

# Package: eDNAjoint (via r-universe)

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**Title** Joint Modeling of Traditional and Environmental DNA Survey Data

**Version** 0.1

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**Description** Models integrate environmental DNA (eDNA) detection data and traditional survey data to jointly estimate species catch rate (see package vignette: [https://bookdown.org/abigailkeller/eDNAjoint\\_vignette/](https://bookdown.org/abigailkeller/eDNAjoint_vignette/)). Models can be used with count data via traditional survey methods (i.e., trapping, electrofishing, visual) and replicated eDNA detection/nondetection data via polymerase chain reaction (i.e., PCR or qPCR) from multiple survey locations. Estimated parameters include probability of a false positive eDNA detection, a site-level covariates that scale the sensitivity of eDNA surveys relative to traditional surveys, and catchability coefficients for traditional gear types. Models are implemented with a Bayesian framework (Markov chain Monte Carlo) using the 'Stan' probabilistic programming language.

**License** GPL-3

**URL** <https://github.com/abigailkeller/eDNAjoint>

**BugReports** <https://github.com/abigailkeller/eDNAjoint/issues>

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` ` srr::srr_stats_roclet"))`

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RcppParallel (>= 5.0.1), rstan (>= 2.26.23), StanHeaders (>= 2.26.22)

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**Repository** <https://ropensci.r-universe.dev>

**RemoteUrl** <https://github.com/ropensci/eDNAjoint>

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eDNAjoint-package      *The 'eDNAjoint' package.*

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## Description

Models integrate environmental DNA (eDNA) detection data and traditional survey data to jointly estimate species catch rate (see **Package Vignette**). Models can be used with count data via traditional survey methods (i.e., trapping, electrofishing, visual) and replicated eDNA detection/nondetection data via polymerase chain reaction (i.e., PCR or qPCR) from multiple survey locations. Estimated parameters include probability of a false positive eDNA detection, a site-level covariates that scale the sensitivity of eDNA surveys relative to traditional surveys, and catchability coefficients for traditional gear types. Models are implemented with a Bayesian framework (Markov chain Monte Carlo) using the 'Stan' probabilistic programming language.

**Author(s)**

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- Ryan P. Kelly <rpkelley@uw.edu> [contributor]

**References**

Stan Development Team (NA). RStan: the R interface to Stan. <https://mc-stan.org>

**See Also**

Useful links:

- <https://github.com/abigailkeller/eDNAjoint>
- Report bugs at <https://github.com/abigailkeller/eDNAjoint/issues>

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detectionCalculate	<i>Calculate the survey effort necessary to detect species presence, given the species expected catch rate.</i>
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**Description**

This function calculates the number of survey effort units necessary to detect species presence using median estimated parameter values from `jointModel()`. Detecting species presence is defined as producing at least one true positive eDNA detection or catching at least one individual. See more examples in the [Package Vignette](#).

**Usage**

```
detectionCalculate(modelfit, mu, cov.val = NULL, probability = 0.9, qPCR.N = 3)
```

**Arguments**

<code>modelfit</code>	An object of class <code>stanfit</code> .
<code>mu</code>	A numeric vector of species densities/capture rates. If multiple traditional gear types are represented in the model, <code>mu</code> is the catch rate of gear type 1.
<code>cov.val</code>	A numeric vector indicating the values of site-level covariates to use for prediction. Default is <code>NULL</code> .
<code>probability</code>	A numeric value indicating the probability of detecting presence. The default is 0.9.
<code>qPCR.N</code>	An integer indicating the number of qPCR replicates per eDNA sample. The default is 3.

**Value**

A summary table of survey efforts necessary to detect species presence, given  $\mu$ , for each survey type.

