

Package ‘eggCounts’

November 9, 2018

Imports boot, coda, utils, testthat, numbers, lattice, rootSolve

Depends R (>= 3.4.0), Rcpp (>= 0.12.0), rstan (>= 2.18.1), methods

Suggests R.rsp, eggCountsExtra, knitr, rmarkdown

VignetteBuilder R.rsp, knitr

Title Hierarchical Modelling of Faecal Egg Counts

Version 2.1-2

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Maintainer Craig Wang <craig.wang@uzh.ch>

Description An implementation of Bayesian hierarchical models for faecal egg count data to assess anthelmintic efficacy. Bayesian inference is done via MCMC sampling using Stan.

SystemRequirements GNU make

Additional_repositories <https://craigwanguzh.github.io/eggCountsExtra-package/>

License GPL (>= 3)

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

LazyLoad yes

ByteCompile true

NeedsCompilation yes

URL <https://www.math.uzh.ch/pages/eggcount/>

RcppModules stan_fit4paired_mod, stan_fit4unpaired_mod, stan_fit4zipaired_mod, stan_fit4ziunpaired_mod, stan_fit4nb_mod, stan_fit4zinb_mod, stan_fit4indefficacy_mod, stan_fit4simple_mod

RoxygenNote 6.0.1

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R topics documented:

eggCounts-package	2
eggs	3
fecrtCI	4
fecr_probs	5
fecr_stan	6
fecr_stanExtra	8
fecr_stanSimple	11
fec_stan	13
getPrior_delta	15
getPrior_mu	16
plotCounts	17
simData1s	18
simData2s	19
stan2mcmc	20

Index 22

eggCounts-package *Hierarchical modelling of faecal egg counts*

Description

This package implements Bayesian hierarchical models for the analysis of faecal egg count data. Bayesian inference is done via efficient MCMC sampling using Stan. Additional models are available externally for handling FECs with potential outliers or bi-modality. The models are in **eggCountsExtra** package hosted on Github.

Details

Package: eggCounts
 Type: Package
 Version: 2.1-2
 Date: 2018-11-07
 License: GPL (>= 3)
 LazyLoad: yes

Author(s)

Craig Wang <craig.wang@uzh.ch>
Michaela Paul

Examples

```
## Not run:  
  
## Citations  
citation('eggCounts')  
  
## History of changes  
file.show(system.file("NEWS", package = "eggCounts"))  
  
## Demonstration  
demo("fecm_stan", package = "eggCounts")  
  
## Install eggCountsExtra  
devtools::install_github("CraigWangUZH/eggCountsExtra")  
  
## End(Not run)
```

eggs

Faecal egg count samples (before and after treatment)

Description

This is an example data set containing 14 eggs per gram (epg) values in sheep before and after anthelmintic treatment of benzimidazole. The correction factor of the diagnostic technique was 50.

Usage

```
data(eggs)
```

Format

A dataframe containing 14 observations.

References

Craig Wang, Paul R. Torgerson, Johan Høglund, Reinhard Furrer, Zero-inflated hierarchical models for faecal egg counts to assess anthelmintic efficacy, *Veterinary Parasitology*, Volume 235, 2017, Pages 20-28.

 fecrtCI

Compute standard FECRT according to WAAVP guidelines

Description

Computes the standard Faecal Egg Count Reduction Test together with approximate confidence interval according to the WAAVP guidelines (Coles et al., 1992, 2006). The function also returns bootstrap confidence intervals.

Usage

```
fecrtCI(egp1, egp2, paired = FALSE, alpha = 0.05, R = 1999)
```

Arguments

egp1	faecal egg counts in untreated animals
egp2	faecal egg counts in treated animals
paired	logical. If TRUE, indicates the samples are paired. Otherwise they are unpaired.
alpha	confidence level of the interval
R	number of bootstrap replicates

Value

A list with	
estimate	the estimated percentage reduction in mean epg
bootCI	corresponding bootstrap confidence interval
approxCI	corresponding approximate confidence interval

Author(s)

Michaela Paul

References

Coles GC, Bauer C, Borgsteede FHM, Geerts S, Klei TR, Taylor MA, Waller, PJ (1992). World Association for the Advancement of Veterinary Parasitology (WAAVP) methods for the detection of anthelmintic resistance in nematodes of veterinary importance, *Veterinary Parasitology*, 44:35-44.

