18.1), StanHeaders (>= 2.18.0)

Suggests knitr, testthat, BiocStyle, rmarkdown

VignetteBuilder knitr

RdMacros lifecycle

biocViews RNASeq, DifferentialExpression, GeneExpression,
    Normalization, Clustering, QualityControl, Sequencing,
    Transcription, Transcriptomics

SystemRequirements GNU make

RoxygenNote 7.1.1

Roxygen list(markdown = TRUE)

BugReports https://github.com/stemangiola/ppcseq/issues

Config/testthat/edition 3

git_url https://git.bioconductor.org/packages/ppcseq

git_branch RELEASE_3_14

git_last_commit 91373e3

git_last_commit_date 2021-10-26

Date/Publication 2022-04-10

Author Stefano Mangiola [aut, cre] (<https://orcid.org/0000-0001-7474-836X>)

Maintainer Stefano Mangiola <mangiolastefano@gmail.com>

R topics documented:

---

ppcseq-package   *The 'ppcseq' package.*

Description

Relative transcript abundance has proven to be a valuable tool for understanding the function of genes in biological systems. For the differential analysis of transcript abundance using RNA sequencing data, the negative binomial model is by far the most frequently adopted. However, common methods that are based on a negative binomial model are not robust to extreme outliers, which we found to be abundant in public datasets. So far, no rigorous and probabilistic methods for detection of outliers have been developed for RNA sequencing data, leaving the identification mostly to visual inspection. Recent advances in Bayesian computation allow large-scale comparison of observed data against its theoretical distribution given in a statistical model. Here we propose ppcseq, a key quality-control tool for identifying transcripts that include outlier data points in differential expression analysis, which do not follow a negative binomial distribution. Applying ppcseq to analyse several publicly available datasets using popular tools, we show that from 3 to 10 percent of differentially abundant transcripts across algorithms and datasets had statistics inflated by the presence of outliers.

Usage

```
data(counts)
```

Value

See documentation

References

Stan Development Team (2019). RStan: the R interface to Stan. R package version 2.19.2. https://mc-stan.org angiola S, Thomas E, Modrak M, Vehtari A, Papenfuss A (2021). "Probabilistic outlier identification for RNA sequencing generalized linear models." NAR Genomics and Bioinformatics_, *3*(1), lqab005. <URL: https://doi.org/10.1093/nargab/lqab005>.

---

counts *counts*

Description

Contains an example dataset for ppcseq, including RNA sequencing

Usage

```
counts
```

Format

An object of class `tbl_df` (inherits from `tbl`, `data.frame`) with 394821 rows and 9 columns.

---

identify_outliers *identify_outliers main*

Description

This function runs the data modeling and statistical test for the hypothesis that a transcript includes outlier biological replicate.

[Maturing]

Usage

```
identify_outliers(
  .data,
  formula = ~1,
  .sample,
  .transcript,
  .abundance,
  .significance,
  .do_check,
  percent_false_positive_genes = 1,
  how_many_negative_controls = 500,
  approximate_posterior_inference = TRUE,
  approximate_posterior_analysis = TRUE,
  draws_after_tail = 10,
  save_generated_quantities = FALSE,
  additional_parameters_to_save = c(),
  cores = detect_cores(),
  pass_fit = FALSE,
  do_check_only_on_detrimental = length(parse_formula(formula)) > 0,
  tol_rel_obj = 0.01,
  just_discovery = FALSE,
  seed = sample(seq_len(length.out = 999999), size = 1),
  adj_prob_theshold_2 = NULL,
  return_fit = FALSE
)
```

Arguments

`.data`	A tibble including a transcript name column | sample name column | read counts column | covariate columns | Pvalue column | a significance column
`formula`	A formula. The sample formula used to perform the differential transcript abundance analysis
`.sample`	A column name as symbol. The sample identifier
`.transcript`	A column name as symbol. The transcript identifier
`.abundance`	A column name as symbol. The transcript abundance (read count)
`.significance`	A column name as symbol. A column with the Pvalue, or other significance measure (preferred Pvalue over false discovery rate)
`.do_check`	A column name as symbol. A column with a boolean indicating whether a transcript was identified as differentially abundant
`percent_false_positive_genes`	
	A real between 0 and 100. It is the aimed percent of transcript being a false positive. For example, percent_false_positive_genes = 1 provide 1 percent of the calls for outlier containing transcripts that has actually not outliers.
`how_many_negative_controls`	
	An integer. How many transcript from the bottom non-significant should be taken for inferring the mean-overdispersion trend.

approximate_posterior_inference

> A boolean. Whether the inference of the joint posterior distribution should be approximated with variational Bayes It confers execution time advantage.

approximate_posterior_analysis

> A boolean. Whether the calculation of the credible intervals should be done semi-analytically, rather than with pure sampling from the posterior. It confers execution time and memory advantage.

draws_after_tail

> An integer. How many draws should on average be after the tail, in a way to inform CI.

save_generated_quantities

> A boolean. Used for development and testing purposes

additional_parameters_to_save

> A character vector. Used for development and testing purposes

cores   An integer. How many cored to be used with parallel calculations.

pass_fit   A boolean. Used for development and testing purposes

do_check_only_on_detrimental

> A boolean. Whether to test only for detrimental outliers (same direction as the fold change). It allows to test for less transcript/sample pairs and therefore higher the probability threshold.

tol_rel_obj   A real. Used for development and testing purposes

just_discovery   A boolean. Used for development and testing purposes

seed   An integer. Used for development and testing purposes

adj_prob_theshold_2

> A boolean. Used for development and testing purposes

return_fit   A boolean

Value

A nested tibble `tbl` with transcript-wise information: sample wise data | plot | ppc samples failed | tot deleterious outliers

Examples

```
library(dplyr)

data("counts")

if(Sys.info()[['sysname']] == "Linux")
result =
  counts %>%
 dplyr::mutate( is_significant = ifelse(symbol %in% c("SLC16A12", "CYP1A1", "ART3"), TRUE, FALSE) ) %>%
 ppcseq::identify_outliers(
formula = ~ Label,
sample, symbol, value,
.significance = PValue,
.do_check  = is_significant,
```

```
percent_false_positive_genes = 1,
tol_rel_obj = 0.01,
approximate_posterior_inference =TRUE,
approximate_posterior_analysis =TRUE,
how_many_negative_controls = 50,
cores=1
)
```

---

plot_credible_intervals

plot_credible interval for theoretical data distributions

Description

Plot the data along the theoretical data distribution.

Usage

```
plot_credible_intervals(.data)
```

Arguments

.data   The tibble returned by identify_outliers

Value

A tibble with an additional plot column

Examples

```
library(dplyr)

data("counts")

if(Sys.info()[['sysname']] == "Linux"){
result =
  counts %>%
 dplyr::mutate( is_significant = ifelse(symbol %in% c("SLC16A12", "CYP1A1", "ART3"), TRUE, FALSE) ) %>%
 ppcseq::identify_outliers(
formula = ~ Label,
sample, symbol, value,
.significance = PValue,
.do_check  = is_significant,
percent_false_positive_genes = 1,
tol_rel_obj = 0.01,
approximate_posterior_inference =TRUE,
approximate_posterior_analysis =TRUE,
how_many_negative_controls = 50,
cores=1
```

```
)

result_plot = result %>% plot_credible_intervals()
}
```

Index