**Note**

Before fitting the model, this function checks to ensure that the function is possible given the inputs. These checks include:

- Input model fit is an object of class 'stanfit'.
- Input  $\mu$  is a numeric vector.
- Input probability is a univariate numeric value.
- If model fit contains alpha, cov.val must be provided.
- Input cov.val is numeric.
- Input cov.val is the same length as the number of estimated covariates.
- Input model fit has converged (i.e. no divergent transitions after warm-up).

If any of these checks fail, the function returns an error message.

**Examples**

```
# Ex. 1: Calculating necessary effort for detection with site-level
# covariates

# Load data
data(gobyData)

# Fit a model including 'Filter_time' and 'Salinity' site-level covariates
fit.cov <- jointModel(data = gobyData, cov = c('Filter_time','Salinity'),
  family = "poisson", p10priors = c(1,20), q = FALSE,
  multicore = FALSE)

# Calculate at the mean covariate values
# (covariates are standardized, so mean = 0)
detectionCalculate(fit.cov$model, mu = seq(from = 0.1, to = 1, by = 0.1),
  cov.val = c(0,0), qPCR.N = 3)

# Calculate mu_critical at salinity 0.5 z-scores greater than the mean
detectionCalculate(fit.cov$model, mu = seq(from = 0.1, to = 1, by = 0.1),
  cov.val = c(0,0.5), qPCR.N = 3)

# Ex. 2: Calculating necessary effort for detection with multiple traditional
# gear types

# Load data
data(greencrabData)

# Fit a model with no site-level covariates
fit.q <- jointModel(data = greencrabData, cov = NULL, family = "negbin",
  p10priors = c(1,20), q = TRUE, multicore = FALSE)
```

```
# Calculate
detectionCalculate(fit.q$model, mu = seq(from = 0.1, to = 1, by = 0.1),
                  cov.val = NULL, qPCR.N = 3)

# Change probability of detecting presence to 0.95
detectionCalculate(fit.q$model, mu = 0.1, cov.val = NULL,
                  probability = 0.95, qPCR.N = 3)
```

---

detectionPlot	<i>Plot the survey effort necessary to detect species presence, given the species expected catch rate.</i>
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### Description

This function plots the number of survey effort units to necessary detect species presence, calculated using median estimated parameter values from `jointModel()`. Detecting species presence is defined as producing at least one true positive eDNA detection or catching at least one individual. See more examples in the [Package Vignette](#).

### Usage

```
detectionPlot(
  modelfit,
  mu.min,
  mu.max,
  cov.val = NULL,
  probability = 0.9,
  qPCR.N = 3
)
```

### Arguments

<code>modelfit</code>	An object of class <code>stanfit</code> .
<code>mu.min</code>	A value indicating the minimum expected species catch rate for plotting. If multiple traditional gear types are represented in the model, <code>mu</code> is the catch rate of gear type 1.
<code>mu.max</code>	A value indicating the maximum expected species catch rate for plotting. If multiple traditional gear types are represented in the model, <code>mu</code> is the catch rate of gear type 1.
<code>cov.val</code>	A numeric vector indicating the values of site-level covariates to use for prediction. Default is <code>NULL</code> .
<code>probability</code>	A numeric value indicating the probability of detecting presence. The default is 0.9.
<code>qPCR.N</code>	An integer indicating the number of qPCR replicates per eDNA sample. The default is 3.

**Value**

A plot displaying survey efforts necessary to detect species presence, given  $\mu$ , for each survey type.

**Note**

Before fitting the model, this function checks to ensure that the function is possible given the inputs. These checks include:

- Input model fit is an object of class 'stanfit'.
- Input mu.min is a numeric value greater than 0.
- Input mu.max is a numeric value.
- If model fit contains alpha, cov.val must be provided.
- Input cov.val is numeric.
- Input cov.val is the same length as the number of estimated covariates.
- Input probability is a univariate numeric value.
- Input model fit has converged (i.e. no divergent transitions after warm-up).

If any of these checks fail, the function returns an error message.