Coles GC, Jackson F, Pomroy WE, Prichard RK, von Samson-Himmelstjerna G, Silvestre A, Taylor MA, Vercruysse J (2006). The detection of anthelmintic resistance in nematodes of veterinary importance, *Veterinary Parasitology*, 136:167-185.

Examples

```
data(eggs)
fecrtCI(eggs$before, eggs$after, paired=TRUE)
```

fecr_probs	<i>Compute the probability of the reduction parameter relative to the given threshold</i>
------------	---

Description

Computes the probability of the reduction parameter's marginal posterior density relative to a threshold.

Usage

```
fecr_probs(stanFit, threshold = 0.95, lessthan = TRUE,
           plot = TRUE, xlab, ylab, main, verbose = TRUE, ...)
```

Arguments

stanFit	A stanfit object from the output of fecr_stan()
threshold	numeric. The default threshold is 0.95 or 95%.
lessthan	logical. If TRUE, the probability less than the threshold is computed. Otherwise greater or equal to the threshold is computed. Default is TRUE.
plot	logical. If TRUE, the posterior density of the reduction is plotted with region less than the threshold shaded.
xlab, ylab, main	Arguments for plotting. Only used if showPlot = TRUE.
verbose	logical, If TRUE, a statement with computed probability is printed out.
...	Additional plotting arguments.

Value

Returns a numeric value indicating the probability in percentage.

Author(s)

Craig Wang

Examples

```
## Not run:
## load sample data
data(eggs)

## apply zero-inflation model to the data vector
model <- fecr_stan(eggs$before, eggs$after, rawCounts=FALSE, preCF=10,
                  paired=TRUE, zeroInflation=TRUE)
fecr_probs(model$stan.samples)

## End(Not run)
```

fecr_stan *Modelling the reduction of faecal egg count data*

Description

Models the reduction in faecal egg counts data with analytical sensitivity-adjusted (un)paired (zero-inflated) Poisson-gamma models (with individual efficacy) using Stan modelling language.

It is computationally several-fold faster compared to conventional MCMC techniques. For the installation instruction of Stan, please read <https://github.com/stan-dev/rstan/wiki/RStan-Getting-Started>.

Usage

```
fecr_stan(preFEC, postFEC, rawCounts = FALSE, preCF = 50, postCF = preCF,
paired = TRUE, indEfficacy = TRUE, zeroInflation = FALSE,
muPrior, kappaPrior, deltaPrior, phiPrior, deltakappaPrior,
nsamples = 2000, nburnin = 1000, thinning = 1, nchain = 2,
ncore = 1, adaptDelta = 0.95, saveAll = FALSE, verbose = FALSE)
```

Arguments

preFEC	vector of pre-treatment faecal egg counts
postFEC	vector of post-treatment faecal egg counts
rawCounts	logical. If TRUE, preFEC and postFEC correspond to raw counts (as counted on equipment). Otherwise they correspond to calculated eggs (raw counts times correction factor). Defaults to FALSE.
preCF	a positive integer or a vector of positive integers. Pre-treatment correction factor(s)
postCF	a positive integer or a vector of positive integers. Post-treatment correction factor(s)
paired	logical. If TRUE, uses the model for the paired design. Otherwise uses the model for the unpaired design
indEfficacy	logical. If TRUE, uses the paired model allowing for individual efficacy. Only use in combination with paired = TRUE and zeroInflation = FALSE
zeroInflation	logical. If TRUE, uses the model with zero-inflation. Otherwise uses the model without zero-inflation
muPrior	a list with hyper-prior information for the group mean epg parameter μ . The default prior is <code>list(priorDist = "gamma", hyperpars=c(1,0.001))</code> , i.e. a gamma distribution with shape 1 and rate 0.001, its 90% probability mass lies between 51 and 2996
kappaPrior	a list with hyper-prior information for the group dispersion parameter κ . The default prior is <code>list(priorDist = "gamma", hyperpars=c(1,0.7))</code> , i.e. a gamma distribution with shape 1 and rate 0.7, its 90% probability mass lies between 0.1 and 4.3 with a median of 1

