

Package ‘mfa’

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Title Bayesian hierarchical mixture of factor analyzers for modelling genomic bifurcations

Version 1.20.0

Description MFA models genomic bifurcations using a Bayesian hierarchical mixture of factor analysers.

Depends R (>= 3.4.0)

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R topics documented:

| | |
|----------------------------|---|
| calculate_chi | 2 |
| create_synthetic | 3 |

| | |
|-------------------------------------|----|
| empirical_lambda | 4 |
| mfa | 4 |
| plot_chi | 6 |
| plot_dropout_relationship | 7 |
| plot_mfa_autocorr | 7 |
| plot_mfa_trace | 8 |
| print.mfa | 9 |
| summary.mfa | 9 |
| to_ggmcmc | 10 |

| | |
|--------------|-----------|
| Index | 11 |
|--------------|-----------|

| | |
|---------------|---|
| calculate_chi | <i>Calculate posterior chi precision parameters</i> |
|---------------|---|

Description

Calculates a data frame of the MAP estimates of χ .

Usage

```
calculate_chi(m)
```

Arguments

`m` A fit returned from `mfa`

Value

A `data_frame` with one entry for the feature names and one for the MAP estimates of `chi` (using the `posterior.mode` function from `MCMCglmm`).

Examples

```
synth <- create_synthetic(C = 20, G = 5)
m <- mfa(synth$X)
chi_map <- calculate_chi(m)
```

| | |
|------------------|------------------------------|
| create_synthetic | <i>Create synthetic data</i> |
|------------------|------------------------------|

Description

Create synthetic bifurcating data for two branches. Optionally incorporate zero inflation and transient gene expression.

Usage

```
create_synthetic(C = 100, G = 40, p_transient = 0, zero_negative = TRUE,  
  model_dropout = FALSE, lambda = 1)
```

Arguments

| | |
|---------------|---|
| C | Number of cells to simulate |
| G | Number of genes to simulate |
| p_transient | Proportion of genes that exhibit transient expression |
| zero_negative | Logical: should expression generated less than zero be set to zero? This will zero-inflate the data |
| model_dropout | Logical: if true, expression will be set to zero with the exponential dropout formula dependent on the latent expression using dropout parameter lambda |
| lambda | The dropout parameter |

Value

A list with the following entries:

- X A cell-by-feature expression matrix
- branch A vector of length C assigning cells to branches
- pst A vector of pseudotimes for each cell
- k The k parameters
- phi The ϕ parameters
- delta The δ parameters
- p_transient The proportion of genes simulated as transient according to the original function call

Examples

```
synth <- create_synthetic()
```

`empirical_lambda` *Estimate the dropout parameter*

Description

Estimate the dropout parameter

Usage

```
empirical_lambda(y, lower_limit = 0)
```

Arguments

`y` A cell-by-gene expression matrix
`lower_limit` The limit below which expression counts as 'dropout'

Value

The estimated lambda

Examples

```
synth <- create_synthetic(C = 20, G = 5, zero_negative = TRUE, model_dropout = TRUE)
lambda <- empirical_lambda(synth$X)
```

`mfa` *Fit a MFA object*

Description

Perform Gibbs sampling inference for a hierarchical Bayesian mixture of factor analysers to identify bifurcations in single-cell expression data.

Usage

```
mfa(y, iter = 2000, thin = 1, burn = iter/2, b = 2,
    zero_inflation = FALSE, pc_initialise = 1, prop_collapse = 0,
    scale_input = !zero_inflation, lambda = NULL, eta_tilde = NULL,
    alpha = 0.1, beta = 0.1, theta_tilde = 0, tau_eta = 1,
    tau_theta = 1, tau_c = 1, alpha_chi = 0.01, beta_chi = 0.01,
    w_alpha = 1/b, clamp_pseudotimes = FALSE)
```

Arguments

| | |
|--------------------------------|---|
| <code>y</code> | A cell-by-gene single-cell expression matrix or an ExpressionSet object |
| <code>iter</code> | Number of MCMC iterations |
| <code>thin</code> | MCMC samples to thin |
| <code>burn</code> | Number of MCMC samples to throw away |
| <code>b</code> | Number of branches to model |
| <code>zero_inflation</code> | Logical, should zero inflation be enabled? |
| <code>pc_initialise</code> | Which principal component to initialise pseudotimes to |
| <code>prop_collapse</code> | Proportion of Gibbs samples which should marginalise over c |
| <code>scale_input</code> | Logical. If true, input is scaled to have mean 0 variance 1 |
| <code>lambda</code> | The dropout parameter - by default estimated using the function <code>empirical_lambda</code> |
| <code>eta_tilde</code> | Hyperparameter |
| <code>alpha</code> | Hyperparameter |
| <code>beta</code> | Hyperparameter |
| <code>theta_tilde</code> | Hyperparameter |
| <code>tau_eta</code> | Hyperparameter |
| <code>tau_theta</code> | Hyperparameter |
| <code>tau_c</code> | Hyperparameter |
| <code>alpha_chi</code> | Hyperparameter |
| <code>beta_chi</code> | Hyperparameter |
| <code>w_alpha</code> | Hyperparameter |
| <code>clamp_pseudotimes</code> | This clamps the pseudotimes to their initial values and doesn't perform sampling. Should be FALSE except for diagnostics. |

Details

The column names of `Y` are used as feature (gene/transcript) names while the row names are used as cell names. If either of these is undefined then the corresponding names are set to `cell_x` or `feature_y`.

It is recommended the form of `Y` is analogous to log-expression to mitigate the impact of outliers.

In the absence of prior information, three valid local maxima in the posterior likelihood exist (see manuscript). Setting the initial values to a principal component typically fixes sampling to one of them, analogous to specifying a root cell in similar methods.

