

# Package ‘aod’

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**Title** Analysis of Overdispersed Data

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**Depends** R (>= 2.0.0), methods, stats

**Suggests** MASS, boot, lme4

**Description** This package provides a set of functions to analyse overdispersed counts or proportions. Most of the methods are already available elsewhere but are scattered in different packages. The proposed functions should be considered as complements to more sophisticated methods such as generalized estimating equations (GEE) or generalized linear mixed effect models (GLMM).

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**LazyData** yes

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## R topics documented:

aic-class . . . . .	2
AIC-methods . . . . .	3
anova-methods . . . . .	4
antibio . . . . .	5

aod-pkg . . . . .	6
betabin . . . . .	7
coef-methods . . . . .	10
cohorts . . . . .	11
deviance-methods . . . . .	11
df.residual-methods . . . . .	12
dja . . . . .	12
donner . . . . .	13
drs-class . . . . .	15
fitted-methods . . . . .	15
glimML-class . . . . .	16
glimQL-class . . . . .	17
iccbin . . . . .	17
iccbin-class . . . . .	19
invlink . . . . .	20
link . . . . .	21
lizards . . . . .	21
logLik-methods . . . . .	22
mice . . . . .	23
negbin . . . . .	24
orob1 . . . . .	26
orob2 . . . . .	27
predict-methods . . . . .	27
quasibin . . . . .	28
quasipois . . . . .	30
rabbits . . . . .	32
raoscott . . . . .	33
rats . . . . .	34
residuals-methods . . . . .	35
salmonella . . . . .	36
splitbin . . . . .	37
summary,aic-method . . . . .	38
summary.glimML-class . . . . .	39
varbin . . . . .	40
varbin-class . . . . .	42
vcov-methods . . . . .	43
wald.test . . . . .	43
<b>Index</b>	<b>45</b>

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aic-class

*Representation of Objects of Formal Class "aic"*


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## Description

Representation of the output of function AIC.

**Slots**

**istats** A data frame with 3 columns describing the models indicated by the row names:

- **df** number of parameters in the model,
- **AIC** Akaike information criterion for the model (see [AIC](#)),
- **AICc** small-sample corrected Akaike information criterion for the model (see [AIC](#)).

**Methods**

**summary** signature(object = "aic")

**show** signature(object = "aic")

---

AIC-methods

*Akaike Information Criteria*


---

**Description**

Extracts the Akaike information criterion (AIC) and the corrected AIC (AICc) from fitted models of formal class “glimML” and possibly computes derived statistics.

**Usage**

```
## S4 method for signature 'glimML'
AIC(object, ..., k = 2)
```

**Arguments**

**object** fitted model of formal class “glimML” (functions `betabin` or `negbin`).

**...** optional list of fitted models separated by commas.

**k** numeric scalar, with a default value set to 2, thus providing the regular AIC.

**Details**

$AIC = -2 \log\text{-likelihood} + 2 * n_{par}$ , where  $n_{par}$  represents the number of parameters in the fitted model.

$AICc = AIC + 2 * n_{par} * (n_{par} + 1) / (n_{obs} - n_{par} + 1)$ , where  $n_{obs}$  is the number of observations used to compute the log-likelihood. It should be used when the number of fitted parameters is large compared to sample size, i.e., when  $n_{obs} / n_{par} < 40$  (Hurvich and Tsai, 1995).

**Methods**

**glimML** Extracts the AIC and AICc from models of formal class “glimML”, fitted by functions `betabin` and `negbin`.

## References

- Burnham, K.P., Anderson, D.R., 2002. *Model selection and multimodel inference: a practical information-theoretic approach*. New-York, Springer-Verlag, 496 p.
- Hurvich, C.M., Tsai, C.-L., 1995. *Model selection for extended quasi-likelihood models in small samples*. *Biometrics*, 51 (3): 1077-1084.

## See Also

Examples in [betabin](#) and see [AIC](#) in package **stats**.

---

anova-methods

*Likelihood-Ratio Tests for Nested ML Models*

---

## Description

Performs likelihood-ratio tests on nested models. Currently, one method was implemented for beta-binomial models (`betabin`) or negative-binomial models (`negbin`).

## Usage

```
## S4 method for signature 'glimML'
anova(object, ...)
```

## Arguments

<code>object</code>	Fitted model of class “ <code>glimML</code> ”.
<code>...</code>	Further models to be tested or arguments passed to the <code>print</code> function.

## Details

The `anova` method for models of formal class “`glimML`” needs at least 2 nested models of the same type (either beta-binomial or negative-binomial models: they cannot be mixed). The quantity of interest is the deviance difference between the compared models: it is a log-likelihood ratio statistic. Under the null hypothesis that 2 nested models fit the data equally well, the deviance difference has an approximate  $\chi^2$  distribution with degrees of freedom = the difference in the number of parameters between the compared models (Mc Cullagh and Nelder, 1989).

## Value

An object of formal class “`anova.glimML`” with 3 slots:

<code>models</code>	A vector of character strings with each component giving the name of the models and the formulas for the fixed and random effects.
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anova.table	A data frame containing the results. Row names correspond to the models.	
logL	numeric	maximized log-likelihood
k	numeric	number of parameters in the model
AIC	numeric	Akaike information criterion for the model
AICc	numeric	Corrected Akaike information criterion for the model
BIC	numeric	Bayesian information criterion the model
Resid. dev.	numeric	Residual deviance
Resid. Df	numeric	df of the residuals
Test	character	Nested models which are tested
Deviance	numeric	Deviance difference between the 2 models
Df	numeric	df associated with deviance difference
P(> Chi2)	numeric	P value associated with H0.
type	A character chain indicating the kind of fitted model: “BB” for beta-binomial, or “NB” for negative-binomial model.	

### Warning

The comparison between 2 or more models will only be valid if they are fitted to the same data set.

### References

McCullagh, P., Nelder, J.A., 1989. *Generalized linear models*. London, Chapman & Hall, 511 p.  
See Appendix C. Likelihood ratio statistics, p. 476-478.

### See Also

[anova.glm](#), [AIC](#)

### Examples

```
data(orob2)
# likelihood ratio test for the effect of root
fm1 <- betabin(cbind(y, n - y) ~ seed, ~ 1, data = orob2)
fm2 <- betabin(cbind(y, n - y) ~ seed + root, ~ 1, data = orob2)
anova(fm1, fm2)
```

### Description

Hypothetical drug trial to compare the effect of four antibiotics against Shipping fever in calves (Shoukri and Pause, 1999, Table 3.11).

**Usage**

```
data(antibio)
```

**Format**

A data frame with 24 observations on the following 3 variables.

**treatment** A factor with levels 1, 2, 3 and 4

**n** A numeric vector: the number of treated animals within a two-week period.

**y** A numeric vector: the number of deaths at the end of the two weeks.

**References**

Shoukri, M.M., Pause, C.A., 1999, 2nd ed. *Statistical methods for health sciences*. CRC Press, London.

---

aod-pkg

*Analysis of Overdispersed Data*


---

**Description**

This package provides a set of functions to analyse overdispersed counts or proportions. Most of the methods are already available elsewhere but are scattered in different packages. The proposed functions should be considered as complements to more sophisticated methods such as generalized estimating equations (GEE) or generalized linear mixed effect models (GLMM).

**Details**

```
Package:    aod
Version:    1.1-32
Date:       2010-04-02
Depends:    R (>= 2.0.0), methods, stats
Suggests:   MASS, nlme, boot
License:    GPL version 2 or newer
URL:        http://cran.r-project.org/package=aod
LazyLoad:   yes
LazyData:   yes
```

**Index :**

AIC-methods	Akaike Information Criteria
aic-class	Representation of Objects of Formal Class "aic"
anova-method	Likelihood-Ratio Tests for Nested ML Models
antibio	Antibiotics against Shipping Fever in Calves
betabin	Beta-Binomial Model for Proportions

coef-methods	Methods for Function “coef” in Package <b>aod</b>
cohorts	Data set: Age, Period and Cohort Effects for Vital Rates
deviance-methods	Methods for Function deviance in Package <b>aod</b>
df.residual-methods	Methods for Function <code>df.residual</code> in Package <b>aod</b>
dja	Mortality of Djallonke Lambs in Senegal
donner	Test of Proportion Homogeneity using Donner’s Adjustment
drs-class	Representation of Objects of Formal Class “drs”
fitted-methods	Methods for Function <code>fitted</code> in Package <b>aod</b>
glimML-class	Representation of Models of Formal Class “glimML”
glimQL-class	Representation of Models of Formal Class “glimQL”
icc	Intra-Cluster Correlation
icc-class	Representation of Objects of Formal Class “icc”
invlink	Transformation from the Link Scale to the Observation Scale
link	Transformation from the Observation Scale to the Link Scale
lizards	A Comparison of Site Preferences of Two Species of Lizard
logLik-methods	Methods for Functions “logLik” in Package <b>aod</b>
mice	Pregnant Female Mice Experiment
negbin	Negative-Binomial Model for Counts
orob1	Germination Data
orob2	Germination Data
predict-methods	Methods for Function “predict” in Package <b>aod</b>
quasibin	Quasi-Likelihood Model for Proportions
quasipois	Quasi-Likelihood Model for Counts
rabbits	Rabbits Foetuses Survival Experiment
raoscott	Test of Proportion Homogeneity using Rao and Scott’s Adjustment
rats	Rats Diet Experiment
residuals-methods	Residuals for Maximum-Likelihood and Quasi-Likelihood Models
salmonella	Salmonella Reverse Mutagenicity Assay
splitbin	Splits Binomial Data into Bernoulli Data
summary,aic-method	Akaike Information Statistics
summary.glimML-class	Summary of Objects of Class “summary.glimML”
varbin	Mean, Variance and Confidence Interval of a Proportion
varbin-class	Representation of Objects of Formal Class “varbin”
vcov-methods	Methods for Function “vcov” in Package <b>aod</b>
wald.test	Wald Test for Model Coefficients

**Author(s)**

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 Maintainer: Renaud Lancelot

**Description**

Fits a beta-binomial generalized linear model accounting for overdispersion in clustered binomial data  $(n, y)$ .