deltaPrior	a list with hyper-prior information for the reduction parameter δ . The default prior is <code>list(priorDist = "beta", hyperpars=c(1,1))</code> , i.e. a uniform prior between 0 and 1
phiPrior	a list with hyper-prior information for the zero-inflation parameter ϕ . The default prior is <code>list(priorDist = "beta", hyperpars=c(1,1))</code> , i.e. a uniform prior between 0 and 1
deltakappaPrior	a list with hyper-prior information for the shape parameter of reduction δ_κ . Only used if <code>indEfficacy = TRUE</code> . The default prior is <code>list(priorDist = "normal", hyperpars=c(2,1))</code>
nsamples	a positive integer specifying the number of samples for each chain (including burn-in samples)
nburnin	a positive integer specifying the number of burn-in samples
thinning	a positive integer specifying the thinning parameter, the period for saving samples
nchain	a positive integer specifying the number of chains
ncore	a positive integer specifying the number of cores to use when executing the chains in parallel
adaptDelta	the target acceptance rate, a numeric value between 0 and 1
saveAll	logical. If TRUE, posterior samples for all parameters are saved in the <code>stanfit</code> object. If FALSE, only samples for δ , μ , κ and ϕ are saved. Default to FALSE.
verbose	logical. If TRUE, prints progress and debugging information

Details

The first time each model with non-default priors is applied, it can take up to 20 seconds for Stan to compile the model. Currently the function only support prior distributions with two parameters. For a complete list of supported priors and their parameterization, please consult the list of distributions in Stan <http://mc-stan.org/documentation/>.

The default number of samples per chain is 2000, with 1000 burn-in samples. Normally this is sufficient in Stan. If the chains do not converge, one should tune the MCMC parameters until convergence is reached to ensure reliable results.

Value

Prints out the posterior summary of FECR as the reduction, `meanEPG.untreated` as the mean pre-treatment `epg`, and `meanEPG.treated` as the mean after-treatment `epg`. The posterior summary contains the mean, standard deviation (`sd`), 2.5%, 50% and 97.5% percentiles, the 95% highest posterior density interval (`HPDLow95` and `HPDHigh95`) and the posterior mode.

NOTE: we recommend to use the 95% equal-tailed credible interval and the median as summary statistics of reduction for the individual efficacy model. For all other models, we recommend to use the 95% HPD interval and the mode.

The returned value is a list that consists of:

<code>stan.samples</code>	An object of S4 class <code>stanfit</code> representing the fitted results. For more information, please see the <code>stanfit-class</code> in <code>rstan</code> reference manual.
<code>posterior.summary</code>	A data frame that is the same as the printed posterior summary.

Author(s)

Craig Wang

References

Individual efficacy models: Craig Wang, Paul R. Torgerson, Ray M. Kaplan, Melissa M. George, Reinhard Furrer. (2018) Modelling anthelmintic resistance by extending eggCounts package to allow individual efficacy, International Journal for Parasitology: Drugs and Drug Resistance, Volume 8, Pages 386-393. <https://doi.org/10.1016/j.ijpddr.2018.07.003>

Zero-inflation models: Craig Wang, Paul R. Torgerson, Johan Hoggund, Reinhard Furrer. (2017) Zero-inflated hierarchical models for faecal egg counts to assess anthelmintic efficacy, Veterinary Parasitology, Volume 235, Pages 20-28. <http://dx.doi.org/10.1016/j.vetpar.2016.12.007>

Other models: Paul R. Torgerson, Michaela Paul, Reinhard Furrer. (2014) Evaluating faecal egg count reduction using a specifically designed package 'eggCounts' in R and a user friendly web interface, International Journal for Parasitology, Volume 44, Pages 299-303. <http://dx.doi.org/10.1016/j.ijpara.2014.01.005>

See Also

[simData2s](#) for simulating faecal egg counts data with two samples

Examples

```
## Not run:
## load sample data
data(eggs)

## apply paired model with individual efficacy
model <- fecr_stan(eggs$before, eggs$after, rawCounts=FALSE, preCF=50,
                  paired=TRUE, indEfficacy = TRUE)
samples <- stan2mcmc(model$stan.samples)

## End(Not run)
```

 fecr_stanExtra

Modelling the reduction of faecal egg count data using custom models

Description

Models the reduction in faecal egg counts data with custom model formulation using Stan modelling language.

It is computationally several-fold faster compared to conventional MCMC techniques. For the installation instruction of Stan, please read <https://github.com/stan-dev/rstan/wiki/RStan-Getting-Started>.

