Package ‘epitrix’

January 15, 2019

Title      Small Helpers and Tricks for Epidemics Analysis
Version    0.2.2
Description A collection of small functions useful for epidemics analysis and infectious disease modelling. This includes computation of basic reproduction numbers from growth rates, generation of hashed labels to anonymise data, and fitting discretised Gamma distributions.
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License    MIT + file LICENSE
Encoding   UTF-8
LazyData   true
Suggests   testthat, roxygen2, outbreaks, incidence (>= 1.4.1), knitr, rmarkdown
Imports    sodium, distcrete, stringi
RoxygenNote 6.1.1
URL        http://www.reploidemsconsortium.org/epitrix
BugReports https://github.com/reconhub/epitrix/issues
VignetteBuilder knitr
NeedsCompilation no
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Repository  CRAN
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clean_labels

Description
This function standardises labels e.g. used as variable names or character string values, removing non-ascii characters, replacing diacritics (e.g. é, ô) with their closest ascii equivalents, and standardises separating characters. See details for more information on label transformation.

Usage

```
clean_labels(x, sep = "_")
```

Arguments

- **x**: A vector of labels, normally provided as characters.
- **sep**: A character string used as separator, defaulting to `_`.

Details

The following changes are performed:

- all non-ascii characters are removed
- all diacritics are replaced with their non-accentuated equivalents, e.g. 'é', 'ê' and 'è' become 'e', 'ö' becomes 'o', etc.
- all characters are set to lower case
- separators are standardised to the use of a single character provided in `sep` (defaults to `_`); heading and trailing separators are removed.

Author(s)

Thibaut Jombart <thibautjombart@gmail.com>

Examples

```
clean_labels("--This is; A WeIrD**./sêtènçe...")
clean_labels("--This is; A WeIrD**./sêtènçe...", sep = ".")
input <- c("Peter and stëven", "peter-and.stëven", "pëtêr and stëven _")
input
clean_labels(input)
```
Description

These functions perform maximum-likelihood (ML) fitting of a discretised distribution. This is typically useful for describing delays between epidemiological events, such as incubation period (infection to onset) or serial intervals (primary to secondary onsets). The function `optim` is used internally for fitting.

Usage

```r
fit_disc_gamma(x, mu_ini = 1, cv_ini = 1, interval = 1, w = 0, ...)
```

Arguments

- `x`: A vector of numeric data to fit; NAs will be removed with a warning.
- `mu_ini`: The initial value for the mean 'mu', defaulting to 1.
- `cv_ini`: The initial value for the coefficient of variation 'cv', defaulting to 1.
- `interval`: The interval used for discretisation; see `distcrete`.
- `w`: The centering of the interval used for discretisation; see `distcrete`.
- `...`: Further arguments passed to `optim`.

Value

The function returns a list with human-readable parametrisation of the discretised Gamma distribution (mean, sd, cv), convergence indicators, and the discretised Gamma distribution itself as a `distcrete` object (from the `distcrete` package).

Author(s)

Thibaut Jombart <thibautjombart@gmail.com>

See Also

The `distcrete` package for discretising distributions, and `optim` for details on available optimisation procedures.

Examples

```r
## generate data
mu <- 15.3 # days
sigma <- 9.3 # days
cv <- mu / sigma
```
param <- gamma_mucv2shapescale(mu, cv)

if (require(distcrete)) {
  w <- distcrete("gamma", interval = 1,
                 shape = param$shape,
                 scale = param$scale, w = 0)

  x <- w$r(100)
  x

  fit_disc_gamma(x)
}

---

gamma_shapescale2mucv Reparameterise Gamma distributions

Description

These functions permit to use alternate parametrisations for Gamma distributions, from 'shape and scale' to 'mean (mu) and coefficient of variation (cv), and back. gamma_shapescale2mucv does the first conversion, while gamma_mucv2shapescale does the second. The function gamma_log_likelihood is a shortcut for computing Gamma log-likelihood with the alternative parametrisation (mean, cv). See 'details' for a guide of which parametrisation to use.

Usage

gamma_shapescale2mucv(shape, scale)
gamma_mucv2shapescale(mu, cv)
gamma_log_likelihood(x, mu, cv, discrete = TRUE, interval = 1, w = 0,
                      anchor = 0.5)

Arguments

<table>
<thead>
<tr>
<th>Argument</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>shape</td>
<td>The shape parameter of the Gamma distribution.</td>
</tr>
<tr>
<td>scale</td>
<td>The scale parameter of the Gamma distribution.</td>
</tr>
<tr>
<td>mu</td>
<td>The mean of the Gamma distribution.</td>
</tr>
<tr>
<td>cv</td>
<td>The coefficient of variation of the Gamma distribution, i.e. the standard</td>
</tr>
<tr>
<td></td>
<td>deviation divided by the mean.</td>
</tr>
<tr>
<td>x</td>
<td>A vector of data treated as observations drawn from a Gamma distribution,</td>
</tr>
<tr>
<td></td>
<td>for which the likelihood is to be computed.</td>
</tr>
<tr>
<td>discrete</td>
<td>A logical indicating if the distribution should be discretised; TRUE by</td>
</tr>
<tr>
<td></td>
<td>default.</td>
</tr>
<tr>
<td>interval</td>
<td>The interval used for discretisation; see distcrete.</td>
</tr>
<tr>
<td>w</td>
<td>The centering of the interval used for discretisation, defaulting to 0;</td>
</tr>
<tr>
<td></td>
<td>see distcrete.</td>
</tr>
<tr>
<td>anchor</td>
<td>The anchor used for discretisation, i.e. starting point of the discretisation</td>
</tr>
<tr>
<td></td>
<td>process; defaults to 0; see distcrete.</td>
</tr>
</tbody>
</table>
Details

The gamma distribution is described in ?dgamma is parametrised using shape and scale (or rate). However, these parameters are naturally correlated, which make them poor choices whenever trying to fit data to a Gamma distribution. Their interpretation is also less clear than the traditional mean and variance. When fitting the data, or reporting results, it is best to use the alternative parametrisation using the mean ($\mu$) and the coefficient of variation ($cv$), i.e. the standard deviation divided by the mean.