**Examples**

```
# Ex. 1: Calculating necessary effort for detection with site-level
# covariates

# Load data
data(gobyData)

# Fit a model including 'Filter_time' and 'Salinity' site-level covariates
fit.cov <- jointModel(data = gobyData, cov = c('Filter_time','Salinity'),
  family = "poisson", p10priors = c(1,20), q = FALSE,
  multicore = FALSE)

# Plot at the mean covariate values (covariates are standardized, so mean=0)
detectionPlot(fit.cov$model, mu.min = 0.1, mu.max = 1,
  cov.val = c(0,0), qPCR.N = 3)

# Calculate mu_critical at salinity 0.5 z-scores greater than the mean
detectionPlot(fit.cov$model, mu.min = 0.1, mu.max = 1, cov.val = c(0,0.5),
  qPCR.N = 3)

# Ex. 2: Calculating necessary effort for detection with multiple
# traditional gear types

# Load data
data(greencrabData)

# Fit a model with no site-level covariates
fit.q <- jointModel(data = greencrabData, cov = NULL, family = "negbin",
```

```
p10priors = c(1,20), q = TRUE,
multicore = FALSE)

# Calculate
detectionPlot(fit.q$model, mu.min = 0.1, mu.max = 1,
cov.val = NULL, qPCR.N = 3)

# Change probability of detecting presence to 0.95
detectionPlot(fit.q$model, mu.min = 0.1, mu.max = 1, cov.val = NULL,
probability = 0.95, qPCR.N = 3)
```

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gobyData

*gobyData*

---

## Description

gobyData

## Usage

gobyData

## Format

A list with four matrices representing eDNA sampling data (qPCR.N and qPCR.K), seine sampling data (count), and site-level covariate data (site.cov).

**qPCR.N** Total number of eDNA qPCR replicates at each site (row) and eDNA sample replicate (column). Data includes 39 total sites and a maximum of 22 eDNA sample replicates. NA indicates that fewer eDNA samples were collected than the maximum at a site.

**qPCR.K** Total number of positive eDNA qPCR detections at each site (row) and eDNA sample replicate (column). Data includes 39 total sites and a maximum of 22 eDNA sample replicates. NA indicates that fewer eDNA samples were collected than the maximum at a site.

**count** Count of goby individuals in seine samples at each site (row) and seine sample replicate (column). Data includes 39 total sites and a maximum of 22 seine replicates. NA indicates that fewer seine samples were collected than the maximum at a site.

**site.cov** Data representing site-level covariates at each site (row). Data includes mean salinity at a site ('Salinity'), mean time to filter eDNA samples ('Filter\_time'), density of other fish species ('Other\_fishes'), size of habitat ('Hab\_size'), and presence of vegetation ('Veg'). All non-integer covariate data is standardized.

## Source

<https://datadryad.org/stash/dataset/doi:10.5061/dryad.6rs23>

## References

Schmelzle, M.C. and Kinziger, A.P. (2016). Using occupancy modelling to compare environmental DNA to traditional field methods for regional-scale monitoring of an endangered aquatic species. *Molecular Ecology Resources*. 16(4): 895-908.

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greencrabData

*greencrabData*

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## Description

greencrabData

## Usage

greencrabData

## Format

A list with four matrices representing eDNA sampling data (qPCR.N and qPCR.K) and trap sampling data (count and count.type).

**qPCR.N** Total number of eDNA qPCR replicates at each site (row) and eDNA sample replicate (column). Data includes 20 total sites and 5 eDNA sample replicates.

**qPCR.K** Total number of positive eDNA qPCR detections at each site (row) and eDNA sample replicate (column). Data includes 20 total sites and 5 eDNA sample replicates.

**count** Count of green crab individuals in trap samples at each site (row) and trap sample replicate (column). Data includes 20 total sites and a maximum of 420 trap replicates. NA indicates that fewer trap samples were collected than the maximum at a site.

