

# Package: MCMCglmm (via r-universe)

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**Title** MCMC Generalised Linear Mixed Models

**Depends** Matrix, coda, ape

**Imports** corpcor, tensorA, cubature, methods

**Suggests** rgl, combinat, mvtnorm, orthopolynom, MCMCpack, bayesm, msm

**Author** Jarrod Hadfield

**Maintainer** Jarrod Hadfield <j.hadfield@ed.ac.uk>

**Description** Fits Multivariate Generalised Linear Mixed Models (and related models) using Markov chain Monte Carlo techniques (Hadfield 2010 J. Stat. Soft.).

**License** GPL (>= 2)

**URL** <https://github.com/jarroddhadfield/MCMCglmm>

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

**RemoteUrl** <https://github.com/jarroddhadfield/mcmcglmm>

**RemoteRef** HEAD

**RemoteSha** dcea602900c10bf286d4e478729eaf688243c5d0

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**Description**

MCMCg1mm is a package for fitting Generalised Linear Mixed Models using Markov chain Monte Carlo techniques (Hadfield 2009). Most commonly used distributions like the normal and the Poisson are supported together with some useful but less popular ones like the zero-inflated Poisson and the multinomial. Missing values and left, right and interval censoring are accommodated for all traits. The package also supports multi-trait models where the multiple responses can follow different types of distribution. The package allows various residual and random-effect variance structures to be specified including heterogeneous variances, unstructured covariance matrices and random regression (e.g. random slope models). Three special types of variance structure that can be specified are those associated with pedigrees (animal models), phylogenies (the comparative method) and measurement error (meta-analysis).

The package makes heavy use of results in Sorensen & Gianola (2002) and Davis (2006) which taken together result in what is hopefully a fast and efficient routine. Most small to medium sized problems should take seconds to a few minutes, but large problems (> 20,000 records) are possible. My interest is in evolutionary biology so there are also several functions for applying Rice's (2004) tensor analysis to real data and functions for visualising and comparing matrices.

Please read the tutorial vignette("Tutorial", "MCMCg1mm") or the course notes vignette("CourseNotes", "MCMCg1mm")

**Author(s)**

Jarrod Hadfield <j.hadfield@ed.ac.uk>

**References**

- Hadfield, J.D. (2009) MCMC methods for Multi-response Generalised Linear Mixed Models: The MCMCg1mm R Package, *submitted*
- Sorensen, D. & Gianola, D. (2002) Likelihood, Bayesian and MCMC Methods in Quantitative Genetics, Springer
- Davis, T.A. (2006) Direct Methods for Sparse Linear Systems, SIAM
- Rice (2004) Evolutionary Theory: Mathematical and Conceptual Foundations, Sinauer

**Description**

Incidence matrix of levels within a factor

**Usage**

```
at.level(x, level)
```

**Arguments**

```
x          factor
level      factor level
```

**Value**

incidence matrix for level in x

**Author(s)**

Jarrold Hadfield <j.hadfield@ed.ac.uk>

**See Also**

[at.set](#)

**Examples**

```
fac<-gl(3,10,30, labels=letters[1:3])
x<-rnorm(30)
model.matrix(~at.level(fac,"b"):x)
```

---

at.set

*Incidence Matrix of Combined Levels within a Factor*

---

**Description**

Incidence Matrix of Combined Levels within a Factor

**Usage**

```
at.set(x, level)
```

**Arguments**

```
x          factor
level      set of factor levels
```

**Value**

incidence matrix for the set level in x

**Author(s)**

Jarrold Hadfield <j.hadfield@ed.ac.uk>

**See Also**

[at.level](#)

**Examples**

```
fac<-gl(3,10,30, labels=letters[1:3])
x<-rnorm(30)
model.matrix(~at.set(fac,2:3):x)
```

---

BTdata

*Blue Tit Data for a Quantitative Genetic Experiment*

---

**Description**

Blue Tit (*Cyanistes caeruleus*) Data for a Quantitative Genetic Experiment

**Usage**

```
data(BTdata)
```

**Format**

a data frame with 828 rows and 7 columns, with variables tarsus length (tarsus) and colour (back) measured on 828 individuals (animal). The mother of each is also recorded (dam) together with the foster nest (fosternest) in which the chicks were reared. The date on which the first egg in each nest hatched (hatchdate) is recorded together with the sex (sex) of the individuals.

**References**

Hadfield, J.D. et. al. 2007 Journal of Evolutionary Biology 20 549-557

**See Also**

[BTped](#)

---

BTped

*Blue Tit Pedigree*

---

### Description

Blue Tit (*Cyanistes caeruleus*) Pedigree

### Usage

```
data(BTped)
```

### Format

a data frame with 1040 rows and 3 columns, with individual identifier (*animal*) mother identifier (*dam*) and father identifier (*sire*). The first 212 rows are the parents of the 828 offspring from 106 full-sibling families. Parents are assumed to be unrelated to each other and have NA's in the *dam* and *sire* column.

### References

Hadfield, J.D. et. al. 2007 *Journal of Evolutionary Biology* 20 549-557

### See Also

[BTped](#)

---

buildV

*Forms expected (co)variances for GLMMs fitted with MCMCglmm*

---

### Description

Forms the expected covariance structure of link-scale observations for GLMMs fitted with MCMCglmm

### Usage

```
buildV(object, marginal=object$Random$formula, diag=TRUE, it=NULL, posterior="mean", ...)
```

**Arguments**

object	an object of class "MCMCg1mm"
marginal	formula defining random effects to be marginalised
diag	logical; if TRUE the covariances between observations are not calculated
it	integer; optional, MCMC iteration on which covariance structure should be based
posterior	character; if it is NULL should the covariance structure be based on the marginal posterior means ('mean') of the VCV parameters, or the posterior modes ('mode'), or a random draw from the posterior with replacement ('distribution'). If posterior=="all" the posterior distribution of observation variances is returned
...	Further arguments to be passed

**Value**

If diag=TRUE an n by n covariance matrix. If diag=FALSE and posterior!="all" an 1 by n matrix of variances. If posterior=="all" an nit by n matrix of variances (where nit is the number of saved MCMC iterations).