**Usage**

```
betabin(formula, random, data, link = c("logit", "cloglog"), phi.ini = NULL,
        warnings = FALSE, na.action = na.omit, fixpar = list(),
        hessian = TRUE, control = list(maxit = 2000), ...)
```

**Arguments**

formula	A formula for the fixed effects $b$ . The left-hand side of the formula must be of the form <code>cbind(y, n - y)</code> where the modelled probability is $y/n$ .
random	A right-hand formula for the overdispersion parameter(s) $\phi$ .
link	The link function for the mean $p$ : "logit" or "cloglog".
data	A data frame containing the response ( $n$ and $y$ ) and explanatory variable(s).
phi.ini	Initial values for the overdispersion parameter(s) $\phi$ . Default to 0.1.
warnings	Logical to control the printing of warnings occurring during log-likelihood maximization. Default to FALSE (no printing).
na.action	A function name: which action should be taken in the case of missing value(s).
fixpar	A list with 2 components (scalars or vectors) of the same size, indicating which parameters are fixed (i.e., not optimized) in the global parameter vector $(b, \phi)$ and the corresponding fixed values. For example, <code>fixpar = list(c(4, 5), c(0, 0))</code> means that 4th and 5th parameters of the model are set to 0.
hessian	A logical. When set to FALSE, the hessian and the variances-covariances matrices of the parameters are not computed.
control	A list to control the optimization parameters. See <code>optim</code> . By default, set the maximum number of iterations to 2000.
...	Further arguments passed to <code>optim</code> .

**Details**

For a given cluster  $(n, y)$ , the model is:

$$y \sim \lambda \sim \text{Binomial}(n, \lambda)$$

with  $\lambda$  following a Beta distribution  $\text{Beta}(a1, a2)$ .

If  $B$  denotes the beta function, then:

$$P(\lambda) = \frac{\lambda^{a1-1} * (1-\lambda)^{a2-1}}{B(a1, a2)}$$

$$E[\lambda] = \frac{a1}{a1 + a2}$$

$$\text{Var}[\lambda] = \frac{a1 * a2}{(a1 + a2 + 1) * (a1 + a2)^2}$$

The marginal beta-binomial distribution is:

$$P(y) = \frac{C(n, \tilde{y}) * B(a1 + y, a2 + n - y)}{B(a1, \tilde{a}2)}$$

The function uses the parameterization  $p = \frac{a1}{a1+a2} = h(Xb) = h(\eta)$  and  $\phi = \frac{1}{a1+a2+1}$ , where  $h$  is the inverse of the link function (logit or complementary log-log),  $X$  is a design-matrix,  $b$  is a vector of fixed effects,  $\eta = Xb$  is the linear predictor and  $\phi$  is the overdispersion parameter (i.e., the intracluster correlation coefficient, which is here restricted to be positive).

The marginal mean and variance are:

$$E[y] = n * p$$

$$\text{Var}[y] = n * p * (1 - p) * [1 + (n - 1) * \phi]$$

The parameters  $b$  and  $\phi$  are estimated by maximizing the log-likelihood of the marginal model (using the function `optim`). Several explanatory variables are allowed in  $b$ , only one in  $\phi$ .

## Value

An object of formal class “glimML”: see [glimML-class](#) for details.

## Author(s)

Matthieu Lesnoff <matthieu.lesnoff@cirad.fr>, Renaud Lancelot <renaud.lancelot@cirad.fr>

## References

- Crowder, M.J., 1978. *Beta-binomial anova for proportions*. Appl. Statist. 27, 34-37.
- Griffiths, D.A., 1973. *Maximum likelihood estimation for the beta-binomial distribution and an application to the household distribution of the total number of cases of disease*. Biometrics 29, 637-648.
- Prentice, R.L., 1986. *Binary regression using an extended beta-binomial distribution, with discussion of correlation induced by covariate measurement errors*. J.A.S.A. 81, 321-327.
- Williams, D.A., 1975. *The analysis of binary responses from toxicological experiments involving reproduction and teratogenicity*. Biometrics 31, 949-952.

## See Also

[glimML-class](#), [glm](#) and [optim](#)

## Examples

```
data(orob2)
fm1 <- betabin(cbind(y, n - y) ~ seed, ~ 1, data = orob2)
fm2 <- betabin(cbind(y, n - y) ~ seed + root, ~ 1, data = orob2)
fm3 <- betabin(cbind(y, n - y) ~ seed * root, ~ 1, data = orob2)
# show the model
fm1; fm2; fm3
# AIC
AIC(fm1, fm2, fm3)
```

```

summary(AIC(fm1, fm2, fm3), which = "AICc")
# Wald test for root effect
wald.test(b = coef(fm3), Sigma = vcov(fm3), Terms = 3:4)
# likelihood ratio test for root effect
anova(fm1, fm3)
# model predictions
New <- expand.grid(seed = levels(orob2$seed),
                  root = levels(orob2$root))
data.frame(New, predict(fm3, New, se = TRUE, type = "response"))
# Djallonke sheep data
data(dja)
betabin(cbind(y, n - y) ~ group, ~ 1, dja)
# heterogeneous phi
betabin(cbind(y, n - y) ~ group, ~ group, dja,
        control = list(maxit = 1000))
# phi fixed to zero in group TREAT
betabin(cbind(y, n - y) ~ group, ~ group, dja,
        fixpar = list(4, 0))
# glim without overdispersion
summary(glm(cbind(y, n - y) ~ group,
            family = binomial, data = dja))
# phi fixed to zero in both groups
betabin(cbind(y, n - y) ~ group, ~ group, dja,
        fixpar = list(c(3, 4), c(0, 0)))

```

---

coef-methods

*Methods for Function "coef" in Package "aod"*


---

## Description

Extract the fixed-effect coefficients from fitted objects.

## Methods

**ANY** Generic function: see [coef](#).

**glimML** Extract the estimated fixed-effect coefficients from objects of formal class “glimML”. Presently, these objects are generated by functions `betabin` and `negbin`.

**glimQL** Extract the estimated fixed-effect coefficients from objects of formal class “glimQL”. Presently, these objects are generated by functions `quasibin` and `quasipois`.

**Description**

Number of prostate cancer deaths and midperiod population for nonwhites in the USA by age and period. The cohort index  $k$  is related to age and period indices ( $i$  and  $j$ , respectively) by  $k = j + I - i$ , where  $I = \max(i)$  (Holford, 1983, Table 2).

**Usage**

```
data(cohorts)
```

**Format**

A data frame with 49 observations on the following 4 variables.

**period** A factor with levels 1935-, 1940-, ..., 1965-.

**age** A factor with levels 50-, 55-, ..., 80-.

**y** Numeric: the number of prostate cancer deaths.

**n** Numeric: the midperiod population size.

**References**

Holford, T.R., 1983. *The estimation of age, period and cohort effects for vital rates*. Biometrics 39, 311-324.

**Description**

Extracts the deviance fitted models.

**Methods**

**ANY** Generic function: see [deviance](#).

**glimML** Extracts the deviance from models fitted by betabin or negbin.

---

df.residual-methods     *Methods for Function "df.residual" in Package "aod"*

---

### Description

Computes the number of degrees of freedom of the residuals from fitted objects.

### Methods

**ANY** Generic function: see [df.residual](#).

**glimML** Computes the df of residuals for models fitted by betabin or negbin.

**glimQL** Computes the df of residuals for models fitted by quasibin or quasipois.

---

dja     *Mortality of Djallonke Lambs in Senegal*

---

### Description

Field trial to assess the effect of ewes deworming (prevention of gastro-intestinal parasitism) on the mortality of their offspring (age < 1 year). This data set is extracted from a large database on small ruminants production and health in Senegal (Lancelot et al., 1998). Data were collected in a sample of herds in Kolda (Upper Casamance, Senegal) during a multi-site survey (Faugère et al., 1992). See also the references below for a presentation of the follow-up survey (Faugère and Faugère, 1986) and a description of the farming systems (Faugère et al., 1990).

### Usage

```
data(dja)
```

### Format

A data frame with 21 observations on the following 4 variables.

**group** a factor with 2 levels: CTRL and TREAT, indicating the treatment.

**village** a factor indicating the village of the herd.

**herd** a factor indicating the herd.

**n** a numeric vector: the number of animals exposed to mortality.

**trisk** a numeric vector: the exposition time to mortality (in year).

**y** a numeric vector: the number of deaths.

## References

- Faugère, O., Faugère, B., 1986. *Suivi de troupeaux et contrôle des performances individuelles des petits ruminants en milieu traditionnel africain. Aspects méthodologiques*. Rev. Elev. Méd. vét. Pays trop., 39 (1): 29-40.
- Faugère, O., Dockès, A.-C., Perrot, C., Faugère, B., 1990. *L'élevage traditionnel des petits ruminants au Sénégal. I. Pratiques de conduite et d'exploitation des animaux chez les éleveurs de la région de Kolda*. Revue Elev. Méd. vét. Pays trop. 43: 249-259.
- Faugère, O., Tillard, E., Faugère, B., 1992. *Prophylaxie chez les petits ruminants au Sénégal : régionalisation d'une politique nationale de protection sanitaire*. In: B. Rey, S. H. B. Lebbie, L. Reynolds (Ed.), First biennial conference of the African Small Ruminant Research Network, ILCA, 1990, ILRAD, Nairobi, pp. 307-314.
- Lancelot, R., Faye, B., Juanès, X., Ndiaye, M., Pérochon, L., Tillard, E., 1998. *La base de données BAOBAB: un outil pour modéliser la production et la santé des petits ruminants dans les systèmes d'élevage traditionnels au Sénégal*. Revue Elev. Méd. vét. Pays trop., 51 (2): 135-146.