**count.type** Integer indicating the traditional gear type used at each site (row) and trap sample replicate (column). '1' refers to Fukui traps, and '2' refers to Minnow traps. Data includes 20 total sites and a maximum of 420 trap replicates. NA indicates that fewer trap samples were collected than the maximum at a site.

## Source

[doi.org/10.6084/m9.figshare.15117102.v2](https://doi.org/10.6084/m9.figshare.15117102.v2)

## References

Keller, A.G., Grason, E.W., McDonald, P.S., Ramon-Laca, A., Kelly, R.P. (2022). Tracking an invasion front with environmental DNA. *Ecological Applications*. 32(4): e2561. <https://doi.org/10.1002/eap.2561>



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jointModel	<i>Specify and fit joint model using count data from traditional surveys and eDNA qPCR data</i>
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## Description

This function implements a Bayesian model that integrates data from paired eDNA and traditional surveys, as described in Keller et al. (2022) <https://doi.org/10.1002/eap.2561>. The model estimates parameters including the expected species catch rate and the probability of false positive eDNA detection. This function allows for optional model variations, like inclusion of site-level covariates that scale the sensitivity of eDNA sampling relative to traditional sampling, as well as estimation of catchability coefficients when multiple traditional gear types are used. Model is implemented using Bayesian inference using the `rstan` package, which uses Hamiltonian Monte Carlo to simulate the posterior distributions. See more examples in the [Package Vignette](#).

## Usage

```
jointModel(
  data,
  cov = NULL,
  family = "poisson",
  p10priors = c(1, 20),
  q = FALSE,
  phipriors = NULL,
  multicore = TRUE,
  initial_values = NULL,
  n.chain = 4,
  n.iter.burn = 500,
  n.iter.sample = 2500,
  thin = 1,
  adapt_delta = 0.9,
  verbose = TRUE,
  seed = NULL
)
```

## Arguments

data	A list containing data necessary for model fitting. Valid tags are <code>qPCR.N</code> , <code>qPCR.K</code> , <code>count</code> , <code>count.type</code> , and <code>site.cov</code> . <code>qPCR.N</code> and <code>qPCR.K</code> are matrices or data frames with first dimension equal to the number of sites ( <i>i</i> ) and second dimension equal to the maximum number of eDNA samples at a given site ( <i>m</i> ). <code>qPCR.N</code> contains the total number of qPCR replicates per site and eDNA sample, and <code>qPCR.K</code> contains the total number of positive qPCR detections per site and eDNA sample. <code>count</code> is a matrix or data frame of traditional survey count data, with first dimension equal to the number of sites ( <i>i</i> ) and second dimension equal to the maximum number of traditional survey replicates at a given site ( <i>j</i> ). <code>count.type</code> is an optional matrix or data frame of integers indicating gear
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type used in corresponding count data, with first dimension equal to the number of sites ( $i$ ) and second dimension equal to the maximum number of traditional survey replicates at a given site. Values should be integers beginning with 1 (referring to the first gear type) to  $n$  (last gear type). `site.cov` is an optional matrix or data frame of site-level covariate data, with first dimension equal to the number of sites ( $i$ ). `site.cov` should include informative column names. Empty cells should be NA and will be removed during processing. Sites,  $i$ , should be consistent in all qPCR, count, and site covariate data.