**Author(s)**

Jarrod Hadfield <j.hadfield@ed.ac.uk>

**See Also**

[MCMCg1mm](#)

---

commutation

*Commutation Matrix*

---

**Description**

Forms an mn x mn commutation matrix which transforms  $vec(\mathbf{A})$  into  $vec(\mathbf{A}')$ , where  $\mathbf{A}$  is an m x n matrix

**Usage**

commutation(m, n)

**Arguments**

m	integer; number of rows of A
n	integer; number of columns of A

**Value**

Commutation Matrix

**Author(s)**

Jarrold Hadfield <j.hadfield@ed.ac.uk>

**References**

Magnus, J. R. & Neudecker, H. (1979) Annals of Statistics 7 (2) 381-394

**Examples**

```
commutation(2,2)
```

---

dcmvnorm

*Density of a (conditional) multivariate normal variate*

---

**Description**

Density of a (conditional) multivariate normal variate

**Usage**

```
dcmvnorm(x, mean = 0, V = 1, keep=1, cond=(1:length(x))[-keep], log=FALSE)
```

**Arguments**

x	vector of observations
mean	vector of means
V	covariance matrix
keep	vector of integers: observations for which density is required
cond	vector of integers: observations to condition on
log	if TRUE, density p is given as log(p)

**Value**

numeric

**Author(s)**

Jarrold Hadfield <j.hadfield@ed.ac.uk>



**Examples**

```

V1<-cbind(c(1,0.5), c(0.5,1))
dcmvnorm(c(0,2), c(0,0), V=V1, keep=1, cond=2)
# density of x[1]=0 conditional on x[2]=2 given
# x ~ MVN(c(0,0), V1)

dcmvnorm(c(0,2), c(0,0), V=V1, keep=1, cond=NULL)
# density of x[1]=0 marginal to x[2]
dnorm(0,0,1)
# same as univariate density

V2<-diag(2)
dcmvnorm(c(0,2), c(0,0), V=V2, keep=1, cond=2)
# density of x[1]=0 conditional on x[2]=2 given
# x ~ MVN(c(0,0), V2)
dnorm(0,0,1)
# same as univariate density because V2 is diagonal

```

---

Ddivergence

*d-divergence*


---

**Description**

Calculates Ovaskainen's (2008) d-divergence between 2 zero-mean multivariate normal distributions.

**Usage**

```
Ddivergence(CA=NULL, CB=NULL, n=10000)
```

**Arguments**

CA	Matrix A
CB	Matrix B
n	number of Monte Carlo samples for approximating the integral

**Value**

d-divergence

**Note**

In versions of MCMCglmm <2.26 Ovaskainen's (2008) d-divergence was incorrectly calculated.

**Author(s)**

Jarrod Hadfield <j.hadfield@ed.ac.uk>

**References**

Ovaskainen, O. et. al. (2008) Proc. Roy. Soc - B (275) 1635 593-750

**Examples**

```
CA<-rIW(diag(2),10, n=1)
CB<-rIW(diag(2),10, n=1)
Ddivergence(CA, CB)
```

---

Dexpressions	<i>List of unevaluated expressions for (mixed) partial derivatives of fitness with respect to linear predictors.</i>
--------------	--

---

**Description**

Unevaluated expressions for (mixed) partial derivatives of fitness with respect to linear predictors for survival and fecundity.

**Usage**

Dexpressions

**Value**

PW.d0W	Fitness (W) function for the Poisson-Weibull (PW) model.
PW.d1Wds	First Partial derivative of fitness (d1W) with respect to survival (d1s) linear predictor for the Poisson-Weibull (PW) model.
PW.d1Wdf	First Partial derivative of fitness (d1W) with respect to fecundity (d1f) linear predictor for the Poisson-Weibull (PW) model.
PW.d3Wd2sd1f	Mixed third partial derivative of fitness (d3W) with 2nd derivative of survival linear predictor (d2s) and first derivative of fecundity linear predictor (d1f) from the Poisson-Weibull (PW) model.
PW.d3Wdsd2f	and so on ...
PW.d2Wd2f	
PW.d2Wd2s	
PW.d3Wd3s	
PW.d3Wd3f	

**Author(s)**

Jarrod Hadfield <j.hadfield@ed.ac.uk>

**See Also**

[Dtensor](#)

---

Dtensor	<i>Tensor of (mixed) partial derivatives</i>
---------	--

---

**Description**

Forms tensor of (mixed) partial derivatives

**Usage**

```
Dtensor(expr, name=NULL, mu = NULL, m=1, evaluate = TRUE)
```

**Arguments**

expr	'expression'
name	character vector, giving the variable names with respect to which derivatives will be computed. If NULL all variables in the expression will be used
mu	optional: numeric vector, at which the derivatives are evaluated
m	order of derivative
evaluate	logical; if TRUE the derivatives are evaluated at mu, if FALSE the derivatives are left unevaluated

**Value**

Dtensor (list) of unevaluated expression(s) if evaluate=FALSE or a tensor if evaluate=TRUE

**Author(s)**

Jarrold Hadfield [j.hadfield@ed.ac.uk](mailto:j.hadfield@ed.ac.uk)

**References**

Rice, S.H. (2004) Evolutionary Theory: Mathematical and Conceptual Foundations. Sinauer (MA) USA.

**See Also**

[evalDtensor](#), [Dexpressions](#), [D](#)

**Examples**

```
f<-expression(beta_1 + time * beta_2 + u)
Dtensor(f,eval=FALSE)
```

evalDtensor                      *Evaluates a list of (mixed) partial derivatives*

---

### **Description**

Evaluates a list of (mixed) partial derivatives

### **Usage**

```
evalDtensor(x, mu, m=1)
```

### **Arguments**

x	unevaluated (list) of expression(s)
mu	values at which the derivatives are evaluated: names need to match terms in x
m	order of derivative

### **Value**

tensor

### **Author(s)**

Jarrod Hadfield <j.hadfield@ed.ac.uk>

### **See Also**

[Dtensor](#), [D](#)

### **Examples**

```
f<-expression(beta_1 + time*beta_2+u)
Df<-Dtensor(f, eval=FALSE, m=2)
evalDtensor(Df, mu=data.frame(beta_1=0.5, beta_2=1, time=3, u=2.3))
Dtensor(f, mu=c(1,3,1,2.3), m=2)
```

---

gelman.prior                      *Prior Covariance Matrix for Fixed Effects.*

---

### Description

Prior Covariance Matrix for Fixed Effects.