---

 donner

 Test of Proportion Homogeneity using Donner's Adjustment
 

---

## Description

Tests the homogeneity of proportions between  $I$  groups ( $H_0: p_1 = p_2 = \dots = p_I$ ) from clustered binomial data  $(n, y)$  using the adjusted  $\chi^2$  statistic proposed by Donner (1989).

## Usage

```
donner(formula = NULL, response = NULL,
       weights = NULL, group = NULL, data, C = NULL)
```

## Arguments

formula	An optional formula where the left-hand side is either a matrix of the form <code>cbind(y, n-y)</code> , where the modelled probability is $y/n$ , or a vector of proportions to be modelled ( $y/n$ ). In both cases, the right-hand side must specify a single grouping variable. When the left-hand side of the formula is a vector of proportions, the argument <code>weight</code> must be used to indicate the denominators of the proportions.
response	An optional argument indicating either a matrix of the form <code>cbind(y, n-y)</code> , where the modelled probability is $y/n$ , or a vector of proportions to be modelled ( $y/n$ ).
weights	An optional argument used when the left-hand side of <code>formula</code> or <code>response</code> is a vector of proportions: <code>weight</code> is the denominator of the proportion.
group	An optional argument only used when <code>response</code> is used. In this case, this argument is a factor indicating a grouping variable.
data	A data frame containing the response ( $n$ and $y$ ) and the grouping variable.
C	If not <code>NULL</code> , a numerical vector of $I$ cluster correction factors.

## Details

The  $\chi^2$  statistic is adjusted with the correction factor  $C_i$  computed in each group  $i$ . The test statistic is given by:

$$X^2 = \sum_i \frac{(y_i - n_i * p)^2}{C_i * n_i * p * (1 - p)}$$

where  $C_i = 1 + (nA_i - 1) * \rho$ ,  $nA_i$  is a scalar depending on the cluster sizes, and  $\rho$  is the ANOVA estimate of the intra-cluster correlation, assumed common across groups (see Donner, 1989 or Donner et al., 1994). The statistic is compared to a  $\chi^2$  distribution with  $I - 1$  degrees of freedom. Fixed correction factors can be specified with the argument `C`.

## Value

An object of formal class “`drs`”: see [`drs-class`](#) for details. The slot `tab` provides the proportion of successes and the correction factor for each group.

## Author(s)

Matthieu Lesnoff <matthieu.lesnoff@cirad.fr>, Renaud Lancelot <renaud.lancelot@cirad.fr>

## References

- Donner, A., 1989. *Statistical methods in ophthalmology: an adjusted chi-squared approach*. *Biometrics* 45, 605-611.
- Donner, A., 1993. *The comparison of proportions in the presence of litter effects*. *Prev. Vet. Med.* 18, 17-26.
- Donner, A., Eliasziw, M., Klar, N., 1994. *A comparison of methods for testing homogeneity of proportions in teratologic studies*. *Stat. Med.* 13, 1253-1264.

## See Also

[`chisq.test`](#), [`raoscott`](#), [`drs-class`](#)

## Examples

```
data(rats)
donner(formula = cbind(y, n - y) ~ group, data = rats)
donner(formula = y/n ~ group, weights = n, data = rats)
donner(response = cbind(y, n - y), group = group, data = rats)
donner(response = y/n, weights = n, group = group, data = rats)
# standard test
donner(cbind(y, n - y) ~ group, data = rats, C = c(1, 1))
data(antibio)
donner(cbind(y, n - y) ~ treatment, data = antibio)
```

---

 drs-class

*Representation of Objects of Formal Class "drs"*


---

### Description

Representation of the output of functions `donner` and `raoscott`.

### Objects from the Class

Objects can be created by calls of the form `new("drs", ...)` or, more commonly, via the `donner` or `raoscott` functions.

### Slots

`CALL` The call of the function.

`tab` A data frame containing test information. The content of the data frame depends on the type of the function which generated it.

`rho` The ANOVA estimate of the intra-cluster correlation (function `donner`).

`X2` The adjusted  $\chi^2$  statistic.

### Methods

**donner** signature(object = "drs"): see [donner](#).

**raoscott** signature(object = "drs"): see [raoscott](#).

---

 fitted-methods

*Methods for Function "fitted" in Package "aod"*


---

### Description

Extracts the fitted values from models.

### Methods

**ANY** Generic function: see [fitted](#).

**glimML** Extract the fitted values from models of formal class "glimML", presently generated by functions `betabin` and `negbin`.

**glimQL** Extract the fitted values from models of formal class "glimQL", presently generated by functions `quasibin` and `quasibin`.

glimML-class

*Representation of Models of Formal Class "glimML"***Description**

Representation of models of formal class "glimML" fitted by maximum-likelihood method.

**Objects from the Class**

Objects can be created by calls of the form `new("glimML", ...)` or, more commonly, via the functions `betabin` or `negbin`.

**Slots**

`CALL` The call of the function.

`link` The link function used to transform the mean: "logit", "cloglog" or "log".

`method` The type of fitted model: "BB" for beta-binomial and "NB" for negative-binomial models.

`formula` The formula used to model the mean.

`random` The formula used to model the overdispersion parameter  $\phi$ .

`data` Data set to which model was fitted. Different from the original data in case of missing value(s).

`param` The vector of the ML estimated parameters  $b$  and  $\phi$ .

`varparam` The variance-covariance matrix of the ML estimated parameters  $b$  and  $\phi$ .

`fixed.param` The vector of the ML estimated fixed-effect parameters  $b$ .

`random.param` The vector of the ML estimated random-effect (correlation) parameters  $\phi$ .

`logL` The log-likelihood of the fitted model.

`logL.max` The log-likelihood of the maximal model (data).

`dev` The deviance of the model, i.e.,  $-2 * (\logL - \logL.max)$ .

`df.residual` The residual degrees of freedom of the fitted model.

`nbpar` The number of **estimated** parameters, i.e., `nbpar` = total number of parameters - number of fixed parameters. See argument `fixpar` in `betabin` or `negbin`.

`iterations` The number of iterations performed in `optim`.

`code` An integer (returned by `optim`) indicating why the optimization process terminated.

- 1 Relative gradient is close to 0, current iterate is probably solution.
- 2 Successive iterates within tolerance, current iterate is probably solution.
- 3 Last global step failed to locate a point lower than estimate. Either estimate is an approximate local minimum of the function or `steptol` is too small.
- 4 Iteration limit exceeded.
- 5 Maximum step size `stepmax` exceeded 5 consecutive times. Either the function is unbounded below, becomes asymptotic to a finite value from above in some direction or `stepmax` is too small.

msg Message returned by `optim`.

singular.hessian Logical: true when fitting provided a singular hessian, indicating an overparameterized model.

param.ini The initial values provided to the ML algorithm.

na.action A function defining the action taken when missing values are encountered.

---

glimQL-class *Representation of Models of Formal Class "glimQL"*

---

### Description

Representation of models of formal class "glimQL" fitted by quasi-likelihood method.

### Objects from the Class

Objects can be created by calls of the form `new("glimQL", ...)` or, more commonly, via the `quasibin` or `quasipois` functions.

### Slots

CALL The call of the function.

fm A fitted model of class "glm".

phi The overdispersion parameter.

### Methods

**show** `signature(object = "glimQL")`: Main results of "glimQL" models.

---

iccbin *Intra-Cluster Correlation for Binomial Data*

---

### Description

This function calculates point estimates of the intraclass correlation  $\rho$  from clustered binomial data  $(n_1, y_1), (n_2, y_2), \dots, (n_K, y_K)$  (with  $K$  the number of clusters), using a 1-way random effect model. Three estimates, following methods referred to as "A", "B" and "C" in Goldstein et al. (2002), can be obtained.

### Usage

`iccbin(n, y, data, method = c("A", "B", "C"), nAGQ = 1, M = 1000)`

**Arguments**

n	Vector of the denominators of the proportions.
y	Vector of the numerators of the proportions.
data	A data frame containing the variables n and y.
method	A character (“A”, “B” or “C”) defining the calculation method. See Details.
nAGQ	Same as in function <code>glmer</code> of package <b>lme4</b> . Only for methods “A” and “B”. Default to 1.
M	Number of Monte Carlo (MC) replicates used in method “B”. Default to 1000.

**Details**

Before computations, the clustered data are split into binary (0/1) observations  $y_{ij}$  (obs.  $j$  in cluster  $i$ ). The calculation methods are described in Goldstein et al. (2002). Methods “A” and “B” assume a 1-way generalized linear mixed model, and method “C” a 1-way linear mixed model. For “A” and “B”, function `iccbin` uses the logistic binomial-Gaussian model:

$$y_{ij}|p_{ij} \sim \text{Bernoulli}(p_{ij}),$$

$$\text{logit}(p_{ij}) = b_0 + u_i,$$

where  $b_0$  is a constant and  $u_i$  a cluster random effect with  $u_i \sim N(0, s_u^2)$ . The ML estimate of the variance component  $s_u^2$  is calculated with the function `glmer` of package **lme4**. The intra-class correlation  $\rho = \text{Corr}[y_{ij}, y_{ij'}]$  is then calculated with a first-order model linearization around  $E[u_i] = 0$  in method “A”, and with Monte Carlo simulations in method “B”.

For “C”, function `iccbin` provides the common ANOVA (moments) estimate of  $\rho$ . For details, see for instance Donner (1986), Searle et al. (1992) or Ukoumunne (2002).