<code>cov</code>	A character vector indicating the site-level covariates to include in the model. Default value is NULL.
<code>family</code>	The distribution class used to model traditional survey count data. Options include <code>poisson</code> ('poisson'), negative binomial ('negbin'), and <code>gamma</code> ('gamma'). Default value is 'poisson'.
<code>p10priors</code>	A numeric vector indicating beta distribution hyperparameters (alpha, beta) used as the prior distribution for the eDNA false positive probability ( $p_{10}$ ). Default vector is <code>c(1,20)</code> .
<code>q</code>	A logical value indicating whether to estimate a catchability coefficient, $q$ , for traditional survey gear types (TRUE) or to not estimate a catchability coefficient, $q$ , for traditional survey gear types (FALSE). Default value is FALSE.
<code>phipriors</code>	A numeric vector indicating gamma distribution hyperparameters (shape, rate) used as the prior distribution for $\phi$ , the overdispersion in the negative binomial distribution for traditional survey gear data. Used when <code>family = 'negbin'</code> . If <code>family = 'negbin'</code> , then default vector is <code>c(0.25,0.25)</code> , otherwise, default is NULL.
<code>multicore</code>	A logical value indicating whether to parallelize chains with multiple cores. Default is TRUE.
<code>initial_values</code>	A list of lists of initial values to use in MCMC. The length should equal the number of MCMC chains. Initial values can be provided for parameters: $\beta$ , $p_{10}$ (log-scale), $\mu$ , $q$ , $\alpha$ . If no initial values are provided, default random values are drawn.
<code>n.chain</code>	Number of MCMC chains. Default value is 4.
<code>n.iter.burn</code>	Number of warm-up MCMC iterations. Default value is 500.
<code>n.iter.sample</code>	Number of sampling MCMC iterations. Default value is 2500.
<code>thin</code>	A positive integer specifying the period for saving samples. Default value is 1.
<code>adapt_delta</code>	Numeric value between 0 and 1 indicating target average acceptance probability used in <code>rstan::sampling</code> . Default value is 0.9.
<code>verbose</code>	Logical value controlling the verbosity of output (i.e., warnings, messages, progress bar). Default is TRUE.
<code>seed</code>	A positive integer seed used for random number generation in MCMC. Default is NULL, which means the seed is generated from 1 to the maximum integer supported by R.

**Value**

A list of:

- a model object of class stanfit returned by rstan::sampling
- initial values used in MCMC

### Note

Before fitting the model, this function checks to ensure that the model specification is possible given the data files. These checks include:

- All tags in data are valid (i.e., include qPCR.N, qPCR.K, count, count.type, and site.cov).
- Dimensions of qPCR.N and qPCR.K are equal, and dimensions of count and count.type are equal (if count.type provided).
- Number of sites in qPCR and count data are equal.
- All data are numeric (i.e., integer or NA).
- Empty data cells (NA) match in qPCR.N and qPCR.K and in count and count.type.
- family is either 'poisson', 'negbin', or 'gamma'.
- p10priors and p10priors (if used) is a vector of two numeric values.
- site.cov has same number of rows as qPCR.N and count, if present
- site.cov is numeric, if present
- cov values match column names in site.cov, if present

If any of these checks fail, the function returns an error message.

### Examples

```
# Ex. 1: Implementing the joint model

# Load data
data(gobyData)

# Examine data in list
names(gobyData)

# Note that the surveyed sites (rows) should match in all data
dim(gobyData$qPCR.N)[1]
dim(gobyData$count)[1]

# Fit a basic model with paired eDNA and traditional survey data.
# Count data is modeled using a poisson distribution.
fit <- jointModel(data = gobyData, family = "poisson", p10priors = c(1,20),
                  multicore = FALSE)

# Ex. 2: Implementing the joint model with site-level covariates

# With the same data, fit a model including 'Filter_time' and 'Salinity'
# site-level covariates
# These covariates will scale the sensitivity of eDNA sampling relative to
# traditional surveys
# Count data is modeled using a poisson distribution.
fit.cov <- jointModel(data = gobyData, cov = c('Filter_time','Salinity'),
```

```

family = "poisson", p10priors = c(1,20),
multicore = FALSE)

# Ex. 3: Implementing the joint model with multiple traditional gear types

# Load data
data(greencrabData)

# Examine data in list
names(greencrabData)

# Note that the surveyed sites (rows) should match in all data
dim(greencrabData$qPCR.N)[1]
dim(greencrabData$count)[1]

# Fit a model estimating a catchability coefficient for traditional survey
# gear types.
# This model does not assume all traditional survey methods have the same
# catchability.
# Count data is modeled using a negative binomial distribution.
fit.q <- jointModel(data = greencrabData, cov = NULL, family = "negbin",
                    p10priors = c(1,20), q = TRUE, p10priors = c(0.25,0.25),
                    multicore = FALSE, initial_values = NULL,
                    n.chain = 4, n.iter.burn = 500,
                    n.iter.sample = 2500, thin = 1, adapt_delta = 0.9,
                    verbose = TRUE, seed = 123)