Usage

```
fecr_stanExtra(preFEC, postFEC, rawCounts = FALSE, preCF = 50, postCF = preCF,
  modelName = NULL, modelCode = NULL, modelFile = NULL, modelData = NULL,
  nsamples = 2000, nburnin = 1000, thinning = 1, nchain = 2,
  ncore = 1, adaptDelta = 0.95, verbose = FALSE)
```

Arguments

preFEC	vector of pre-treatment faecal egg counts. Not required if modelCode or modelFile is supplied.
postFEC	vector of post-treatment faecal egg counts. Not required if modelCode or modelFile is supplied.
rawCounts	logical. If TRUE, preFEC and postFEC correspond to raw counts (as counted on equipment). Otherwise they correspond to calculated eggs (raw counts times correction factor). Defaults to FALSE. Not required if modelCode or modelFile is supplied.
preCF	a positive integer or a vector of positive integers. Pre-treatment correction factor(s). Not required if modelCode or modelFile is supplied.
postCF	a positive integer or a vector of positive integers. Post-treatment correction factor(s). Not required if modelCode or modelFile is supplied.
modelName	string. One of four available models ("Po", "UPo", "ZIPo", "ZIUPo") from eggCountsExtra package, which corresponds to outlier-adjusted version of paired, unpaired, paired with zero inflation and unpaired with zero inflation models. Not required if modelCode or modelFile is supplied.
modelCode	stan model code. Not required when modelName or modelFile is supplied.
modelFile	stan model file with file extension '*.stan'. Not required when modelName or modelCode is supplied.
modelData	stan data list. A named list or environment providing the data for the model, or a character vector for all the names of objects used as data. Not required when modelName is supplied.
nsamples	a positive integer specifying the number of samples for each chain (including burn-in samples)
nburnin	a positive integer specifying the number of burn-in samples
thinning	a positive integer specifying the thinning parameter, the period for saving samples
nchain	a positive integer specifying the number of chains
ncore	a positive integer specifying the number of cores to use when executing the chains in parallel
adaptDelta	the target acceptance rate, a numeric value between 0 and 1
verbose	logical. If TRUE, prints progress and debugging information

Details

If `modelName` is one of `c("Po", "UPo", "ZIPo", "ZIUPo")`, then outlier-adjusted models are used.

- In paired models, outliers are those counts with `postFEC > preFEC`. Outlier weights are assigned as the inverse of `postFEC/preFEC`,
- In unpaired models, outliers are those counts with `postFEC` greater than the 95th percentile of a Poisson distribution, where the Poisson mean is computed based on the mean of `postFEC` excluding `postFEC > Q3 + 1.5*IQR`. `Q3` is the 75th percentile and `IQR` is the interquartile range. The lowest outlier weight is assigned as 0.01, and other outliers assigned proportionally.
- In both cases, non-outliers are assigned with outlier weight = 1.

The first time each model is applied, it can take up to 20 seconds for Stan to compile the model.

The default number of samples per chain is 2000, with 1000 burn-in samples. Normally this is sufficient in Stan. If the chains do not converge, one should tune the MCMC parameters until convergence is reached to ensure reliable results.

Value

Prints out the posterior summary of FECR as the reduction, `meanEPG.untreated` as the mean pre-treatment `epg`, and `meanEPG.treated` as the mean after-treatment `epg`. The posterior summary contains the mean, standard deviation (`sd`), 2.5%, 50% and 97.5% percentiles, the 95% highest posterior density interval (`HPDLow95` and `HPDHigh95`) and the posterior mode.

The returned value is a list that consists of:

<code>stan.model</code>	A object of class <code>stanmodel</code> that was used.
<code>stan.samples</code>	An object of S4 class <code>stanfit</code> representing the fitted results. For more information, please see the <code>stanfit-class</code> in <code>rstan</code> reference manual.
<code>posterior.summary</code>	A data frame that is the same as the printed posterior summary, not available for custom models.