**Value**

An object of formal class “iccbin”, with 3 slots:

CALL	The call of the function.
features	A character vector summarizing the main features of the method used.
rho	The point estimate of the intraclass correlation $\rho$ .

**Author(s)**

Matthieu Lesnoff <matthieu.lesnoff@cirad.fr>, Renaud Lancelot <renaud.lancelot@cirad.fr>

**References**

- Donner A., 1986, *A review of inference procedures for the intraclass correlation coefficient in the one-way random effects model*. International Statistical Review 54, 67-82.
- Searle, S.R., Casella, G., McCulloch, C.E., 1992. *Variance components*. Wiley, New York.
- Ukoumunne, O. C., 2002. *A comparison of confidence interval methods for the intraclass correlation coefficient in cluster randomized trials*. Statistics in Medicine 21, 3757-3774.
- Golstein, H., Browne, H., Rasbash, J., 2002. *Partitioning variation in multilevel models*. Understanding Statistics 1(4), 223-231.

**See Also**

[iccbin-class](#), [glmer](#)

**Examples**

```

data(rats)
tmp <- rats[rats$group == "TREAT", ]
# A: glmm (model linearization)
iccbin(n, y, data = tmp, method = "A")
iccbin(n, y, data = tmp, method = "A", nAGQ = 10)
# B: glmm (Monte Carlo)
iccbin(n, y, data = tmp, method = "B")
iccbin(n, y, data = tmp, method = "B", nAGQ = 10, M = 1500)
# C: lmm (ANOVA moments)
iccbin(n, y, data = tmp, method = "C")

## Not run:
# Example of confidence interval calculation with nonparametric bootstrap
require(boot)
foo <- function(X, ind) {
  n <- X$n[ind]
  y <- X$y[ind]
  X <- data.frame(n = n, y = y)
  iccbin(n = n, y = y, data = X, method = "C")@rho[1]
}
res <- boot(data = tmp[, c("n", "y")], statistic = foo, R = 500, sim = "ordinary", stype = "i")
res
boot.ci(res, conf = 0.95, type = "basic")

## End(Not run)

```

---

iccbin-class

*Representation of Objects of Formal Class "iccbin"*

---

**Description**

Representation of the output of function `iccbin`.

**Objects from the Class**

Objects can be created by calls of the form `new("iccbin", ...)` or, more commonly, via the function `iccbin`.

**Slots**

CALL The call of the function.

features A character vector summarizing the main features of the method used.

rho A numeric scalar giving the intra-cluster correlation.

**Methods**

**icc** signature(object = "iccbin"): see [iccbin](#).

---

 invlink

---

*Transformation from the Link Scale to the Observation Scale*


---

**Description**

The function transforms a variable from the link scale to the observation scale: probability or count.

**Usage**

```
invlink(x, type = c("cloglog", "log", "logit"))
```

**Arguments**

x A vector of real numbers.

type A character string. Legal values are "cloglog", "log" and "logit".

**Value**

$anti - logit(x) = exp(x)/(1 + exp(x))$

$anti - cloglog(x) = 1 - exp(-exp(x))$

**See Also**

[link](#)

**Examples**

```
x <- seq(-5, 5, length = 100)
plot(x, invlink(x, type = "logit"),
     type = "l", lwd = 2, ylab = "Probability")
lines(x, invlink(x, type = "cloglog"), lty = 2, lwd = 2)
grid(col = "black")
legend(-5, 1, legend = c("alogit(x)", "acloglog(x)"),
      lty = c(1, 2), bg = "white")
```

---

link	<i>Transformation from the Observation Scale to the Link Scale</i>
------	--

---

**Description**

The function transforms a variable from the observation scale (probability or count) to the link scale.

**Usage**

```
link(x, type = c("cloglog", "log", "logit"))
```

**Arguments**

x	A vector of real numbers.
type	A character string. Legal values are “cloglog”, “log” and “logit”.

**Value**

$$\text{logit}(x) = \log(x/(1-x))$$

$$\text{cloglog}(x) = \log(-\log(1-x))$$
**See Also**

[invlink](#)

**Examples**

```
x <- seq(.001, .999, length = 100)
plot(x, link(x, type = "logit"),
     type = "l", lwd = 2, ylab = "link(proba.)")
lines(x, link(x, type = "cloglog"), lty = 2, lwd = 2)
grid(col = "black")
legend(0, 6, legend = c("logit(x)", "cloglog(x)"),
      lty = c(1, 2), bg = "white")
```

---

lizards	<i>A Comparison of Site Preferences of Two Species of Lizard</i>
---------	--

---

**Description**

“These data describe the daytime habits of two species of lizards, *grahami* and *opalinus*. They were collected by observing occupied sites or perches and recording the appropriate description, namely species involved, time of the day, height and diameter of the perch and whether the site was sunny or shaded. Time of the day is recorded as early, mid-day or late.” (McCullagh and Nelder, 1989, p.129).

**Usage**

```
data(lizards)
```

**Format**

A data frame with 24 observations on the following 6 variables.

**Site** A factor with levels Sun and Shade.

**Diameter** A factor with levels D <= 2 and D > 2 (inches).

**Height** A factor with levels H < 5 and H >= 5 (feet).

**Time** A factor with levels Early, Mid-day and Late.

**grahami** A numeric vector giving the observed sample size for *grahami* lizards.

**opalinus** A numeric vector giving the observed sample size for *opalinus* lizards.

**Details**

The data were originally published in Fienberg (1970).

**Source**

McCullagh, P., Nelder, J.A., 1989. *Generalized linear models*. London, Chapman & Hall, 511 p.

**References**

Fienberg, S.E., 1970. *The analysis of multidimensional contingency tables*. Ecology 51: 419-433.

**Examples**

```
data(lizards)
```

---

 logLik-methods

*Methods for Functions "logLik" in Package "aod"*


---

**Description**

Extracts the maximized log-likelihood from fitted models of formal class "glimML".

**Usage**

```
## S4 method for signature 'glimML'
logLik(object, ...)
```

**Arguments**

**object** A fitted model of formal class "glimML" (functions betabin or negbin).  
**...** Other arguments passed to methods.

**Value**

A numeric scalar with 2 attributes: “df” (number of parameters in the model) and “nobs” (number of observations = degrees of freedom of the residuals + number of parameters in the model).

**Methods**

**ANY** Generic function: see [logLik](#).

**glimML** Extract the maximized log-likelihood from models of formal class “glimML”, fitted by functions `betabin` and `negbin`.

**See Also**

[logLik](#) in package `stats`.

---

mice

*Pregnant Female Mice Experiment*

---

**Description**

Unpublished laboratory data on the proportion of affected fetuses in two groups (control and treatment) of 10 pregnant female mice (Kupper and Haseman, 1978, p. 75).

**Usage**

```
data(mice)
```

**Format**

A data frame with 20 observations on the following 3 variables.

**group** a factor with levels CTRL and TREAT

**n** a numeric vector: the total number of fetuses.

**y** a numeric vector: the number of affected fetuses.

**References**

Kupper, L.L., Haseman, J.K., 1978. *The use of a correlated binomial model for the analysis of a certain toxicological experiments*. *Biometrics* 34, 69-76.

negbin

*Negative-Binomial Model for Counts***Description**

The function fits a negative-binomial log linear model accounting for overdispersion in counts  $y$ .

**Usage**

```
negbin(formula, random, data, phi.ini = NULL, warnings = FALSE,
       na.action = na.omit, fixpar = list(),
       hessian = TRUE, control = list(maxit = 2000), ...)
```

**Arguments**

formula	A formula for the fixed effects. The left-hand side of the formula must be the counts $y$ i.e., positive integers ( $y \geq 0$ ). The right-hand side can involve an offset term.
random	A right-hand formula for the overdispersion parameter(s) $\phi$ .
data	A data frame containing the response ( $y$ ) and explanatory variable(s).
phi.ini	Initial values for the overdispersion parameter(s) $\phi$ . Default to 0.1.
warnings	Logical to control printing of warnings occurring during log-likelihood maximization. Default to FALSE (no printing).
na.action	A function name. Indicates which action should be taken in the case of missing value(s).
fixpar	A list with 2 components (scalars or vectors) of the same size, indicating which parameters are fixed (i.e., not optimized) in the global parameter vector $(b, \phi)$ and the corresponding fixed values. For example, <code>fixpar = list(c(4, 5), c(0, 0))</code> means that 4th and 5th parameters of the model are set to 0.
hessian	A logical. When set to FALSE, the hessian and the variances-covariances matrices of the parameters are not computed.
control	A list to control the optimization parameters. See <a href="#">optim</a> . By default, set the maximum number of iterations to 2000.
...	Further arguments passed to <a href="#">optim</a> .

**Details**

For a given count  $y$ , the model is:

$$y \sim \lambda \sim \text{Poisson}(\lambda)$$

with  $\lambda$  following a Gamma distribution  $Gamma(r, \sim\theta)$ .  
If  $G$  denote the gamma function, then:

$$P(\lambda) = r^{-\theta} * \lambda^{\theta-1} * \frac{\exp(-\frac{\lambda}{r})}{G(\theta)}$$

$$E[\lambda] = \theta * r$$

$$Var[\lambda] = \theta * r^2$$

The marginal negative-binomial distribution is:

$$P(y) = G(y + \theta) * \left(\frac{1}{1+r}\right)^\theta * \frac{\left(\frac{r}{1+r}\right)^y}{y! * G(\theta)}$$

The function uses the parameterization  $\mu = \theta * r = \exp(Xb) = \exp(\eta)$  and  $\phi = 1/\theta$ , where  $X$  is a design-matrix,  $b$  is a vector of fixed effects,  $\eta = Xb$  is the linear predictor and  $\phi$  the overdispersion parameter.