```

---

jointSelect

*Perform model selection using leave one out cross validation of model objects*

---

### Description

This function performs leave one out cross validation of a list of model fits using functions in the `loo` package, as described in Vehtari, Gelman, and Gabry (2017) [doi:10.1007/s11222-016-9696-4](https://doi.org/10.1007/s11222-016-9696-4). Compare models fit using `jointModel()` or models fits using `traditionalModel()`. See more examples in the [Package Vignette](#).

### Usage

```
jointSelect(modelfits)
```

### Arguments

`modelfits` A list containing model fits of class `stanfit`.

**Value**

A matrix of delta elpd (expected log pointwise predictive density) between model fits. Function is performed using the loo package.

**Note**

Before model selection, this function makes the following check:

- Input is a list of model fits of class 'stanfit'.
- All models compared were fit with either `jointModel()` or all with `traditionalModel()`.

If any of these checks fail, the function returns an error message.

**Examples**

```
data(greencrabData)

# Fit a model without estimating a catchability coefficient for traditional
# survey gear types.
# This model assumes all traditional survey methods have the same
# catchability.
# Count data is modeled using a poisson distribution.
fit.no.q <- jointModel(data = greencrabData, family = "poisson",
                      p10priors = c(1,20), q = FALSE, multicore = FALSE)

# Fit a model estimating a catchability coefficient for traditional
# survey gear types.
# This model does not assume all traditional survey methods have the
# same catchability.
# Gear type 1 is used as the reference gear type.
# Count data is modeled using a negative binomial distribution.
fit.q <- jointModel(data = greencrabData, family = "negbin",
                   p10priors = c(1,20), q = TRUE, multicore = FALSE)

# Perform model selection
jointSelect(modelfits = list(fit.no.q$model, fit.q$model))
```

---

jointSummarize

*Summarize posterior distributions of model parameters.*

---

**Description**

This function summarizes the posterior distributions of specified parameters from a model fit. Summary includes mean, sd, and specified quantiles, as well as effective sample size (`n_eff`) and `Rhat` for estimated parameters. See more examples in the [Package Vignette](#).

**Usage**

```
jointSummarize(modelfit, par = "all", probs = c(0.025, 0.975), digits = 3)
```

**Arguments**

<code>modelfit</code>	An object of class <code>stanfit</code> .
<code>par</code>	A character vector of parameter names. The default is <code>'all'</code> .
<code>probs</code>	A numeric vector of quantiles of interest. The default is <code>c(0.025,0.975)</code> .
<code>digits</code>	An integer indicating the number of decimal places to round values in summary table. Default value is 3.

**Value**

A summary table of parameter estimates.

**Note**

Before fitting the model, this function checks to ensure that the function is possible given the inputs. These checks include:

- Input model fit is an object of class `'stanfit'`.
- Input `probs` is a numeric vector.
- Input `par` is a character vector.
- Input `par` are present in fitted model.
- Input model fit has converged (i.e. no divergent transitions after warm-up).

If any of these checks fail, the function returns an error message.