### Usage

```
gelman.prior(formula, data, scale=1, intercept=scale, singular.ok=FALSE)
```

### Arguments

formula	formula for the fixed effects.
data	data.frame.
intercept	prior standard deviation for the intercept
scale	prior standard deviation for regression parameters
singular.ok	logical: if FALSE linear dependencies in the fixed effects are removed. if TRUE they are left in an estimated, although all information comes from the prior

### Details

Gelman et al. (2008) suggest that the input variables of a categorical regression are standardised and that the associated regression parameters are assumed independent in the prior. Gelman et al. (2008) recommend a scaled t-distribution with a single degree of freedom (scaled Cauchy) and a scale of 10 for the intercept and 2.5 for the regression parameters. If the degree of freedom is infinity (i.e. a normal distribution) then a prior covariance matrix  $B\Sigma V$  can be defined for the regression parameters without input standardisation that corresponds to a diagonal prior  $\mathbf{D}$  for the regression parameters had the inputs been standardised. The diagonal elements of  $\mathbf{D}$  are set to  $\text{scale}^2$  except the first which is set to  $\text{intercept}^2$ . With logistic regression  $D = \pi^2/3 + \sigma^2$  gives a prior that is approximately flat on the probability scale, where  $\sigma^2$  is the total variance due to the random effects. For probit regression it is  $D = 1 + \sigma^2$ .

### Value

prior covariance matrix

### Author(s)

Jarrod Hadfield <j.hadfield@ed.ac.uk>

### References

Gelman, A. et al. (2008) The Annals of Applied Statistics 2 4 1360-1383

**Examples**

```

dat<-data.frame(y=c(0,0,1,1), x=gl(2,2))
# data with complete separation

#####
# probit regression #
#####

prior1<-list(
  B=list(mu=c(0,0), V=gelman.prior(~x, data=dat, scale=sqrt(1+1))),
  R=list(V=1,fix=1))

m1<-MCMCg1mm(y~x, prior=prior1, data=dat, family="ordinal", verbose=FALSE)

c2<-1
p1<-pnorm(m1$Sol[,1]/sqrt(1+c2)) # marginal probability when x=1

#####
# logistic regression #
#####

prior2<-list(B=list(mu=c(0,0), V=gelman.prior(~x, data=dat, scale=sqrt(pi^2/3+1))),
  R=list(V=1,fix=1))

m2<-MCMCg1mm(y~x, prior=prior2, data=dat, family="categorical", verbose=FALSE)

c2 <- (16 * sqrt(3))/(15 * pi))^2
p2<-plogis(m2$Sol[,1]/sqrt(1+c2)) # marginal probability when x=1

plot(mcmc.list(p1,p2))

```

---

inverseA

---

*Inverse Relatedness Matrix and Phylogenetic Covariance Matrix*


---

**Description**

Henderson (1976) and Meuwissen and Luo (1992) algorithm for inverting relatedness matrices, and Hadfield and Nakagawa (2010) algorithm for inverting phylogenetic covariance matrices.

**Usage**

```

inverseA(pedigree=NULL, nodes="ALL", scale=TRUE, reduced=FALSE,
  tol = .Machine$double.eps^0.5)

```

**Arguments**

pedigree	ordered pedigree with 3 columns: id, dam and sire, or a phylo object.
nodes	"ALL" calculates the inverse for all individuals/nodes. For phylogenies "TIPS" calculates the inverse for the species tips only, and for pedigrees a vector of id's can be passed which inverts the relatedness matrix for that subset.
scale	logical: should a phylogeny (needs to be ultrametric) be scaled to unit length (distance from root to tip)?
reduced	logical: should childless nodes be dropped from the inverse and the pedigree/phylogeny representation be reduced?
tol	numeric: differences in branch length smaller than this are ignored when assessing whether a tree is ultrametric.

**Value**

Ainv	inverse as sparseMatrix
inbreeding	inbreeding coefficients/branch lengths
pedigree	pedigree/pedigree representation of phylogeny

**Author(s)**

Jarrod Hadfield <j.hadfield@ed.ac.uk>

**References**

- Henderson, C.R. (1976) *Biometrics* 32 (1) 69:83  
 Quaas, R. L. and Pollak, E. J. (1980) *Journal of Animal Science* 51:1277-1287.  
 Meuwissen, T.H.E and Luo, Z. (1992) *Genetic Selection Evolution* 24 (4) 305:313  
 Hadfield, J.D. and Nakagawa, S. (2010) *Journal of Evolutionary Biology* 23 494-508

**Examples**

```
data(bird.families)
Ainv<-inverseA(bird.families)
```

---

 knorm

*(Mixed) Central Moments of a Multivariate Normal Distribution*

---

**Description**

Forms a tensor of (mixed) central moments of a multivariate normal distribution

**Usage**

```
knorm(V, k)
```

**Arguments**

V (co)variance matrix  
k kth central moment, must be even

**Value**

tensor

**Author(s)**

Jarrod Hadfield <j.hadfield@ed.ac.uk>

**References**

Schott, J.R.(2003) Journal of Multivariate Analysis 87 (1) 177-190

**See Also**

[dnorm](#)

**Examples**

```
V<-diag(2)
knorm(V,2)
knorm(V,4)
```

---

KPPM

*Kronecker Product Permutation Matrix*

---

**Description**

Forms an  $m \times k \times m \times k$  Kronecker Product Permutation Matrix

**Usage**

```
KPPM(m, k)
```

**Arguments**

m integer  
k integer

**Value**

Matrix



**Author(s)**

Jarrold Hadfield <j.hadfield@ed.ac.uk>

**References**

Schott, J.R.(2003) Journal of Multivariate Analysis 87 (1) 177-190

**Examples**

KPPM(2,3)

---

 krzanowski.test

*Krzanowski's Comparison of Subspaces*


---

**Description**

Calculates statistics of Krzanowski's comparison of subspaces.

**Usage**

```
krzanowski.test(CA, CB, vecsA, vecsB, corr = FALSE, ...)
```

**Arguments**

CA	Matrix A
CB	Matrix B
vecsA	Vector of integers indexing the eigenvectors determining the subspace of A
vecsB	Vector of integers indexing the eigenvectors determining the subspace of B
corr	logical; if TRUE the variances of A and B are standardised
...	further arguments to be passed

**Value**

sumofS	metric for overall similarity with 0 indicating no similarity and a value of length(vecsA) for identical subspaces
angles	angle in degrees between each best matched pair of vectors
bisector	vector that lies between each best matched pair of vectors

**Author(s)**

Jarrold Hadfield <j.hadfield@ed.ac.uk>

**References**

Krzanowski, W.J. (2000) Principles of Multivariate Analysis. OUP

**Examples**

```
CA<-rIW(diag(5),10, n=1)
CB<-rIW(diag(5),10, n=1)
krzanowski.test(CA, CB, vecsA=1:2, vecsB=1:2)
krzanowski.test(CA, CA, vecsA=1:2, vecsB=1:2)
```