The marginal mean and variance are:

$$E[y] = \mu$$

$$Var[y] = \mu + \phi * \mu^2$$

The parameters  $b$  and  $\phi$  are estimated by maximizing the log-likelihood of the marginal model (using the function `optim()`). Several explanatory variables are allowed in  $b$ . Only one is allowed in  $\phi$ .

An offset can be specified in the formula argument to model rates  $y/T$ . The offset and the marginal mean are  $\log(T)$  and  $\mu = \exp(\log(T) + \eta)$ , respectively.

## Value

An object of formal class “glimML”: see [glimML-class](#) for details.

## Author(s)

Matthieu Lesnoff <matthieu.lesnoff@cirad.fr>, Renaud Lancelot <renaud.lancelot@cirad.fr>

## References

Lawless, J.F., 1987. *Negative binomial and mixed Poisson regression*. The Canadian Journal of Statistics, 15(3): 209-225.

## See Also

[glimML-class](#), [glm](#) and [optim](#),  
[glm.nb](#) in the recommended package **MASS**,  
[gnlr](#) in package **gnlm** available at [www.luc.ac.be/~jlindsey/rcode.html](http://www.luc.ac.be/~jlindsey/rcode.html).

### Examples

```
# without offset
data(salmonella)
negbin(y ~ log(dose + 10) + dose, ~ 1, salmonella)
library(MASS) # function glm.nb in MASS
fm.nb <- glm.nb(y ~ log(dose + 10) + dose,
               link = log, data = salmonella)
coef(fm.nb)
1 / fm.nb$theta # theta = 1 / phi
c(logLik(fm.nb), AIC(fm.nb))
# with offset
data(dja)
negbin(y ~ group + offset(log(trisk)), ~ group, dja)
# phi fixed to zero in group TREAT
negbin(y ~ group + offset(log(trisk)), ~ group, dja,
      fixpar = list(4, 0))
# glim without overdispersion
summary(glm(y ~ group + offset(log(trisk)),
           family = poisson, data = dja))
# phi fixed to zero in both groups
negbin(y ~ group + offset(log(trisk)), ~ group, dja,
      fixpar = list(c(3, 4), c(0, 0)))
```

---

orob1

*Germination Data*

---

### Description

[Data describing the germination] “for seed *Orobancha cernua* cultivated in three dilutions of a bean root extract. The mean proportions of the three sets are 0.142, 0.872 and 0.842, and the overall mean is 0.614.” (Crowder, 1978, Table 1).

### Usage

```
data(orob1)
```

### Format

A data frame with 16 observations on the following 3 variables.

**dilution** a factor with 3 levels: 1/1, 1/25 and 1/625.

**n** a numeric vector: the number of seeds exposed to germination.

**y** a numeric vector: the number of seeds which actually germinated.

### References

Crowder, M.J., 1978. *Beta-binomial anova for proportions*. Appl. Statist. 27, 34-37.

---

 orob2

*Germination Data*


---

### Description

“A 2 x 2 factorial experiment comparing 2 types of seed and 2 root extracts. There are 5 or 6 replicates in each of the 4 treatment groups, and each replicate comprises a number of seeds varying between 4 and 81. The response variable is the proportion of seeds germinating in each replicate.” (Crowder, 1978, Table 3).

### Usage

```
data(orob2)
```

### Format

A data frame with 21 observations on the following 4 variables.

**seed** a factor with 2 levels: 073 and 075.

**root** a factor with 2 levels BEAN and CUCUMBER.

**n** a numeric vector: the number of seeds exposed to germination.

**y** a numeric vector: the number of seeds which actually germinated.

### References

Crowder, M.J., 1978. *Beta-binomial anova for proportions*. Appl. Statist. 27, 34-37.

---

 predict-methods

*Methods for Function "predict" in Package "aod"*


---

### Description

“predict” methods for fitted models generated by functions in package **aod**.

### Usage

```
## S4 method for signature 'glimML'
predict(object, newdata = NULL,
        type = c("response", "link"), se.fit = FALSE, ...)
## S4 method for signature 'glimQL'
predict(object, newdata = NULL,
        type = c("response", "link"), se.fit = FALSE, ...)
```

**Arguments**

object	A fitted model of formal class “glimML” (functions <code>betabin</code> or <code>negbin</code> ) or “glimQL” (functions <code>quasibin</code> or <code>quasipois</code> ).
newdata	A <code>data.frame</code> providing all the explanatory variables necessary for predictions.
type	A character string indicating the scale on which predictions are made: either “response” for predictions on the observation scale, or “link” for predictions on the scale of the link.
se.fit	A logical scalar indicating whether pointwise standard errors should be computed for the predictions.
...	Other arguments passed to methods.

**Methods**

**glimML** Compute predictions for models of formal class “glimML”, presently generated by functions `betabin` and `negbin`. See the examples for these functions.

**glimQL** Compute predictions for models of formal class “glimQL”, presently generated by the functions `quasibin` and `quasipois`. See the examples for these functions.

**See Also**

[predict.glm](#)

---

quasibin

*Quasi-Likelihood Model for Proportions*

---

**Description**

The function fits the generalized linear model “II” proposed by Williams (1982) accounting for overdispersion in clustered binomial data  $(n, y)$ .

**Usage**

```
quasibin(formula, data, link = c("logit", "cloglog"), phi = NULL, tol = 0.001)
```

**Arguments**

formula	A formula for the fixed effects. The left-hand side of the formula must be of the form <code>cbind(y, n - y)</code> where the modelled probability is $y/n$ .
link	The link function for the mean $p$ : “logit” or “cloglog”.
data	A data frame containing the response ( $n$ and $y$ ) and explanatory variable(s).
phi	When <code>phi</code> is <code>NULL</code> (the default), the overdispersion parameter $\phi$ is estimated from the data. Otherwise, its value is considered as fixed.
tol	A positive scalar (default to 0.001). The algorithm stops at iteration $r + 1$ when the condition $\chi^2[r + 1] - \chi^2[r] \leq tol$ is met by the $\chi^2$ statistics .

## Details

For a given cluster  $(n, y)$ , the model is:

$$y \sim \lambda \sim \text{Binomial}(n, \lambda)$$

with  $\lambda$  a random variable of mean  $E[\lambda] = p$  and variance  $\text{Var}[\lambda] = \phi * p * (1 - p)$ .

The marginal mean and variance are:

$$E[y] = p$$

$$\text{Var}[y] = p * (1 - p) * [1 + (n - 1) * \phi]$$

The overdispersion parameter  $\phi$  corresponds to the intra-cluster correlation coefficient, which is here restricted to be positive.

The function uses the function `glm` and the parameterization:  $p = h(Xb) = h(\eta)$ , where  $h$  is the inverse of a given link function,  $X$  is a design-matrix,  $b$  is a vector of fixed effects and  $\eta = Xb$  is the linear predictor.

The estimate of  $b$  maximizes the quasi log-likelihood of the marginal model. The parameter  $\phi$  is estimated with the moment method or can be set to a constant (a regular *glim* is fitted when  $\phi$  is set to zero). The literature recommends to estimate  $\phi$  from the saturated model. Several explanatory variables are allowed in  $b$ . None is allowed in  $\phi$ .

## Value

An object of formal class “glimQL”: see [glimQL-class](#) for details.

## Author(s)

Matthieu Lesnoff <matthieu.lesnoff@cirad.fr>, Renaud Lancelot <renaud.lancelot@cirad.fr>

## References

- Moore, D.F., 1987, *Modelling the extraneous variance in the presence of extra-binomial variation*. Appl. Statist. 36, 8-14.  
 Williams, D.A., 1982, *Extra-binomial variation in logistic linear models*. Appl. Statist. 31, 144-148.

## See Also

[glm](#), [geese](#) in the contributed package **geepack**, [glm.binomial.disp](#) in the contributed package **dispmod**.

## Examples

```
data(orob2)
fm1 <- glm(cbind(y, n - y) ~ seed * root,
           family = binomial, data = orob2)
fm2 <- quasibin(cbind(y, n - y) ~ seed * root,
               data = orob2, phi = 0)
fm3 <- quasibin(cbind(y, n - y) ~ seed * root,
               data = orob2)
rbind(fm1 = coef(fm1), fm2 = coef(fm2), fm3 = coef(fm3))
```

```

# show the model
fm3
# dispersion parameter and goodness-of-fit statistic
c(phi = fm3@phi,
  X2 = sum(residuals(fm3, type = "pearson")^2))
# model predictions
predfm1 <- predict(fm1, type = "response", se = TRUE)
predfm3 <- predict(fm3, type = "response", se = TRUE)
New <- expand.grid(seed = levels(orob2$seed),
  root = levels(orob2$root))
predict(fm3, New, se = TRUE, type = "response")
data.frame(orob2, p1 = predfm1$fit,
  se.p1 = predfm1$se.fit,
  p3 = predfm3$fit,
  se.p3 = predfm3$se.fit)
fm4 <- quasibin(cbind(y, n - y) ~ seed + root,
  data = orob2, phi = fm3@phi)
# Pearson's chi-squared goodness-of-fit statistic
# compare with fm3's X2
sum(residuals(fm4, type = "pearson")^2)

```

---

quasipois

*Quasi-Likelihood Model for Counts*


---

## Description

The function fits the log linear model (“Procedure II”) proposed by Breslow (1984) accounting for overdispersion in counts  $y$ .

## Usage

```
quasipois(formula, data, phi = NULL, tol = 0.001)
```

## Arguments

formula	A formula for the fixed effects. The left-hand side of the formula must be the counts $y$ i.e., positive integers ( $y \geq 0$ ). The right-hand side can involve an offset term.
data	A data frame containing the response ( $y$ ) and explanatory variable(s).
phi	When phi is NULL (the default), the overdispersion parameter $\phi$ is estimated from the data. Otherwise, its value is considered as fixed.
tol	A positive scalar (default to 0.001). The algorithm stops at iteration $r + 1$ when the condition $\chi^2[r + 1] - \chi^2[r] \leq tol$ is met by the $\chi^2$ statistics .