**Examples**

```
data(greencrabData)

# Fit a model
modelfit <- jointModel(data = greencrabData, family = "negbin", q = TRUE,
                      multicore = FALSE)

# Create summary table of all parameters
jointSummarize(modelfit$model)

# Summarize just 'p10' parameter
jointSummarize(modelfit$model, par = "p10", probs = c(0.025, 0.975),
              digits = 3)
```

---

muCritical	Calculate mu_critical
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---

### Description

This function uses the full posterior distributions of parameters estimated by `jointModel()` to calculate `mu_critical`, or the expected catch rate at which the probabilities of a false positive eDNA detection and true positive eDNA detection are equal. See more examples in the [Package Vignette](#).

### Usage

```
muCritical(modelfit, cov.val = NULL, ci = 0.9)
```

### Arguments

<code>modelfit</code>	An object of class <code>stanfit</code>
<code>cov.val</code>	A numeric vector indicating the values of site-level covariates to use for prediction. Default is <code>NULL</code> .
<code>ci</code>	Credible interval calculated using highest density interval (HDI). Default is 0.9 (i.e., 90% credible interval).

### Value

A list with median `mu_critical` and lower and upper bounds on the credible interval. If multiple gear types are used, a table of `mu_critical` and lower and upper credible interval bounds is returned with one column for each gear type.

### Note

Before fitting the model, this function checks to ensure that the function is possible given the inputs. These checks include:

- Input model fit is an object of class `'stanfit'`.
- Input credible interval is a univariate numeric value greater than 0 and less than 1.
- Input model fit contains `p10` parameter.
- If model fit contains `alpha`, `cov.val` must be provided.
- Input `cov.val` is numeric.
- Input `cov.val` is the same length as the number of estimated covariates.
- Input model fit has converged (i.e. no divergent transitions after warm-up).

If any of these checks fail, the function returns an error message.

**Examples**

```
# Ex. 1: Calculating mu_critical with site-level covariates

# Load data
data(gobyData)

# Fit a model including 'Filter_time' and 'Salinity' site-level covariates
fit.cov <- jointModel(data = gobyData, cov = c('Filter_time','Salinity'),
                     family = "poisson", p10priors = c(1,20), q = FALSE,
                     multicore = FALSE)

# Calculate mu_critical at the mean covariate values (covariates are
# standardized, so mean = 0)
muCritical(fit.cov$model, cov.val = c(0,0), ci = 0.9)

# Calculate mu_critical at habitat size 0.5 z-scores greater than the mean
muCritical(fit.cov$model, cov.val = c(0,0.5), ci = 0.9)

# Ex. 2: Calculating mu_critical with multiple traditional gear types

# Load data
data(greencrabData)

# Fit a model with no site-level covariates
fit.q <- jointModel(data = greencrabData, cov = NULL, family = "negbin",
                   p10priors = c(1,20), q = TRUE, multicore = FALSE)

# Calculate mu_critical
muCritical(fit.q$model, cov.val = NULL, ci = 0.9)
```

---

traditionalModel

*Specify and fit model using count data from traditional, non eDNA surveys*


---

**Description**

This function implements a Bayesian model that estimates expected species catch rate using count data from traditional, non eDNA surveys. When multiple traditional gear types are used, an optional variation allows estimation of catchability coefficients, which scale the catchability of gear types relative to the expected catch rate of a reference gear type. Model is implemented using Bayesian inference using the `rstan` package, which uses Hamiltonian Monte Carlo to simulate the posterior distributions. See more examples in the [Package Vignette](#).

**Usage**

```
traditionalModel(
  data,
```



```

family = "poisson",
q = FALSE,
phipriors = NULL,
multicore = TRUE,
initial_values = NULL,
n.chain = 4,
n.iter.burn = 500,
n.iter.sample = 2500,
thin = 1,
adapt_delta = 0.9,
verbose = TRUE,
seed = NULL
)