---

kunif

*Central Moments of a Uniform Distribution*

---

**Description**

Returns the central moments of a uniform distribution

**Usage**

```
kunif(min, max, k)
```

**Arguments**

min, max            lower and upper limits of the distribution. Must be finite.  
k                    k central moment, must be even

**Value**

kth central moment

**Author(s)**

Jarrold Hadfield <j.hadfield@ed.ac.uk>

**See Also**

[dunif](#)

**Examples**

```
kunif(-1,1,4)
y<-runif(1000,-1,1)
mean((y-mean(y))^4)
```

---

list2bdiag	<i>Forms the direct sum from a list of matrices</i>
------------	---

---

**Description**

Forms a block-diagonal matrix from a list of matrices

**Usage**

```
list2bdiag(x)
```

**Arguments**

x                    list of square matrices

**Value**

matrix

**Author(s)**

Jarrod Hadfield <j.hadfield@ed.ac.uk>

**Examples**

```
M<-list(rIW(diag(3), 10), rIW(diag(2), 10))
list2bdiag(M)
```

---

MCMCglmm	<i>Multivariate Generalised Linear Mixed Models</i>
----------	---

---

**Description**

Markov chain Monte Carlo Sampler for Multivariate Generalised Linear Mixed Models with special emphasis on correlated random effects arising from pedigrees and phylogenies (Hadfield 2010). Please read the course notes: vignette("CourseNotes", "MCMCglmm") or the overview vignette("Overview", "MCMCglmm")

**Usage**

```
MCMCglmm(fixed, random=NULL, rcov=~units, family="gaussian", mev=NULL,
data, start=NULL, prior=NULL, tune=NULL, pedigree=NULL, nodes="ALL",
scale=TRUE, nitt=13000, thin=10, burnin=3000, pr=FALSE,
pl=FALSE, verbose=TRUE, DIC=TRUE, singular.ok=FALSE, saveX=TRUE,
saveZ=TRUE, saveXL=TRUE, slice=FALSE, ginverse=NULL, trunc=FALSE,
theta_scale=NULL, saveWS=TRUE)
```

**Arguments**

fixed	<code>formula</code> for the fixed effects, multiple responses are passed as a matrix using <code>cbind</code>
random	<code>formula</code> for the random effects. Multiple random terms can be passed using the <code>+</code> operator, and in the most general case each random term has the form <code>variance.function(formula):linking.function(random.terms)</code> . Currently, the only <code>variance.functions</code> available are <code>idv</code> , <code>idh</code> , <code>us</code> , <code>cor[]</code> and <code>ante[]</code> . <code>idv</code> fits a constant variance across all components in <code>formula</code> . Both <code>idh</code> and <code>us</code> fit different variances across each component in <code>formula</code> , but <code>us</code> will also fit the covariances. <code>corg</code> fixes the variances along the diagonal to one and <code>corgh</code> fixes the variances along the diagonal to those specified in the prior. <code>cors</code> allows correlation submatrices. <code>ante[]</code> fits ante-dependence structures of different order (e.g <code>ante1</code> , <code>ante2</code> ), and the number can be prefixed by a <code>c</code> to hold all regression coefficients of the same order equal. The number can also be suffixed by a <code>v</code> to hold all innovation variances equal (e.g <code>antec2v</code> has 3 parameters). The <code>formula</code> can contain both factors and numeric terms (i.e. random regression) although it should be noted that the intercept term is suppressed. The (co)variances are the (co)variances of the <code>random.terms</code> effects. Currently, the only <code>linking.functions</code> available are <code>mm</code> and <code>str</code> . <code>mm</code> fits a multimembership model where multiple random terms are separated by the <code>+</code> operator. <code>str</code> allows covariances to exist between multiple random terms that are also separated by the <code>+</code> operator. In both cases the levels of all multiple random terms have to be the same. For simpler models the <code>variance.function(formula)</code> and <code>linking.function(random.terms)</code> can be omitted and the model syntax has the simpler form <code>~random1+random2+...</code> . There are two reserved variables: <code>units</code> which index rows of the response variable and <code>trait</code> which index columns of the response variable
rcov	<code>formula</code> for residual covariance structure. This has to be set up so that each data point is associated with a unique residual. For example a multi-response model might have the R-structure defined by <code>~us(trait):units</code>
family	optional character vector of trait distributions. Currently, "gaussian", "poisson", "categorical", "multinomial", "ordinal", "threshold", "exponential", "geometric", "cengaussian", "cenpoisson", "cenexponential", "zipoisson", "zapoisson", "ztpoisson", "hupoisson", "zibinomial", "threshold", "nzbinom", "ncst", "msst", "hubinomial", "ztmb" and "ztmultinomial" are supported, where the prefix "cen" means censored, the prefix "zi" means zero inflated, the prefix "za" means zero altered, the prefix "zt" means zero truncated and the prefix "hu" means hurdle. If NULL, data needs to contain a family column.
mev	optional vector of measurement error variances for each data point for random effect meta-analysis.
data	<code>data.frame</code>
start	optional list having 5 possible elements: R (R-structure) G (G-structure) and Liab (latent variables or liabilities) should contain the starting values where G itself is also a list with as many elements as random effect components. The element QUASI should be logical: if TRUE starting latent variables are obtained

heuristically, if FALSE then they are sampled from a Z-distribution. The element  $r$  should be between -1 and 1 and determines the correlation between the starting latent variables and the ordered latent variables (ordered by the response variable): the default is 0.8.