## Details

For a given count  $y$ , the model is:

$$y \sim \lambda \sim \text{Poisson}(\lambda)$$

with  $\lambda$  a random variable of mean  $E[\lambda] = \mu$  and variance  $\text{Var}[\lambda] = \phi * \mu^2$ .  
The marginal mean and variance are:

$$E[y] = \mu$$

$$\text{Var}[y] = \mu + \phi * \mu^2$$

The function uses the function `glm` and the parameterization:  $\mu = \exp(Xb) = \exp(\eta)$ , where  $X$  is a design-matrix,  $b$  is a vector of fixed effects and  $\eta = Xb$  is the linear predictor.

The estimate of  $b$  maximizes the quasi log-likelihood of the marginal model. The parameter  $\phi$  is estimated with the moment method or can be set to a constant (a regular *glim* is fitted when  $\phi$  is set to 0). The literature recommends to estimate  $\phi$  with the saturated model. Several explanatory variables are allowed in  $b$ . None is allowed in  $\phi$ .

An offset can be specified in the argument `formula` to model rates  $y/T$  (see examples). The offset and the marginal mean are  $\log(T)$  and  $\mu = \exp(\log(T) + \eta)$ , respectively.

## Value

An object of formal class “`glimQL`”: see [glimQL-class](#) for details.

## Author(s)

Matthieu Lesnoff <matthieu.lesnoff@cirad.fr>, Renaud Lancelot <renaud.lancelot@cirad.fr>

## References

Breslow, N.E., 1984. *Extra-Poisson variation in log-linear models*. Appl. Statist. 33, 38-44.  
Moore, D.F., Tsiatis, A., 1991. *Robust estimation of the variance in moment methods for extra-binomial and extra-poisson variation*. Biometrics 47, 383-401.

## See Also

[glm](#), [negative.binomial](#) in the recommended package **MASS**, [geese](#) in the contributed package **geepack**, [glm.poisson.disp](#) in the contributed package **dispmod**.

## Examples

```
# without offset
data(salmonella)
quasipois(y ~ log(dose + 10) + dose,
          data = salmonella)
quasipois(y ~ log(dose + 10) + dose,
          data = salmonella, phi = 0.07180449)
summary(glm(y ~ log(dose + 10) + dose,
            family = poisson, data = salmonella))
quasipois(y ~ log(dose + 10) + dose,
          data = salmonella, phi = 0)
```

```

# with offset
data(cohorts)
i <- cohorts$age ; levels(i) <- 1:7
j <- cohorts$period ; levels(j) <- 1:7
i <- as.numeric(i); j <- as.numeric(j)
cohorts$cohort <- j + max(i) - i
cohorts$cohort <- as.factor(1850 + 5 * cohorts$cohort)
fm1 <- quasipois(y ~ age + period + cohort + offset(log(n)),
                data = cohorts)

fm1
quasipois(y ~ age + cohort + offset(log(n)),
          data = cohorts, phi = fm1@phi)

```

---

rabbits

*Rabbits Foetuses Survival Experiment*


---

### Description

Experimental data for analyzing the effect of an increasing dose of a compound on the proportion of live foetuses affected. Four treatment-groups were considered: control “C”, low dose “L”, medium dose “M” and high dose “H”. The animal species used in the experiment was banded Dutch rabbit (Paul, 1982, Table 1).

### Usage

```
data(rabbits)
```

### Format

A data frame with 84 observations on the following 3 variables.

**group** a factor with levels C, H, L and M

**n** a numeric vector: the total number of foetuses.

**y** a numeric vector: the number of affected foetuses.

### References

Paul, S.R., 1982. *Analysis of proportions of affected foetuses in teratological experiments*. *Biometrics* 38, 361-370.

raoscott

*Test of Proportion Homogeneity using Rao and Scott's Adjustment***Description**

Tests the homogeneity of proportions between  $I$  groups ( $H_0: p_1 = p_2 = \dots = p_I$ ) from clustered binomial data  $(n, y)$  using the adjusted  $\chi^2$  statistic proposed by Rao and Scott (1993).

**Usage**

```
raoscott(formula = NULL, response = NULL, weights = NULL,
         group = NULL, data, pooled = FALSE, deff = NULL)
```

**Arguments**

formula	An optional formula where the left-hand side is either a matrix of the form <code>cbind(y, n-y)</code> , where the modelled probability is $y/n$ , or a vector of proportions to be modelled ( $y/n$ ). In both cases, the right-hand side must specify a single grouping variable. When the left-hand side of the formula is a vector of proportions, the argument <code>weight</code> must be used to indicate the denominators of the proportions.
response	An optional argument: either a matrix of the form <code>cbind(y, n-y)</code> , where the modelled probability is $y/n$ , or a vector of proportions to be modelled ( $y/n$ ).
weights	An optional argument used when the left-hand side of <code>formula</code> or <code>response</code> is a vector of proportions: <code>weight</code> is the denominator of the proportions.
group	An optional argument only used when <code>response</code> is used. In this case, this argument is a factor indicating a grouping variable.
data	A data frame containing the response ( $n$ and $y$ ) and the grouping variable.
pooled	Logical indicating if a pooled design effect is estimated over the $I$ groups.
deff	A numerical vector of $I$ design effects.

**Details**

The method is based on the concepts of design effect and effective sample size.

The design effect in each group  $i$  is estimated by  $deff_i = vratio_i/vbin_i$ , where  $vratio_i$  is the variance of the ratio estimate of the probability in group  $i$  (Cochran, 1999, p. 32 and p. 66) and  $vbin_i$  is the standard binomial variance. A pooled design effect (i.e., over the  $I$  groups) is estimated if argument `pooled = TRUE` (see Rao and Scott, 1993, Eq. 6). Fixed design effects can be specified with the argument `deff`.

The  $deff_i$  are used to compute the effective sample sizes  $nadj_i = n_i/deff_i$ , the effective numbers of successes  $yadj_i = y_i/deff_i$  in each group  $i$ , and the overall effective proportion  $padj = \sum_i yadj_i / \sum_i deff_i$ . The test statistic is obtained by substituting these quantities in the usual  $\chi^2$  statistic, yielding:

$$X^2 = \sum_i \frac{(yadj_i - nadj_i * padj)^2}{nadj_i * padj * (1 - padj)}$$

which is compared to a  $\chi^2$  distribution with  $I - 1$  degrees of freedom.

### Value

An object of formal class “drs”: see [drs-class](#) for details. The slot tab provides the proportion of successes, the variances of the proportion and the design effect for each group.

### Author(s)

Matthieu Lesnoff <matthieu.lesnoff@cirad.fr>, Renaud Lancelot <renaud.lancelot@cirad.fr>

### References

Cochran, W.G., 1999, 2nd ed. *Sampling techniques*. John Wiley & Sons, New York.  
 Rao, J.N.K., Scott, A.J., 1992. *A simple method for the analysis of clustered binary data*. *Biometrics* 48, 577-585.

### See Also

[chisq.test](#), [donner](#), [iccbin](#), [drs-class](#)

### Examples

```
data(rats)
# deff by group
raoscott(cbind(y, n - y) ~ group, data = rats)
raoscott(y/n ~ group, weights = n, data = rats)
raoscott(response = cbind(y, n - y), group = group, data = rats)
raoscott(response = y/n, weights = n, group = group, data = rats)
# pooled deff
raoscott(cbind(y, n - y) ~ group, data = rats, pooled = TRUE)
# standard test
raoscott(cbind(y, n - y) ~ group, data = rats, deff = c(1, 1))
data(antibio)
raoscott(cbind(y, n - y) ~ treatment, data = antibio)
```

---

rats

*Rats Diet Experiment*

---

### Description

“Weil (1970) in Table 1 gives the results from an experiment comprising two treatments. One group of 16 pregnant female rats was fed a control diet during pregnancy and lactation, the diet of a second group of 16 pregnant females was treated with a chemical. For each litter the number  $n$  of pups alive at 4 days and the number  $x$  of pups that survived the 21 day lactation period were recorded.” (Williams, 1975, p. 951).

**Usage**

```
data(rats)
```

**Format**

A data frame with 32 observations on the following 3 variables.

**group** A factor with levels CTRL and TREAT

**n** A numeric vector: the number of pups alive at 4 days.

**y** A numeric vector: the number of pups that survived the 21 day lactation.

**Source**

Williams, D.A., 1975. *The analysis of binary responses from toxicological experiments involving reproduction and teratogenicity*. Biometrics 31, 949-952.

**References**

Weil, C.S., 1970. *Selection of the valid number of sampling units and a consideration of their combination in toxicological studies involving reproduction, teratogenesis or carcinogenesis*. *Fd. Cosmet. Toxicol.* 8, 177-182.

---

residuals-methods      *Residuals for Maximum-Likelihood and Quasi-Likelihood Models*

---

**Description**

Residuals of models fitted with functions `betabin` and `negbin` (formal class “`glimML`”), or `quasibin` and `quasipois` (formal class “`glimQL`”).

**Usage**

```
## S4 method for signature 'glimML'
residuals(object, type = c("pearson", "response"), ...)
## S4 method for signature 'glimQL'
residuals(object, type = c("pearson", "response"), ...)
```

**Arguments**

**object** Fitted model of formal class “`glimML`” or “`glimQL`”.

**type** Character string for the type of residual: “`pearson`” (default) or “`response`”.

**...** Further arguments to be passed to the function, such as `na.action`.

**Details**

For models fitted with betabin or quasibin, Pearson's residuals are computed as:

$$\frac{y - n * \hat{p}}{\sqrt{n * \hat{p} * (1 - \hat{p}) * (1 + (n - 1) * \hat{\phi})}}$$

where  $y$  and  $n$  are respectively the numerator and the denominator of the response,  $\hat{p}$  is the fitted probability and  $\hat{\phi}$  is the fitted overdispersion parameter. When  $n = 0$ , the residual is set to 0. Response residuals are computed as  $y/n - \hat{p}$ .