The hyper-parameter `eta_tilde` represents the expected expression in the absence of any actual expression measurements. While a Bayesian purist might reason this based on knowledge of the measurement technology, simply taking the mean of the input matrix in an Empirical Bayes style seems reasonable.

The degree of shrinkage of the factor loading matrices to a common value is given by the gamma prior on `chi`. The mean of this is α_chi / β_chi while the variance α_chi / β_chi^2 . Therefore, to obtain higher levels of shrinkage increase `alpha_chi` with respect to `beta_chi`.

The collapsed Gibbs sampling option given by `collapse` involves marginalising out `c` (the factor loading intercepts) when updating the branch assignment parameters `gamma` which tends to soften the branch assignments.

If zero inflation is enabled using the `zero_inflation` parameter then scaling should *not* be enabled.

Value

An S3 structure with the following entries:

- `traces` A list of iteration-by-dim trace matrices for several important variables
- `iter` Number of iterations
- `thin` Thinning applied
- `burn` Burn period at the start of MCMC
- `b` Number of branches modelled
- `prop_collapse` Proportion of updates for `gamma` that are collapsed
- `N` Number of cells
- `G` Number of features (genes/transcripts)
- `feature_names` Names of features
- `cell_names` Names of cells

Examples

```
synth <- create_synthetic(C = 20, G = 5)
m <- mfa(synth$X)
```

plot_chi

Plot posterior precision parameters

Description

Plot posterior precision parameters

Usage

```
plot_chi(m, nfeatures = m$G)
```

Arguments

| | |
|------------------------|--------------------------------------|
| <code>m</code> | A fit returned from <code>mfa</code> |
| <code>nfeatures</code> | Top number of |

Value

A `ggplot2` bar-plot showing the map estimates of χ^{-1}

Examples

```
synth <- create_synthetic(C = 20, G = 5)
m <- mfa(synth$X)
plot_chi(m)
```

```
plot_dropout_relationship
```

Plot the dropout relationship

Description

Plot the dropout relationship

Usage

```
plot_dropout_relationship(y, lambda = empirical_lambda(y))
```

Arguments

| | |
|--------|-------------------------------|
| y | The input data matrix |
| lambda | The estimated value of lambda |

Value

A ggplot2 plot showing the estimated dropout relationship

Examples

```
synth <- create_synthetic(C = 20, G = 5, zero_negative = TRUE, model_dropout = TRUE)
lambda <- empirical_lambda(synth$X)
plot_dropout_relationship(synth$X, lambda)
```

```
plot_mfa_autocorr
```

Plot MFA autocorrelation

Description

Plots the autocorrelation of the posterior log-likelihood.

Usage

```
plot_mfa_autocorr(m)
```

Arguments

| | |
|---|-------------------------|
| m | A fit returned from mfa |
|---|-------------------------|

Value

A ggplot2 plot returned by the ggmcmc package plotting the autocorrelation of the posterior log-likelihood.

Examples

```
synth <- create_synthetic(C = 20, G = 5)
m <- mfa(synth$X)
plot_mfa_autocorr(m)
```

plot_mfa_trace

Plot MFA trace

Description

Plots the trace of the posterior log-likelihood.

Usage

```
plot_mfa_trace(m)
```

Arguments

m A fit returned from mfa

Value

A ggplot2 plot plotting the trace of the posterior log-likelihood.

Examples

```
synth <- create_synthetic(C = 20, G = 5)
m <- mfa(synth$X)
plot_mfa_trace(m)
```

| | |
|-----------|-------------------------|
| print.mfa | <i>Print an mfa fit</i> |
|-----------|-------------------------|

Description

Print an mfa fit

Usage

```
## S3 method for class 'mfa'  
print(x, ...)
```

Arguments

| | |
|-----|----------------------------|
| x | An MFA fit returned by mfa |
| ... | Additional arguments |

Value

A string representation of an mfa object.

Examples

```
synth <- create_synthetic(C = 20, G = 5)  
m <- mfa(synth$X)  
print(m)
```

| | |
|-------------|-----------------------------|
| summary.mfa | <i>Summarise an mfa fit</i> |
|-------------|-----------------------------|

Description

Returns summary statistics of an mfa fit, including MAP pseudotime and branch allocations along with uncertainties.

Usage

```
## S3 method for class 'mfa'  
summary(object, ...)
```

Arguments

| | |
|--------|--------------------------------------|
| object | An MFA fit returned by a call to mfa |
| ... | Additional arguments |

Value

A data_frame with the following columns:

- pseudotime The MAP pseudotime estimate
- branch The MAP branch estimate
- branch_certainty The proportion of traces for which the cell is assigned to its MAP branch
- pseudotime_lower The lower bound on the 95 (HPD) credible interval
- pseudotime_upper The upper bound on the 95

Examples

```
synth <- create_synthetic(C = 20, G = 5)
m <- mfa(synth$X)
ms <- summary(m)
```

to_ggmcmc

Turn a trace list to a ggmcmc object

Description

Turn a trace list to a ggmcmc object

Usage

```
to_ggmcmc(g)
```

Arguments

g A list of trace matrices

Value

The trace list converted into a ggs object for input to ggmcmc.

Index

`calculate_chi`, [2](#)
`create_synthetic`, [3](#)

`empirical_lambda`, [4](#)

`mfa`, [4](#)

`plot_chi`, [6](#)
`plot_dropout_relationship`, [7](#)
`plot_mfa_autocorr`, [7](#)
`plot_mfa_trace`, [8](#)
`print.mfa`, [9](#)

`summary.mfa`, [9](#)

`to_ggmcmc`, [10](#)