prior	optional list of prior specifications having 4 possible elements: R (R-structure) G (G-structure), B (fixed effects) and S (theta_scale parameter). B and S are lists containing the expected value ( $\mu$ ) and a (co)variance matrix (V) representing the strength of belief: the defaults are $B\mu=S\mu=0$ and $BV=SV=I*1e+10$ , where where I is an identity matrix of appropriate dimension. The priors for the variance structures (R and G) are lists with the expected (co)variances (V) and degree of belief parameter ( $\nu$ ) for the inverse-Wishart, and also the mean vector ( $\alpha.\mu$ ) and covariance matrix ( $\alpha.V$ ) for the redundant working parameters. The defaults are $\nu=0$ , $V=1$ , $\alpha.\mu=0$ , and $\alpha.V=0$ . When $\alpha.V$ is non-zero, parameter expanded algorithms are used.
tune	optional list with elements mh_V and/or mh_weights mh_V should be a list with as many elements as there are R-structure terms with each element being the (co)variance matrix defining the proposal distribution for the associated latent variables. If NULL an adaptive algorithm is used which ceases to adapt once the burn-in phase has finished. mh_weights should be equal to the number of latent variables and acts as a scaling factor for the proposal standard deviations.
pedigree	ordered pedigree with 3 columns id, dam and sire or a phylo object. This argument is retained for back compatibility - see ginverse argument for a more general formulation.
nodes	pedigree/phylogeny nodes to be estimated. The default, "ALL" estimates effects for all individuals in a pedigree or nodes in a phylogeny (including ancestral nodes). For phylogenies "TIPS" estimates effects for the tips only, and for pedigrees a vector of ids can be passed to nodes specifying the subset of individuals for which animal effects are estimated. Note that all analyses are equivalent if omitted nodes have missing data but by absorbing these nodes the chain mix better. However, the algorithm may be less numerically stable and may iterate slower, especially for large phylogenies.
scale	logical: should the phylogeny (needs to be ultrametric) be scaled to unit length (distance from root to tip)?
nitt	number of MCMC iterations
thin	thinning interval
burnin	burnin
pr	logical: should the posterior distribution of random effects be saved?
pl	logical: should the posterior distribution of latent variables be saved?
verbose	logical: if TRUE MH diagnostics are printed to screen
DIC	logical: if TRUE deviance and deviance information criterion are calculated
singular.ok	logical: if FALSE linear dependencies in the fixed effects are removed. if TRUE they are left in an estimated, although all information comes from the prior
saveX	logical: save fixed effect design matrix
saveZ	logical: save random effect design matrix

saveXL	logical: save structural parameter design matrix
slice	logical: should slice sampling be used? Only applicable for binary traits with independent residuals
ginverse	a list of sparse inverse matrices ( $\mathbf{A}^{-1}$ ) that are proportional to the covariance structure of the random effects. The names of the matrices should correspond to columns in data that are associated with the random term. All levels of the random term should appear as rownames for the matrices.
trunc	logical: should latent variables in binary models be truncated to prevent under/overflow (+/-20 for categorical/multinomial models and +/-7 for threshold/probit models)?
theta_scale	optional list of 4 possible elements specifying a set of location effects (fixed or random) that are to be scaled by the parameter theta_scale for the subset of observations which have level level in factor factor: factor, level, fixed (position of fixed terms to be scaled) and random (position of random effect components).
saveWS	logical: save design matrix for scaled effects.

**Value**

Sol	Posterior Distribution of MME solutions, including fixed effects
VCV	Posterior Distribution of (co)variance matrices
CP	Posterior Distribution of cut-points from an ordinal model
Liab	Posterior Distribution of latent variables
Fixed	list: fixed formula and number of fixed effects
Random	list: random formula, dimensions of each covariance matrix, number of levels per covariance matrix, and term in random formula to which each covariance belongs
Residual	list: residual formula, dimensions of each covariance matrix, number of levels per covariance matrix, and term in residual formula to which each covariance belongs
Deviance	deviance $-2*\log(p(y ...))$
DIC	deviance information criterion
X	sparse fixed effect design matrix
Z	sparse random effect design matrix
XL	sparse structural parameter design matrix
error.term	residual term for each datum
family	distribution of each datum
Tune	(co)variance matrix of the proposal distribution for the latent variables
meta	logical; was mev passed?
Wscale	sparse design matrix for scaled terms.

**Author(s)**

Jarrod Hadfield <j.hadfield@ed.ac.uk>

## References

- General analyses: Hadfield, J.D. (2010) *Journal of Statistical Software* 33 2 1-22
- Phylogenetic analyses: Hadfield, J.D. & Nakagawa, S. (2010) *Journal of Evolutionary Biology* 23 494-508
- Background Sorensen, D. & Gianola, D. (2002) Springer

## See Also

[mcmc](#)

## Examples

```
# Example 1: univariate Gaussian model with standard random effect

data(PlodiaPO)
model1<-MCMCgImm(PO~1, random=~FSfamily, data=PlodiaPO, verbose=FALSE,
  nitt=1300, burnin=300, thin=1)
summary(model1)

# Example 2: univariate Gaussian model with phylogenetically correlated
# random effect

data(bird.families)

phylo.effect<-rbv(bird.families, 1, nodes="TIPS")
phenotype<-phylo.effect+rnorm(dim(phylo.effect)[1], 0, 1)

# simulate phylogenetic and residual effects with unit variance

test.data<-data.frame(phenotype=phenotype, taxon=row.names(phenotype))

Ainv<-inverseA(bird.families)$Ainv

# inverse matrix of shared phylogenetic history

prior<-list(R=list(V=1, nu=0.002), G=list(G1=list(V=1, nu=0.002)))

model2<-MCMCgImm(phenotype~1, random=~taxon, ginverse=list(taxon=Ainv),
  data=test.data, prior=prior, verbose=FALSE, nitt=1300, burnin=300, thin=1)

plot(model2$VCV)
```

## Description

Sets up design matrix for measurement error models.

**Usage**

```
me(formula, error=NULL, group=NULL, type="classical")
```

**Arguments**

formula	formula for the fixed effects.
error	character; name of column in <code>data.frame</code> in which standard error (type="classical" or type="berkson") or miscalssification error (type="dclassical") is stored.
group	name of column in <code>data.frame</code> in which groups are stored. Rows of the design matrix with the same group level are assumed to pertain to the same obsevation of the covariate that is measured with error.
type	character; one of type="classical", type="berkson", type="dclassical" or type="dberkson" (see details)

**Value**

design matrix, with a prior distribution attribute

**Author(s)**

Jarrod Hadfield <j.hadfield@ed.ac.uk>

---

mult.memb

*Design Matrices for Multiple Membership Models*

---

**Description**

Forms design matrices for multiple membership models

**Usage**

```
mult.memb(formula)
```

**Arguments**

formula	formula
---------	---------

**Details**

Currently `mult.memb` can only usefully be used inside an `idv` variance function. The formula usually contains several factors that have the same factor levels.

**Value**

design matrix



**Author(s)**

Jarrod Hadfield <j.hadfield@ed.ac.uk>

**Examples**

```
fac1<-factor(sample(letters[1:3], 5, TRUE), levels=letters[1:3])
fac2<-factor(sample(letters[1:3], 5, TRUE), levels=letters[1:3])
cbind(fac1, fac2)
mult.memb(~fac1+fac2)
```

---

path

*Design Matrix for Path Analyses*

---

**Description**

Forms design matrix for path analyses that involve paths within residual blocks

**Usage**

```
path(cause=NULL, effect=NULL, k)
```

**Arguments**

cause	integer; index of predictor ‘trait’ within residual block
effect	integer; index of response ‘trait’ within residual block
k	integer; dimension of residual block

**Value**

design matrix

**Note**

For more general path analytic models see [sir](#) which allows paths to exist between responses that are not in the same residual block. However, [sir](#) does not handle non-Gaussian or missing responses. Note that path models involving non-Gaussian data are defined on the link scale which may not always be appropriate.