For models fitted with negbin or quasipois, Pearson's residuals are computed as:

$$\frac{y - \hat{y}}{\sqrt{\hat{y} + \hat{\phi} * \hat{y}^2}}$$

where  $y$  and  $\hat{y}$  are the observed and fitted counts, respectively. Response residuals are computed as  $y - \hat{y}$ .

**Value**

A numeric vector of residuals.

**Author(s)**

Matthieu Lesnoff <matthieu.lesnoff@cirad.fr>, Renaud Lancelot <renaud.lancelot@cirad.fr>

**See Also**

[residuals.glm](#)

**Examples**

```
data(oro2)
fm <- betabin(cbind(y, n - y) ~ seed, ~ 1,
             link = "logit", data = oro2)
#Pearson's chi-squared goodness-of-fit statistic
sum(residuals(fm, type = "pearson")^2)
```

---

salmonella

*Salmonella Reverse Mutagenicity Assay*

---

**Description**

“Data for our third example were compiled by Margolin et al. (1981) from an Ames *Salmonella* reverse mutagenicity assay. Table 1 shows the number of revertant colonies observed on each of 3 replicate plates tested at each of 6 dose levels of quinoline.” (Breslow, 1984, Table 1).

**Usage**

```
data(salmonella)
```

**Format**

A data frame with 18 observations on the following 2 variables.

**dose** a numeric vector: the dose level of quinoline (microgram per plate).

**y** a numeric vector: the number of revertant colonies of TA98 *Salmonella*.

**Source**

Breslow, N.E., 1984. *Extra-Poisson variation in log-linear models*. Applied Statistics 33(1), 38-44.

**References**

Margolin, B.H., Kaplan, N., Zeiger, E., 1981. *Statistical analysis of the Ames Salmonella / microsome test*. Proc. Natl Acad. Sci. USA 76, 3779-3783.

---

 splitbin

*Split Grouped Data Into Individual Data*


---

**Description**

The function splits grouped data and optional covariates into individual data. Two types of grouped data are managed by splitbin:

- Grouped data with weights;
- Grouped data of form cbind(success, failure).

When weights, successes or failures involve non-integer numbers, these numbers are rounded before splitting.

**Usage**

```
splitbin(formula, data, id = "idbin")
```

**Arguments**

formula	A formula. The left-hand side represents grouped data. The right-hand side defines the covariates. See examples for syntax.
data	A data frame where all the variables described in formula are found.
id	An optional character string naming the identifier (= grouping factor). Default to "idbin".

**Value**

A data frame built according to the formula and function used in the call.

**Examples**

```
# Grouped data with weights
mydata <- data.frame(
  success = c(0, 1, 0, 1),
  f1 = c("A", "A", "B", "B"),
  f2 = c("C", "D", "C", "D"),
  n = c(4, 2, 1, 3)
)
mydata
splitbin(formula = n ~ f1, data = mydata)$tab
splitbin(formula = n ~ f1 + f2 + success, data = mydata)$tab

# Grouped data of form "cbind(success, failure)"
mydata <- data.frame(
  success = c(4, 1),
  failure = c(1, 2),
  f1 = c("A", "B"),
  f2 = c("C", "D")
)
mydata$n <- mydata$success + mydata$failure
mydata
splitbin(formula = cbind(success, failure) ~ 1, data = mydata)$tab
splitbin(formula = cbind(success, failure) ~ f1 + f2, data = mydata)$tab
splitbin(formula = cbind(success, n - success) ~ f1 + f2, data = mydata)$tab
splitbin(formula = cbind(success, n - 0.5 * failure - success) ~ f1 + f2,
  data = mydata)$tab
```

---

summary,aic-method      *Akaike Information Statistics*

---

**Description**

Computes Akaike difference and Akaike weights from an object of formal class "aic".

**Usage**

```
## S4 method for signature 'aic'
summary(object, which = c("AIC", "AICc"))
```

**Arguments**

**object**            An object of formal class "aic".

**which**            A character string indicating which information criterion is selected to compute Akaike difference and Akaike weights: either "AIC" or "AICc".

## Methods

**summary** The models are ordered according to AIC or AICc and 3 statistics are computed:

- the *Akaike difference*  $\Delta$ : the change in AIC (or AICc) between successive (ordered) models,
- the *Akaike weight*  $W$ : when  $r$  models are compared,  $W = e^{-0.5*\Delta} / \sum_r e^{-\frac{1}{2}*\Delta}$ ,
- the *cumulative Akaike weight cum.W*: the Akaike weights sum to 1 for the  $r$  models which are compared.

## References

- Burnham, K.P., Anderson, D.R., 2002. *Model selection and multimodel inference: a practical information-theoretic approach*. New-York, Springer-Verlag, 496 p.
- Hurvich, C.M., Tsai, C.-L., 1995. *Model selection for extended quasi-likelihood models in small samples*. *Biometrics*, 51 (3): 1077-1084.

## See Also

Examples in [betabin](#) and [AIC](#) in package **stats**.

---

summary.glimML-class    *Summary of Objects of Class "summary.glimML"*

---

## Description

Summary of a model of formal class "glimML" fitted by betabin or negbin.

## Objects from the Class

Objects can be created by calls of the form `new("summary.glimML", ...)` or, more commonly, via the `summary` or `show` method for objects of formal class "glimML".

## Slots

**object** An object of formal class "glimML".

**Coef** A data frame containing the estimates, standard error, z and P values for the fixed-effect coefficients which were *estimated* by the fitting function.

**FixedCoef** A data frame containing the values of the fixed-effect coefficients which were *set* to a fixed value.

**Phi** A data frame containing the estimates, standard error, z and P values for the overdispersion coefficients which were *estimated* by the fitting function. Because the overdispersion coefficients are  $> 0$ , P values correspond to unilateral tests.

**FixedPhi** A data frame containing the values of the overdispersion coefficients which were *set* to a fixed value.

## Methods

```
show signature(object = "summary.glimML")
show signature(object = "glimML")
summary signature(object = "glimML")
```

## Examples

```
data(orob2)
fm1 <- betabin(cbind(y, n - y) ~ seed, ~ 1, data = orob2)
# show objects of class "glimML"
fm1
# summary for objects of class "glimML"
res <- summary(fm1)
res@Coef
# show objects of class "summary.glimML"
res
```

---

varbin

*Mean, Variance and Confidence Interval of a Proportion*

---

## Description

This function computes the mean and variance of a proportion from clustered binomial data  $(n, y)$ , using various methods. Confidence intervals are computed using a normal approximation, which might be inappropriate when the proportion is close to 0 or 1.

## Usage

```
varbin(n, y, data, alpha = 0.05, R = 5000)
```

## Arguments

n	The denominator of the proportion.
y	The numerator of the proportion.
data	A data frame containing the data.
alpha	The significance level for the confidence intervals. Default to 0.05, providing 95% CI's.
R	The number of bootstrap replicates to compute the bootstrap mean and variance.

## Details

Five methods are used for the estimations. Let us consider  $N$  clusters of sizes  $n_1, \dots, n_N$  with observed responses (counts)  $y_1, \dots, y_N$ . We note  $p_i = y_i/n_i$  the observed proportions ( $i = 1, \dots, N$ ). An underlying assumption is that the theoretical proportion is homogeneous across the clusters.

**Binomial method:** the proportion and its variance are estimated as  $p = \frac{\sum_i y_i}{\sum_i n_i}$  and  $\frac{p*(1-p)}{\sum_i n_i - 1}$ , respectively.

**Ratio method:** the one-stage cluster sampling formula is used to estimate the variance of the ratio estimate (see Cochran, 1999, p. 32 and p. 66). The proportion is estimated as above ( $p$ ).

**Arithmetic method:** the proportion is estimated as  $p_A = \frac{1}{N} \sum_i \frac{y_i}{n_i}$ , with estimated variance  $\frac{\sum_i (p_i - p_A)^2}{N*(N-1)}$ .

**Jackknife method:** the proportion  $p_J$  is the arithmetic mean of the pseudovalues  $pv_i$ , with estimated variance  $\frac{\sum_i (pv_i - p_J)^2}{N*(N-1)}$  (Gladen, 1977, Paul, 1982).

**Bootstrap method:**  $R$  samples of size  $N$  are drawn with equal probability from the initial sample  $(p_1, \dots, p_N)$  (Efron and Tibshirani, 1993). The bootstrap estimate  $p_B$  and its estimated variance are the arithmetic mean and the empirical variance (computed with denominator  $R - 1$ ) of the  $R$  binomial estimates, respectively.

## Value

An object of formal class “varbin”, with 5 slots:

CALL	The call of the function.
tab	A 4-column data frame giving for each estimation method the mean, variance, upper and lower limits of the $(1 - \alpha)$ confidence interval.
boot	A numeric vector containing the R bootstrap replicates of the proportion. Might be used to compute other kinds of CI's for the proportion.
alpha	The significance level used to compute the $(1 - \alpha)$ confidence intervals.
features	A numeric vector with 3 components summarizing the main features of the data: $N$ = number of clusters, $n$ = number of subjects, $y$ = number of cases.

The “show” method displays the slot tab described above, substituting the standard error to the variance.