Author(s)

Craig Wang

Examples

```
## Not run:
library(eggCountsExtra)
data(eggs) ## load sample data

## apply paired model with outliers
model1 <- fecr_stanExtra(eggs$before, eggs$after, rawCounts=FALSE,
  preCF=10, modelName = "Po")
samples <- stan2mcmc(model1$stan.samples)
fecr_probs(model1$stan.samples, threshold = 0.99)

## apply a very simple custom model
code <- "data{
```

```

    int J; // number of animals
    int y_before[J]; // after treatment McMaster count
    int y_after[J]; // before treatment McMaster count
  }
  parameters{
    real<lower=0> mu;
    real<lower=0,upper=1> delta;
  }
  model{
    mu ~ gamma(1,0.001);
    delta ~ beta(1,1);
    y_before ~ poisson(mu);
    y_after ~ poisson(mu*delta);
  }"

  dat <- list(J = nrow(eggs), y_before = eggs$before,
             y_after = eggs$after)
  model2 <- fecr_stanExtra(modelCode = code, modelData = dat)

  ## End(Not run)

```

fecr_stanSimple	<i>Modelling the reduction of faecal egg count data using a simple Bayesian model</i>
-----------------	---

Description

Models the reduction in faecal egg counts data with simple Bayesian model formulation using Stan modelling language. The model is for paired design only, and it assumes analytical sensitivity adjusted Poisson distribution for the observed egg counts.

It is computationally several-fold faster compared to conventional MCMC techniques. For the installation instruction of Stan, please read: [Stan Installation](#).

Usage

```

fecr_stanSimple(preFEC, postFEC, rawCounts = FALSE, preCF = 50, postCF = preCF,
  muPrior, deltaPrior, nsamples = 2000, nburnin = 1000, thinning = 1, nchain = 2,
  ncore = 1, adaptDelta = 0.95, saveAll = FALSE, verbose = FALSE)

```

Arguments

preFEC	vector of pre-treatment faecal egg counts
postFEC	vector of post-treatment faecal egg counts
rawCounts	logical. If TRUE, preFEC and postFEC correspond to raw counts (as counted on equipment). Otherwise they correspond to calculated eggs (raw counts times correction factor). Defaults to FALSE.
preCF	a positive integer or a vector of positive integers. Pre-treatment correction factor(s)

postCF	a positive integer or a vector of positive integers. Post-treatment correction factor(s)
muPrior	a list with hyper-prior information for the group mean epg parameter μ . The default prior is <code>list(priorDist = "gamma", hyperpars=c(1, 0.001))</code> , i.e. a gamma distribution with shape 1 and rate 0.001, its 90% probability mass lies between 51 and 2996
deltaPrior	a list with hyper-prior information for the reduction parameter δ . The default prior is <code>list(priorDist = "beta", hyperpars=c(1, 1))</code> , i.e. a uniform prior between 0 and 1
nsamples	a positive integer specifying the number of samples for each chain (including burn-in samples)
nburnin	a positive integer specifying the number of burn-in samples
thinning	a positive integer specifying the thinning parameter, the period for saving samples
nchain	a positive integer specifying the number of chains
ncore	a positive integer specifying the number of cores to use when executing the chains in parallel
adaptDelta	the target acceptance rate, a numeric value between 0 and 1
saveAll	logical. If TRUE, posterior samples for all parameters are saved in the <code>stanfit</code> object. If FALSE, only samples for δ , μ , κ and ϕ are saved. Default to FALSE.
verbose	logical. If TRUE, prints progress and debugging information

Details

The first time each model with non-default priors is applied, it can take up to 20 seconds for Stan to compile the model. Currently the function only support prior distributions with two parameters. For a complete list of supported priors and their parameterization, please consult the list of distributions in [Stan](#).

The default number of samples per chain is 2000, with 1000 burn-in samples. Normally this is sufficient in Stan. If the chains do not converge, one should tune the MCMC parameters until convergence is reached to ensure reliable results.

Value

Prints out the posterior summary of FECR as the reduction, `meanEPG.untreated` as the mean pre-treatment epg, and `meanEPG.treated` as the mean after-treatment epg. The posterior summary contains the mean, standard deviation (sd), 2.5%, 50% and 97.5% percentiles, the 95% highest posterior density interval (HPDLow95 and HPDHigh95) and the posterior mode.

NOTE: we recommend to use the 95% equal-tailed credible interval and the median as summary statistics of reduction for the individual efficacy model. For all other models, we recommend to use the 95% HPD interval and the mode.

The returned value is a list that consists of:

<code>stan.samples</code>	An object of S4 class <code>stanfit</code> representing the fitted results. For more information, please see the stanfit-class in rstan reference manual.
<code>posterior.summary</code>	A data frame that is the same as the printed posterior summary.