**Author(s)**

Jarrod Hadfield <j.hadfield@ed.ac.uk>

**See Also**

[sir](#)

**Examples**

```
path(1, 2,2)
```

pkk

*Probability that all multinomial categories have a non-zero count.*

---

**Description**

Calculates the probability that all categories in a multinomial have a non-zero count.

**Usage**

```
pkk(prob, size)
```

**Arguments**

prob	numeric non-negative vector of length K, specifying the probability for the K classes; is internally normalized to sum 1. Infinite and missing values are not allowed.
size	integer, say N, specifying the total number of objects that are put into K boxes in the typical multinomial experiment.

**Value**

probability that there is at least one object in each of the K boxes

**Author(s)**

Jarrod Hadfield <j.hadfield@ed.ac.uk>

**Examples**

```
p<-runif(4)
pkk(p, 10)
```

---

PlodiaPO*Phenoloxidase measures on caterpillars of the Indian meal moth.*

---

**Description**

Phenoloxidase measures on caterpillars of the Indian meal moth (*Plodia interpunctella*).

**Usage**

```
data(PlodiaPO)
```

**Format**

a data frame with 511 rows and 3 columns, with variables indicating full-sib family (FSfamily), phenoloxidase measures (PO), and plate (plate). PO has undergone a Box-Cox power transformation of 0.141

**Source**

Tidbury H & Boots M (2007) University of Sheffield

**See Also**

[PlodiaR](#), [PlodiaRB](#)

---

PlodiaR

*Resistance of Indian meal moth caterpillars to the granulosis virus PiGV.*

---

**Description**

Resistance of Indian meal moth (*Plodia interpunctella*) caterpillars to the granulosis virus PiGV.

**Usage**

```
data(PlodiaR)
```

**Format**

a data frame with 50 rows and 5 columns, with variables indicating full-sib family (FSfamily), date of egg laying (date\_laid) and assaying (date\_Ass), and the number of individuals from the family that were experimentally infected with the virus Infected and the number of those that pupated Pupated. These full-sib family identifiers also relate to the full-sib family identifiers in PlodiaPO

**Source**

Tidbury H & Boots M (2007) University of Sheffield

**See Also**

[PlodiaRB](#), [PlodiaPO](#)

---

PlodiaRB	<i>Resistance (as a binary trait) of Indian meal moth caterpillars to the granulosis virus PiGV.</i>
----------	--

---

**Description**

Resistance (as a binary trait) of Indian meal moth (*Plodia interpunctella*) caterpillars to the granulosis virus PiGV.

**Usage**

```
data(PlodiaRB)
```

**Format**

a data frame with 784 rows and 4 columns, with variables indicating full- sib family (FSfamily), date of egg laying (date\_laid) and assaying (date\_Ass), and a binary variable indicating whether an individual was resistant (Pupated) to an experimental infection of the virus. These data are identical to those in the data.frame PlodiaR except each family-level binomial variable has been expanded into a binary variable for each individual.

**Source**

Tidbury H & Boots M (2007) University of Sheffield

**See Also**

[PlodiaR](#), [PlodiaPO](#)

---

plot.MCMCglmm	<i>Plots MCMC chains from MCMCglmm using plot.mcmc</i>
---------------	--

---

**Description**

plot method for class "MCMCglmm".

**Usage**

```
## S3 method for class 'MCMCglmm'
plot(x, random=FALSE, ...)
```

**Arguments**

x	an object of class "MCMCglmm"
random	logical; should saved random effects be plotted
...	Further arguments to be passed

**Author(s)**

Jarrold Hadfield <j.hadfield@ed.ac.uk>

**See Also**

plot.mcmc, [MCMCglmm](#)

---

plotsubspace

*Plots covariance matrices*

---

**Description**

Represents covariance matrices as 3-d ellipsoids using the `rgl` package. Covariance matrices of dimension greater than 3 are plotted on the subspace defined by the first three eigenvectors.

**Usage**

```
plotsubspace(CA, CB=NULL, corr = FALSE, shadeCA = TRUE,  
             shadeCB = TRUE, axes.lab = FALSE, ...)
```

**Arguments**

CA	Matrix
CB	Optional second matrix
corr	If TRUE the covariance matrices are transformed into correlation matrices
shadeCA	If TRUE the ellipsoid is solid, if FALSE the ellipsoid is wireframe
shadeCB	If TRUE the ellipsoid is solid, if FALSE the ellipsoid is wireframe
axes.lab	If TRUE the axes are labelled with the eigenvectors
...	further arguments to be passed

**Details**

The matrix `CA` is always red, and the matrix `CB` if given is always blue. The subspace is defined by the first three eigenvectors of `CA`, and the percentage of variance for each matrix along these three dimensions is given in the plot title.

**Author(s)**

Jarrold Hadfield <j.hadfield@ed.ac.uk> with code taken from the `rgl` package

**See Also**

[rgl](#)

**Examples**

```
if(requireNamespace("rgl")!=FALSE){
  G1<-rIW(diag(4),10)
  G2<-G1*1.2
  # plotsubspace(G1, G2, shadeCB=FALSE)
  # commented out because of problems with rgl
}
```

---

posterior.ante

*Posterior distribution of ante-dependence parameters*

---

**Description**

Posterior distribution of ante-dependence parameters

**Usage**

```
posterior.ante(x,k=1)
```

**Arguments**

x	mcmc object of (co)variances stacked column-wise
k	order of the ante-dependence structure

**Value**

posterior ante-dependence parameters (innovation variances followed by regression coefficients)

**Author(s)**

Jarrod Hadfield <j.hadfield@ed.ac.uk>

**See Also**

[posterior.cor](#), [posterior.evals](#), [posterior.inverse](#)

**Examples**

```
v<-rIW(diag(2),10, n=1000)
plot(posterior.ante(mcmc(v),1))
```

---

posterior.cor	<i>Transforms posterior distribution of covariances into correlations</i>
---------------	---

---

**Description**

Transforms posterior distribution of covariances into correlations

**Usage**

```
posterior.cor(x)
```

**Arguments**

x                    mcmc object of (co)variances stacked column-wise

**Value**

posterior correlation matrices

**Author(s)**

Jarrod Hadfield <j.hadfield@ed.ac.uk>

**See Also**

[posterior.evals](#), [posterior.inverse](#), [posterior.ante](#)