## Author(s)

Matthieu Lesnoff <matthieu.lesnoff@cirad.fr>, Renaud Lancelot <renaud.lancelot@cirad.fr>

## References

- Cochran, W.G., 1999, 3th ed. *Sampling techniques*. Wiley, New York.
- Efron, B., Tibshirani, R., 1993. *An introduction to the bootstrap*. Chapman and Hall, London.
- Gladen, B., 1977. *The use of the jackknife to estimate proportions from toxicological data in the presence of litter effects*. JASA 74(366), 278-283.
- Paul, S.R., 1982. *Analysis of proportions of affected fetuses in teratological experiments*. Biometrics 38, 361-370.

**See Also**

[varbin-class](#), [boot](#)

**Examples**

```
data(rabbits)
varbin(n, y, rabbits[rabbits$group == "M", ])
by(rabbits,
  list(group = rabbits$group),
  function(x) varbin(n = n, y = y, data = x, R = 1000))
```

---

varbin-class

*Representation of Objects of Formal Class "varbin"*


---

**Description**

Representation of the output of function `varbin` used to estimate proportions and their variance under various distribution assumptions.

**Objects from the Class**

Objects can be created by calls of the form `new("varbin", ...)` or, more commonly, via the function `varbin`.

**Slots**

`CALL` The call of the function.

`tab` A data frame containing the estimates, their variance and the confidence limits.

`pboot` A numeric vector containing the bootstrap replicates.

`alpha` The  $\alpha$  level to compute confidence intervals.

`features` A named numeric vector summarizing the design.

**Methods**

**varbin** signature(object = "varbin"): see [varbin](#).

vcov-methods

*Methods for Function "vcov" in Package "aod"***Description**

Extract the approximate var-cov matrix of estimated coefficients from fitted models.

**Methods**

**ANY** Generic function: see [vcov](#).

**glimML** Extract the var-cov matrix of estimated coefficients for fitted models of formal class “glimML”.

**glimQL** Extract the var-cov matrix of estimated coefficients for fitted models of formal class “glimQL”.

**geese** Extract the var-cov matrix of estimated coefficients for fitted models of class “geese” (contributed package **geepack**).

**geeglm** Extract the var-cov matrix of estimated coefficients for fitted objects of class “geeglm” (contributed package **geepack**).

wald.test

*Wald Test for Model Coefficients***Description**

Computes a Wald  $\chi^2$  test for 1 or more coefficients, given their variance-covariance matrix.

**Usage**

```
wald.test(Sigma, b, Terms = NULL, L = NULL, H0 = NULL,
          df = NULL, verbose = FALSE)
## S3 method for class 'wald.test'
print(x, digits = 2, ...)
```

**Arguments**

Sigma	A var-cov matrix, usually extracted from one of the fitting functions (e.g., <code>lm</code> , <code>glm</code> , ...).
b	A vector of coefficients with var-cov matrix Sigma. These coefficients are usually extracted from one of the fitting functions available in R (e.g., <code>lm</code> , <code>glm</code> , ...).
Terms	An optional integer vector specifying which coefficients should be <i>jointly</i> tested, using a Wald $\chi^2$ or <i>F</i> test. Its elements correspond to the columns or rows of the var-cov matrix given in Sigma. Default is NULL.

L	An optional matrix conformable to b, such as its product with b i.e., <code>L %*% b</code> gives the linear combinations of the coefficients to be tested. Default is NULL.
H0	A numeric vector giving the null hypothesis for the test. It must be as long as Terms or must have the same number of columns as L. Default to 0 for all the coefficients to be tested.
df	A numeric vector giving the degrees of freedom to be used in an <i>F</i> test, i.e. the degrees of freedom of the residuals of the model from which b and Sigma were fitted. Default to NULL, for no <i>F</i> test. See the section <b>Details</b> for more information.
verbose	A logical scalar controlling the amount of output information. The default is FALSE, providing minimum output.
x	Object of class "wald.test"
digits	Number of decimal places for displaying test results. Default to 2.
...	Additional arguments to print.

### Details

The key assumption is that the coefficients asymptotically follow a (multivariate) normal distribution with mean = model coefficients and variance = their var-cov matrix.

One (and only one) of Terms or L must be given. When L is given, it must have the same number of columns as the length of b, and the same number of rows as the number of linear combinations of coefficients. When df is given, the  $\chi^2$  Wald statistic is divided by  $m$  = the number of linear combinations of coefficients to be tested (i.e., `length(Terms)` or `nrow(L)`). Under the null hypothesis H0, this new statistic follows an  $F(m, df)$  distribution.

### Value

An object of class `wald.test`, printed with `print.wald.test`.

### References

- Diggle, P.J., Liang, K.-Y., Zeger, S.L., 1994. Analysis of longitudinal data. Oxford, Clarendon Press, 253 p.
- Draper, N.R., Smith, H., 1998. Applied Regression Analysis. New York, John Wiley & Sons, Inc., 706 p.

### See Also

[vcov](#)

### Examples

```
data(orob2)
fm <- quasibin(cbind(y, n - y) ~ seed * root, data = orob2)
# Wald test for the effect of root
wald.test(b = coef(fm), Sigma = vcov(fm), Terms = 3:4)
```

# Index

- \*Topic **classes**
  - aic-class, 2
  - drs-class, 15
  - glimML-class, 16
  - glimQL-class, 17
  - iccbin-class, 19
  - summary.glimML-class, 39
  - varbin-class, 42
- \*Topic **datagen**
  - splitbin, 37
- \*Topic **datasets**
  - antibio, 5
  - cohorts, 11
  - dja, 12
  - lizards, 21
  - mice, 23
  - orob1, 26
  - orob2, 27
  - rabbits, 32
  - rats, 34
  - salmonella, 36
- \*Topic **htest**
  - donner, 13
  - iccbin, 17
  - raoscott, 33
  - varbin, 40
  - wald.test, 43
- \*Topic **math**
  - invlink, 20
  - link, 21
- \*Topic **methods**
  - AIC-methods, 3
  - coef-methods, 10
  - deviance-methods, 11
  - df.residual-methods, 12
  - fitted-methods, 15
  - logLik-methods, 22
  - predict-methods, 27
  - summary,aic-method, 38
  - vcov-methods, 43
- \*Topic **package**
  - aod-pkg, 6
- \*Topic **regression**
  - anova-methods, 4
  - betabin, 7
  - negbin, 24
  - quasibin, 28
  - quasipois, 30
  - residuals-methods, 35
- AIC, 3–5, 39
- AIC,glimML-method (AIC-methods), 3
- aic-class, 2
- AIC-methods, 3
- anova,glimML-method (anova-methods), 4
- anova-methods, 4
- anova.glimML-class (anova-methods), 4
- anova.glm, 5
- antibio, 5
- aod (aod-pkg), 6
- aod-pkg, 6
- betabin, 4, 7, 16, 28, 39
- boot, 42
- chisq.test, 14, 34
- coef, 10
- coef,glimML-method (coef-methods), 10
- coef,glimQL-method (coef-methods), 10
- coef-methods, 10
- cohorts, 11
- deviance, 11
- deviance,glimML-method (deviance-methods), 11
- deviance-methods, 11
- df.residual, 12
- df.residual,glimML-method (df.residual-methods), 12

- df.residual, glimQL-method  
(df.residual-methods), 12
- df.residual-methods, 12
- dja, 12
- donner, 13, 15, 34
- drs-class, 14, 34
- drs-class, 15
- fitted, 15
- fitted, glimML-method (fitted-methods),  
15
- fitted, glimQL-method (fitted-methods),  
15
- fitted-methods, 15
- geeglm-class (vcov-methods), 43
- geese, 29, 31
- geese-class (vcov-methods), 43
- glimML-class, 9, 25
- glimML-class, 16
- glimQL-class, 29, 31
- glimQL-class, 17
- glm, 9, 25, 29, 31
- glm.binomial.disp, 29
- glm.nb, 25
- glm.poisson.disp, 31
- glmer, 19
- iccbin, 17, 20, 34
- iccbin-class, 19
- iccbin-class, 19
- invlink, 20, 21
- link, 20, 21
- lizards, 21
- logLik, 23
- logLik, glimML-method (logLik-methods),  
22
- logLik-methods, 22
- mice, 23
- negative.binomial, 31
- negbin, 16, 24, 28
- optim, 8, 9, 24, 25
- orob1, 26
- orob2, 27
- predict, glimML-method  
(predict-methods), 27
- predict, glimQL-method  
(predict-methods), 27
- predict-methods, 27
- predict.glm, 28
- print.wald.test (wald.test), 43
- quasibin, 28, 28
- quasipois, 30
- rabbits, 32
- raoscott, 14, 15, 33
- rats, 34
- residuals, glimML-method  
(residuals-methods), 35
- residuals, glimQL-method  
(residuals-methods), 35
- residuals-methods, 35
- residuals.glm, 36
- salmonella, 36
- show, aic-method (summary, aic-method), 38
- show, anova.glimML-method  
(anova-methods), 4
- show, donner-class (donner), 13
- show, drs-method (drs-class), 15
- show, glimML-class  
(summary.glimML-class), 39
- show, glimML-method (glimML-class), 16
- show, glimQL-method (glimQL-class), 17
- show, iccbin-method (iccbin), 17
- show, raoscott-class (raoscott), 33
- show, summary.glimML-method  
(summary.glimML-class), 39
- show, varbin-class (varbin), 40
- show, varbin-method (varbin-class), 42
- splitbin, 37
- summary, aic-method, 38
- summary, glimML-method  
(summary.glimML-class), 39
- summary.glimML-class, 39
- varbin, 40, 42
- varbin-class, 42
- varbin-class, 42
- vcov, 43, 44
- vcov, geeglm-method (vcov-methods), 43
- vcov, geese-method (vcov-methods), 43
- vcov, glimML-method (vcov-methods), 43
- vcov, glimQL-method (vcov-methods), 43

vcov-methods, [43](#)

wald.test, [43](#)