Author(s)

Tea Isler Craig Wang

See Also[simData2s](#) for simulating faecal egg counts data with two samples**Examples**

```
## Not run:
## load sample data
data(eggs)

## apply paired model with individual efficacy
model <- fecr_stanSimple(eggs$before, eggs$after, rawCounts=FALSE, preCF=10)
samples <- stan2mcmc(model$stan.samples)

## End(Not run)
```

fec_stan

*Modelling of faecal egg count data (one-sample case)***Description**

Models faecal egg counts data in a one-sample case with (zero-inflated) Poisson-gamma model formulation using Stan modelling language. It is computationally several-fold faster compared to conventional MCMC techniques. For the installation instruction of Stan, please read <https://github.com/stan-dev/rstan/wiki/RStan-Getting-Started>.

Usage

```
fec_stan(fec, rawCounts = FALSE, CF = 50, zeroInflation = TRUE,
         muPrior, kappaPrior, phiPrior,
         nsamples = 2000, nburnin = 1000, thinning = 1, nchain = 2,
         ncore = 1, adaptDelta = 0.95, saveAll = FALSE, verbose = FALSE)
```

Arguments

fec	vector of faecal egg counts
rawCounts	logical. If true, preFEC and postFEC correspond to raw counts (as counted on equipment). Otherwise they correspond to calculated eggs (raw counts times correction factor). Defaults to FALSE.
CF	a positive integer or a vector of positive integers. Correction factor(s)
zeroInflation	logical. If true, uses the model with zero-inflation. Otherwise uses the model without zero-inflation

muPrior	a list with hyper-prior information for the group mean epg parameter μ . The default prior is <code>list(priorDist = "gamma", hyperpars=c(1, 0.001))</code> , i.e. a gamma distribution with shape 1 and rate 0.001, its 90% probability mass lies between 51 and 2996
kappaPrior	a list with hyper-prior information for the group dispersion parameter κ . The default prior is <code>list(priorDist = "gamma", hyperpars=c(1, 0.7))</code> , i.e. a gamma distribution with shape 1 and rate 0.7, its 90% probability mass lies between 0.1 and 4.3 with a median of 1
phiPrior	a list with hyper-prior information for zero-inflation parameter. The default prior is <code>list(priorDist = "beta", hyperpars=c(1, 1))</code> , i.e. a uniform prior between 0 and 1
nsamples	a positive integer specifying the number of samples for each chain (including burn-in samples)
nburnin	a positive integer specifying the number of burn-in samples
thinning	a positive integer specifying the thinning parameter, the period for saving samples
nchain	a positive integer specifying the number of chains
ncore	a positive integer specifying the number of cores to use when executing the chains in parallel
adaptDelta	the target acceptance rate, a numeric value between 0 and 1
saveAll	logical. If TRUE, posterior samples for all parameters are saved in the <code>stanfit</code> object. If FALSE, only samples for μ , κ and ϕ are saved. Default to FALSE.
verbose	logical. If true, prints progress and debugging information

Details

The first time each non-default model is applied, it can take up to 20 seconds for Stan to compile the model. Currently the function only support prior distributions with two parameters. For a complete list of supported priors and their parameterization, please consult the list of distributions in Stan <http://mc-stan.org/documentation/>.

The default number of samples per chain is 2000, with 1000 burn-in samples. Normally this is sufficient in Stan. If the chains do not converge, one should tune the MCMC parameters until convergence is reached to ensure reliable results.

Value

Prints out summary of meanEPG as the posterior mean epg. The posterior summary contains the mean, standard deviation (sd), 2.5%, 50% and 97.5% percentiles, the 95% highest posterior density interval (HPDLow95 and HPDHigh95) and the posterior mode. NOTE: we recommend to use the 95% HPD interval and the mode for further statistical analysis.

The returned value is a list that consists of:

<code>stan.samples</code>	An object of S4 class <code>stanfit</code> representing the fitted results. For more information, please see the stanfit-class in rstan reference manual.
<code>posterior.summary</code>	A data frame that is the same as the printed posterior summary.

Author(s)

Craig Wang

See Also[simData1s](#) for simulating faecal egg count data with one sample**Examples**

```
## Not run:
## load the sample data
data(eggs)

## apply zero-inflation model
model <- fec_stan(eggs$before, rawCounts=FALSE, CF=50)
samples <- stan2mcmc(model$stan.samples)

## End(Not run)
```

`getPrior_delta`*Get prior parameters from Beta distribution*

Description

Compute the shape parameters from a Beta distribution for μ based on some prior belief.