**Examples**

```
v<-rIW(diag(2),3, n=1000)
hist(posterior.cor(mcmc(v))[,2])
```

---

posterior.evals	<i>Posterior distribution of eigenvalues</i>
-----------------	--

---

**Description**

Posterior distribution of eigenvalues

**Usage**

```
posterior.evals(x)
```

**Arguments**

x                    mcmc object of (co)variances stacked column-wise

**Value**

posterior eigenvalues

**Author(s)**

Jarrold Hadfield <j.hadfield@ed.ac.uk>

**See Also**

[posterior.cor](#), [posterior.inverse](#), [posterior.ante](#)

**Examples**

```
v<-rIW(diag(2),3, n=1000)
hist(posterior.ivals(mcmc(v))[,2])
```

---

posterior.inverse	<i>Posterior distribution of matrix inverse</i>
-------------------	---

---

**Description**

Posterior distribution of matrix inverse

**Usage**

```
posterior.inverse(x)
```

**Arguments**

x                    mcmc object of (co)variances stacked column-wise

**Value**

posterior of inverse (co)variance matrices

**Author(s)**

Jarrold Hadfield <j.hadfield@ed.ac.uk>

**See Also**

[posterior.cor](#), [posterior.ivals](#), [posterior.ante](#)

**Examples**

```
v<-rIW(diag(2),3, n=1000)
plot(posterior.inverse(mcmc(v)))
```



---

posterior.mode	<i>Estimates the marginal parameter modes using kernel density estimation</i>
----------------	---

---

### Description

Estimates the marginal parameter modes using kernel density estimation

### Usage

```
posterior.mode(x, adjust=0.1, ...)
```

### Arguments

x	mcmc object
adjust	numeric, passed to <a href="#">density</a> to adjust the bandwidth of the kernel density
...	other arguments to be passed

### Value

modes of the kernel density estimates

### Author(s)

Jarrod Hadfield <j.hadfield@ed.ac.uk>

### See Also

[density](#)

### Examples

```
v<-rIW(as.matrix(1),10, n=1000)
hist(v)
abline(v=posterior.mode(mcmc(v)), col="red")
```

---

predict.MCMCglmm      *Predict method for GLMMs fitted with MCMCglmm*

---

### Description

Predicted values for GLMMs fitted with MCMCglmm

### Usage

```
## S3 method for class 'MCMCglmm'
predict(object, newdata=NULL, marginal=object$Random$formula,
        type="response", interval="none", level=0.95, it=NULL,
        posterior="all", verbose=FALSE, approx="numerical", ...)
```

### Arguments

object	an object of class "MCMCglmm"
newdata	An optional data frame in which to look for variables with which to predict
marginal	formula defining random effects to be marginalised
type	character; either "terms" (link scale) or "response" (data scale)
interval	character; either "none", "confidence" or "prediction"
level	A numeric scalar in the interval (0,1) giving the target probability content of the intervals.
it	integer; optional, MCMC iteration on which predictions should be based
posterior	character; should marginal posterior predictions be calculated ("all"), or should they be made conditional on the marginal posterior means ("mean") of the parameters, the posterior modes ("mode"), or a random draw from the posterior ("distribution").
verbose	logical; if TRUE, warnings are issued with newdata when the original model has fixed effects that do not appear in newdata and/or newdata has random effects not present in the original model.
approx	character; for distributions for which the mean cannot be calculated analytically what approximation should be used: numerical integration (numerical; slow), second order Taylor expansion (taylor2) and for logistic models approximations presented in Diggle (2004) (diggle) and McCulloch and Searle (2001) (mcculloch)
...	Further arguments to be passed

### Value

Expectation and credible interval

### Author(s)

Jarrod Hadfield <j.hadfield@ed.ac.uk>

**References**

- Diggle P, et al. (2004). Analysis of Longitudinal Data. 2nd Edition. Oxford University Press.  
 McCulloch CE and Searle SR (2001). Generalized, Linear and Mixed Models. John Wiley & Sons, New York.

**See Also**

[MCMCg1mm](#)

---

 prunePed

*Pedigree pruning*


---

**Description**

Creates a subset of a pedigree by retaining the ancestors of a specified subset of individuals

**Usage**

```
prunePed(pedigree, keep, make.base=FALSE)
```

**Arguments**

pedigree	pedigree with id in column 1 dam in column 2 and sire in column 3
keep	individuals in pedigree for which the ancestors should be retained
make.base	logical: should ancestors that do not provide additional information be discarded?

**Value**

subsetting pedigree

**Note**

If the individuals in keep are the only phenotyped individuals for some analysis then some non-phenotyped individuals can often be discarded if they are not responsible for pedigree links between phenotyped individuals. In the simplest case (make.base=FALSE) all ancestors of phenotyped individuals will be retained, although further pruning may be possible using make.base=TRUE. In this case all pedigree links that do not connect phenotyped individuals are discarded resulting in some individuals becoming part of the base population. In terms of variance component and fixed effect estimation pruning the pedigree should have no impact on the target posterior distribution, although convergence and mixing may be better because there is less missing data.

**Author(s)**

Jarrod Hadfield <j.hadfield@ed.ac.uk> + Michael Morrissey

---

 Ptensor

*Tensor of Sample (Mixed) Central Moments*


---

**Description**

Forms a tensor of sample (mixed) central moments

**Usage**

Ptensor(x, k)

**Arguments**

x                    matrix; traits in columns samples in rows  
 k                    kth central moment

**Value**

tensor

**Author(s)**

Jarrod Hadfield <j.hadfield@ed.ac.uk>

**Examples**

```
n<-1000
y<-matrix(rnorm(n), n/2, 2)
Ptensor(y,2)
cov(y)*((n-1)/n)
```

---

 rbv

*Random Generation of MVN Breeding Values and Phylogenetic Effects*


---

**Description**

Random Generation of MVN Breeding Values and Phylogenetic Effects

**Usage**

rbv(pedigree, G, nodes="ALL", scale=TRUE, ggroups=NULL, gmeans=NULL)

**Arguments**

pedigree	ordered pedigree with 3 columns id, dam and sire or a phylo object.
G	(co)variance matrix
nodes	effects for pedigree/phylogeny nodes to be returned. The default, nodes="ALL" returns effects for all individuals in a pedigree or nodes in a phylogeny (including ancestral nodes). For phylogenies nodes="TIPS" returns effects for the tips only, and for pedigrees a vector of ids can be passed to nodes specifying the subset of individuals for which animal effects are returned.
scale	logical: should a phylogeny (needs to be ultrametric) be scaled to unit length (distance from root to tip)?
ggroups	optional; vector of genetic groups
gmeans	matrix of mean breeding value for genetic groups (rows) by traits (columns)

**Value**

matrix of breeding values/phylogenetic effects

**Author(s)**

Jarrold Hadfield <j.hadfield@ed.ac.uk>

**Examples**

```
data(bird.families)
bv<-rbv(bird.families, diag(2))
```

---

residuals.MCMCglmm      *Residuals form a GLMM fitted with MCMCglmm*

---

**Description**

residuals method for class "MCMCglmm".