Value

A named list containing 'shape' and 'scale', or mean ('mean') and coefficient of variation ('cv').

Author(s)

Code by Anne Cori (<a.cori@imperial.ac.uk>, packaging by Thibaut Jombart <thibautjombart@gmail.com>)

Examples

```r
## set up some parameters
mu <- 10
cv <- 1

## transform into shape scale
tmp <- gamma_mucv2shapescale(mu, cv)
shape <- tmp$shape
scale <- tmp$scale

## recover original parameters when applying the revert function
gamma_shapescale2mucv(shape, scale) # compare with mu, cv

## empirical validation:
## check mean / cv of a sample derived using rgamma with
## shape and scale computed from mu and cv
gamma_sample <- rgamma(n = 10000, shape = shape, scale = scale)
mean(gamma_sample) # compare to mu
sd(gamma_sample) / mean(gamma_sample) # compare to cv
```
hash_names  

Anonymise data using scrypt

Description

This function uses the scrypt algorithm from libsodium to anonymise data, based on user-indicated data fields. Data fields are concatenated first, then each entry is hashed. The function can either return a full detailed output, or short labels ready to use for 'anonymised data'. Before concatenation (using "_" as a separator) to form labels, inputs are modified using clean_labels.

Usage

hash_names(..., size = 6, full = TRUE, salt = NULL)

Arguments

... Data fields to be hashed.
size The number of characters retained in the hash.
full A logical indicating if the a full output should be returned as a data.frame, including original labels, shortened hash, and full hash.
salt An optional object that can be coerced to a character to be used to ‘salt’ the hashing algorithm (see details). Ignored if NULL (default).

Details

The argument salt should be used for salting the algorithm, i.e. adding an extra input to the input fields (the 'salt') to change the resulting hash and prevent identification of individuals via pre-computed hash tables.

It is highly recommend to choose a secret, random salt in order make it harder for an attacker to decode the hash.

Author(s)

Thibaut Jombart <thibautjombart@gmail.com>, Dirk Shchumacher <mail@dirk-schumacher.net>

See Also

clean_labels, used to clean labels prior to hashing.

Examples

```r
first_name <- c("Jane", "Joe", "Raoul")
last_name <- c("Doe", "Smith", "Dupont")
age <- c(25, 69, 36)

hash_names(first_name, last_name, age)
```
Transform a growth rate into a reproduction number

Description

The function `r2R0` can be used to transform a growth rate into a reproduction number estimate, given a generation time distribution. This uses the approach described in Wallinga and Lipsitch (2007, Proc Roy Soc B 274:599–604) for empirical distributions. The function `lm2R0_sample` generates a sample of R0 values from a log-linear regression of incidence data stored in a `lm` object.

Usage

```r
r2R0(r, w, trunc = 1000)

lm2R0_sample(x, w, n = 100, trunc = 1000)
```

Arguments

- `r` A vector of growth rate values.
- `w` The serial interval distribution, either provided as a `distcrete` object, or as a numeric vector containing probabilities of the mass functions.
- `trunc` The number of time units (most often, days), used for truncating `w`, whenever a `distcrete` object is provided. Defaults to 1000.
- `x` A `lm` object storing a linear regression of log-incidence over time.
- `n` The number of draws of R0 values, defaulting to 100.

Details

It is assumed that the growth rate (`r`) is measured in the same time unit as the serial interval (`w` is the SI distribution, starting at time 0).

Author(s)

Code by Anne Cori <a.cori@imperial.ac.uk>, packaging by Thibaut Jombart <thibautjombart@gmail.com>
Examples

## Ebola estimates of the SI distribution from the first 9 months of
## West-African Ebola outbreak

\[
\mu <- 15.3 \text{ # days} \\
\sigma <- 9.3 \text{ # days} \\
\text{param} <- \text{gamma_mucv2shape}(\mu, \sigma / \mu)
\]

if (require(distcrete)) {
  \[w <- \text{distcrete}(\text{"gamma"}, \text{interval} = 1,\]
  \[\text{shape} = \text{param}\$\text{shape},\]
  \[\text{scale} = \text{param}\$\text{scale}, \text{w} = 0)\]
  \r2R0(c(-1, -0.001, 0, 0.001, 1), w)
}

## Use simulated Ebola outbreak and 'incidence' to get a log-linear
## model of daily incidence.

if (require(outbreaks) \& require/incidence) {
  \[i <- \text{incidence}(\text{ebola_sim}\$\text{linelist}\$\text{date_of_onset})\]
  \text{plot}(i)
  \[f <- \text{fit}(i[1:100])\]
  \[f\]
  \text{plot}(i[1:150], \text{fit} = f)
  \text{R0 <- lm2R0_sample(f}\$\text{model}, \text{w})
  \text{hist(R0,}\ 	ext{col} = \text{"grey"}, \text{border} = \text{"white"}, \text{main} = \text{"Distribution of R0")}
  \text{summary(R0)}
}

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