Usage

```
getPrior_delta(lower, upper, p = 0.7, mode, conc, plot = TRUE)
```

Arguments

<code>lower, upper, p</code>	numeric. The lower and upper threshold where p probability density mass lies in a beta distribution.
<code>mode, conc</code>	numeric. The mode and concentration parameters of a beta distribution. Higher concentration indicates smaller variance.
<code>plot</code>	logical. If TRUE, the prior distribution is plotted after parameters are found.

Details

`multiroot` function from **rootSolve** package is used to compute the parameters.

Value

Returns Gamma prior parameters for μ .

Author(s)

Tea Isler
Craig Wang

Examples

```
getPrior_delta(lower = 0.6, upper = 0.9, p = 0.8)
```

getPrior_mu

Get prior parameters from Gamma distribution

Description

Compute the shape and rate parameters from a Gamma distribution for μ based on some prior belief about its cumulative distribution function.

Usage

```
getPrior_mu(x, px, y, py, s1 = 1, s2 = 0.001, plot = TRUE)
```

Arguments

x, px, y, py	numeric. Threshold of some prior belief about true epg. There is px probability that the true epg is below x, and there is py probability that the true epg is below y
s1, s2	numeric. Starting values.
plot	logical. If TRUE, the prior distribution is plotted after parameters are found.

Details

multiroot function from **rootSolve** package is used to compute the parameters.

Value

Returns Gamma prior parameters for μ .

Author(s)

Tea Isler
Craig Wang

Examples

```
getPrior_mu(x = 200, px = 0.3, y = 500, py = 0.8)
```

plotCounts	<i>Plot faecal egg count data</i>
------------	-----------------------------------

Description

Plot egg count data to reflect changes between before and after treatment.

Usage

```
plotCounts(data, paired=TRUE, points=TRUE, points.method="jitter",  
           xlabel="", ylabel="Faecal egg counts [epg]", ...)
```

Arguments

data	a data frame with two columns, the first column is before treatment counts, the second column is after treatment counts.
paired	logical. If true, uses the plot for the paired design. Otherwise uses the plot for the unpaired design.
points	logical. If true, add individual points for unpaired plot. Omitted if paired is TRUE.
points.method	a quoted keyword to be used to separate coincident points if points is TRUE. The default method "overplot" causes such points to be overplotted, but it is also possible to specify "jitter" to jitter the points, or "stack" have coincident points stacked.
xlabel	label of x-axis.
ylabel	label of y-axis.
...	Additional arguments for function xyplot if paired is TRUE, for function boxplot otherwise.

Details

For paired data, a xyplot is used. For unpaired data, a grouped boxplot is used.

Value

A plot is returned based on the arguments.

Author(s)

Craig Wang

Examples

```
data(eggs)  
plotCounts(eggs[,c("before", "after")], paired = TRUE)
```

simData1s *Simulate faecal egg count data (1-sample situation)*

Description

Simulates (zero-inflated) egg count data

Usage

```
simData1s(n = 10, mean = 500, kappa = 0.5, phi = 1,
          f = 50, rounding = TRUE, seed = NULL)
```

Arguments

n	sample size (number of faeces collected)
mean	true number of eggs per gram (epg)
kappa	overdispersion parameter, $\kappa \rightarrow \infty$ corresponds to Poisson distribution
phi	prevalence, i.e. proportion of infected animals, between 0 and 1
f	correction factor of the egg counting technique, either an integer or a vector of integers with length n
rounding	logical. If TRUE, the Poisson mean for the raw counts is rounded. The rounding applies since the mean epg is frequently reported as an integer value. For more information, please see Details.
seed	an integer that will be used in a call to set.seed before simulation. If NULL, a random seed is allocated.

Details

In the simulation of raw (master) counts, it follows a Poisson distribution with some mean. The mean is frequently rounded down if it has a very low value and `rounding = TRUE`, hence there expects to be a negative bias overall when $\mu < 150$. Set `rounding = FALSE` if one does not wish to have any bias in the simulated counts.

Value

A data.frame with three columns, namely the observed epg (obs), number of eggs counted on microscope slide (master) and true epg in the sample(true).