```

### Arguments

data	A list containing data necessary for model fitting. Valid tags are count and count.type. count is a matrix or data frame of traditional survey count data, with first dimension equal to the number of sites (i) and second dimension equal to the maximum number of traditional survey replicates at a given site (j). count.type is an optional matrix or data frame of integers indicating gear type (k) used in corresponding count data, with first dimension equal to the number of sites (i) and second dimension equal to the maximum number of traditional survey replicates at a given site (j). Values should be integers beginning with 1 (referring to the first gear type) to n (last gear type). Empty cells should be NA and will be removed during processing. Sites, i, should be consistent in all count data.
family	The distribution class used to model traditional survey count data. Options include poisson ('poisson'), negative binomial ('negbin'), and gamma ('gamma'). Default value is 'poisson'.
q	A logical value indicating whether to estimate a catchability coefficient, q, for traditional survey gear types (TRUE) or to not estimate a catchability coefficient, q, for traditional survey gear types (FALSE). Default value is FALSE.
phipriors	A numeric vector indicating gamma distribution hyperparameters (shape, rate) used as the prior distribution for phi, the overdispersion in the negative binomial distribution for traditional survey gear data. Used when family = 'negbin.' If family = 'negbin', then default vector is c(0.25,0.25), otherwise, default is NULL.
multicore	A logical value indicating whether to parallelize chains with multiple cores. Default is TRUE.
initial_values	A list of lists of initial values to use in MCMC. The length should equal the number of MCMC chains. Initial values can be provided for parameters: mu and q. If no initial values are provided, default random values are drawn.
n.chain	Number of MCMC chains. Default value is 4.
n.iter.burn	Number of warm-up MCMC iterations. Default value is 500.
n.iter.sample	Number of sampling MCMC iterations. Default value is 2500.

<code>thin</code>	A positive integer specifying the period for saving samples. Default value is 1.
<code>adapt_delta</code>	Numeric value between 0 and 1 indicating target average acceptance probability used in <code>rstan::sampling</code> . Default value is 0.9.
<code>verbose</code>	Logical value controlling the verbosity of output (i.e., warnings, messages, progress bar). Default is TRUE.
<code>seed</code>	A positive integer seed used for random number generation in MCMC. Default is NULL, which means the seed is generated from 1 to the maximum integer supported by R.

**Value**

A list of:

- a model object of class `stanfit` returned by `rstan::sampling`
- initial values used in MCMC

**Note**

Before fitting the model, this function checks to ensure that the model specification is possible given the data files. These checks include:

- All tags in data are valid (i.e., include `count` and `count.type`).
- Number of sites in `count` and `count.type` data are equal.
- All data are numeric (i.e., integer or NA).
- Empty data cells (NA) match in `count` and `count.type`.
- family is 'poisson', 'negbin', or 'gamma'.
- `phipriors` (if used) is a vector of two numeric values.

If any of these checks fail, the function returns an error message.

**Examples**

```
# Load data
data(greencrabData)

# Examine data in list
# This function uses only traditional survey count data and optionally
# the count type data
names(greencrabData)

# Note that the surveyed sites (rows) should match in the data
dim(greencrabData$count)[1]
dim(greencrabData$count.type)[1]

# Fit a model without estimating a catchability coefficient for traditional
# survey gear types.
# This model assumes all traditional survey methods have the same
# catchability.
# Count data is modeled using a poisson distribution.
```

```
fit.no.q <- traditionalModel(data = greencrabData, family = "poisson",
                             q = FALSE, phipriors = NULL, multicore = FALSE,
                             verbose = TRUE)

# Fit a model estimating a catchability coefficient for traditional survey
# gear types.
# This model does not assume all traditional survey methods have the same
# catchability.
# Count data is modeled using a negative binomial distribution.
fit.q <- traditionalModel(data = greencrabData, family = "negbin", q = TRUE,
                          phipriors = c(0.25,0.25), multicore = FALSE,
                          initial_values = NULL, n.chain = 4,
                          n.iter.burn = 500, n.iter.sample = 2500, thin = 1,
                          adapt_delta = 0.9, verbose = TRUE, seed = 123)
```

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