Author(s)

Craig Wang
Michaela Paul

See Also

[fec_stan](#) for analyzing faecal egg count data with one sample

Examples

```
fec <- simData1s(n=10, mean=500, kappa=0.5, phi=0.7)
```

simData2s	<i>Simulate faecal egg count data (2-sample situation)</i>
-----------	--

Description

Generates two samples of (zero-inflated) egg count data

Usage

```
simData2s(n = 10, preMean = 500, delta = 0.1, kappa = 0.5,
  deltaShape = NULL, phiPre = 1, phiPost = phiPre, f = 50,
  paired = TRUE, rounding = TRUE, seed = NULL)
```

Arguments

n	sample size (number of animals)
preMean	true pre-treatment epg
delta	proportion of epg left after treatment, between 0 and 1. $1 - \delta$ is reduction in mean after treatment, $\delta = 0.1$ indicates a 90% reduction
kappa	overdispersion parameter, $\kappa \rightarrow \infty$ corresponds to Poisson distribution
deltaShape	shape parameter for the distribution of reductions. If NULL, the same reduction is applied to the latent true epg of each animal.
phiPre	pre-treatment prevalence (i.e. proportion of infected animals), between 0 and 1
phiPost	post-treatment prevalence, between 0 and 1
f	correction factor of the egg counting technique, either an integer or a vector of integers with length n
paired	logical. If TRUE, paired samples are simulated. Otherwise unpaired samples are simulated.
rounding	logical. If TRUE, the Poisson mean for the raw counts is rounded. The rounding applies since the mean epg is frequently reported as an integer value. For more information, please see Details.
seed	an integer that will be used in a call to set.seed before simulation. If NULL, a random seed is allocated.

Details

In the simulation of raw (master) counts, it follows a Poisson distribution with some mean. The mean is frequently rounded down if it has a very low value and `rounding = TRUE`, there expects to be a up to 3-10% positive bias in the mean reduction when $\mu < 150$ and $\delta < 0.1$. Set `rounding = FALSE` if one does not wish to have any bias.

Value

A data.frame with six columns, namely the observed epg (obs), number of eggs counted on microscope slide (master) and true epg in the sample (true) for both pre- and post- treatment.

Author(s)

Craig Wang
Michaela Paul

See Also

[fecr_stan](#) for analyzing faecal egg count data with two samples

Examples

```
fec <- simData2s(n=10, preMean=500, delta=0.1, kappa=0.5)

## show the positive bias when the true reduction should be 95%
set.seed(1)
fec <- simData2s(n=1e5, preMean=150, delta=0.05, kappa=0.5)
1-mean(fec$masterPost)/mean(fec$masterPre)
```

stan2mcmc

Convert a Stanfit object to MCMC object

Description

Converts a stanfit object into a MCMC object for easier analysis.

Usage

```
stan2mcmc(stanFit)
```

Arguments

stanFit A stanfit object from the output of either fecr_stan() or fec_stan()

Details

The output can be analyzed as a typical MCMC object with the functions from the coda package.
NOTE: The resulting MCMC object does not contain warm-up samples and is already thinned.

Value

A MCMC object with a list of relevant parameters.

Author(s)

Craig Wang

Examples

```
## Not run:
data(eggs)

## apply zero-inflation model for the paired design
model <- fecr_stan(eggs$before, eggs$after, rawCounts = FALSE, indEfficacy = FALSE,
                  preCF = 10, paired = TRUE, zeroInflation = TRUE)
samples <- stan2mcmc(model$stan.samples)
summary(samples)

## End(Not run)
```

Index

*Topic **datasets**

eggs, [3](#)

*Topic **modelling**

fec_stan, [13](#)

fecr_probs, [5](#)

fecr_stan, [6](#)

fecr_stanExtra, [8](#)

fecr_stanSimple, [11](#)

fecrtCI, [4](#)

stan2mcmc, [20](#)

*Topic **package**

eggCounts-package, [2](#)

*Topic **simulation**

simData1s, [18](#)

simData2s, [19](#)

boxplot, [17](#)

eggCounts (eggCounts-package), [2](#)

eggCounts-package, [2](#)

eggs, [3](#)

fec_stan, [13](#), [18](#)

fecr_probs, [5](#)

fecr_stan, [6](#), [20](#)

fecr_stanExtra, [8](#)

fecr_stanSimple, [11](#)

fecrtCI, [4](#)

getPrior_delta, [15](#)

getPrior_mu, [16](#)

plotCounts, [17](#)

simData1s, [15](#), [18](#)

simData2s, [8](#), [13](#), [19](#)

stan2mcmc, [20](#)

xyplot, [17](#)