**Usage**

```
## S3 method for class 'MCMCglmm'
residuals(object, type = c("deviance", "pearson", "working",
                          "response", "partial"), ...)
```

**Arguments**

object	an object of class "MCMCglmm"
type	the type of residuals which should be returned. The alternatives are: "deviance" (default), "pearson", "working", "response", and "partial".
...	Further arguments to be passed

**Value**

vector of residuals

**Author(s)**

Jarrod Hadfield <j.hadfield@ed.ac.uk>

**See Also**

[residuals](#), [MCMCglmm](#)

---

rIW

*Random Generation from the Conditional Inverse Wishart Distribution*

---

**Description**

Samples from the inverse Wishart distribution, with the possibility of conditioning on a diagonal submatrix

**Usage**

```
rIW(V, nu, fix=NULL, n=1, CM=NULL)
```

**Arguments**

V	Expected (co)varaince matrix as nu tends to infinity
nu	degrees of freedom
fix	optional integer indexing the partition to be conditioned on
n	integer: number of samples to be drawn
CM	matrix: optional matrix to condition on. If not given, and fix!=NULL, V <sub>22</sub> is conditioned on

**Details**

If  $\mathbf{W}^{-1}$  is a draw from the inverse Wishart, `fix` indexes the diagonal element of  $\mathbf{W}^{-1}$  which partitions  $\mathbf{W}^{-1}$  into 4 submatrices. `fix` indexes the upper left corner of the lower diagonal matrix and it is this matrix that is conditioned on.

For example partitioning  $\mathbf{W}^{-1}$  such that

$$\mathbf{W}^{-1} = \begin{bmatrix} \mathbf{W}^{-1}_{11} & \mathbf{W}^{-1}_{12} \\ \mathbf{W}^{-1}_{21} & \mathbf{W}^{-1}_{22} \end{bmatrix}$$

`fix` indexes the upper left corner of  $\mathbf{W}^{-1}_{22}$ . If `CM!=NULL` then  $\mathbf{W}^{-1}_{22}$  is fixed at `CM`, otherwise  $\mathbf{W}^{-1}_{22}$  is fixed at `V22`. For example, if `dim(V)=4` and `fix=2` then  $\mathbf{W}^{-1}_{11}$  is a 1X1 matrix and  $\mathbf{W}^{-1}_{22}$  is a 3X3 matrix.

**Value**

if  $n = 1$  a matrix equal in dimension to  $V$ , if  $n > 1$  a matrix of dimension  $n \times \text{length}(V)$

**Note**

In versions of MCMCglmm  $> 1.10$  the arguments to `rIW` have changed so that they are more intuitive in the context of MCMCglmm. Following the notation of Wikipedia ([https://en.wikipedia.org/wiki/Inverse-Wishart\\_distribution](https://en.wikipedia.org/wiki/Inverse-Wishart_distribution)) the inverse scale matrix  $\Psi = (V * \nu)$ . In earlier versions of MCMCglmm ( $< 1.11$ )  $\Psi = V^{-1}$ . Although the old parameterisation is consistent with the `riwish` function in MCMCpack and the `rwishart` function in bayesm it is inconsistent with the prior definition for MCMCglmm. The following pieces of code are sampling from the same distributions:

<code>riwish(nu, nu*V)</code>	from MCMCpack
<code>rwishart(nu, solve(nu*V))\$IW</code>	from bayesm
<code>rIW(nu, solve(nu*V))</code>	from MCMCglmm $< 1.11$
<code>rIW(V, nu)</code>	from MCMCglmm $\geq 1.11$

**Author(s)**

Jarrold Hadfield <j.hadfield@ed.ac.uk>

**References**

Korsgaard, I.R. et. al. 1999 Genetics Selection Evolution 31 (2) 177:181

**See Also**

`rwishart`, `rwish`

**Examples**

```
nu<-10
V<-diag(4)
rIW(V, nu, fix=2)
```

---

rtcmvnorm

*Random Generation from a Truncated Conditional Normal Distribution*

---

**Description**

Samples from the Truncated Conditional Normal Distribution

**Usage**

```
rtcmvnorm(n = 1, mean = 0, V = 1, x=0, keep=1, lower = -Inf, upper = Inf)
```

**Arguments**

n	integer: number of samples to be drawn
mean	vector of means
V	covariance matrix
x	vector of observations to condition on
keep	element of x to be sampled
lower	left truncation point
upper	right truncation point

**Value**

vector

**Author(s)**

Jarrold Hadfield <j.hadfield@ed.ac.uk>

**Examples**

```
par(mfrow=c(2,1))
V1<-cbind(c(1,0.5), c(0.5,1))
x1<-rtcmvnorm(10000, c(0,0), V=V1, c(0,2), keep=1, lower=-1, upper=1)
x2<-rtnorm(10000, 0, 1, lower=-1, upper=1)
plot(density(x1), main="Correlated conditioning observation")
lines(density(x2), col="red")
# denisties of conditional (black) and unconditional (red) distribution
# when the two variables are correlated (r=0.5)

V2<-diag(2)
x3<-rtcmvnorm(10000, c(0,0), V=V2, c(0,2), keep=1, lower=-1, upper=1)
x4<-rtnorm(10000, 0, 1, lower=-1, upper=1)
plot(density(x3), main="Uncorrelated conditioning observation")
lines(density(x4), col="red")
# denisties of conditional (black) and unconditional (red) distribution
# when the two variables are uncorrelated (r=0)
```

---

rtnorm

*Random Generation from a Truncated Normal Distribution*

---

**Description**

Samples from the Truncated Normal Distribution

**Usage**

```
rtnorm(n = 1, mean = 0, sd = 1, lower = -Inf, upper = Inf)
```



**Arguments**

n	integer: number of samples to be drawn
mean	vector of means
sd	vector of standard deviations
lower	left truncation point
upper	right truncation point

**Value**

vector

**Author(s)**

Jarrod Hadfield <j.hadfield@ed.ac.uk>

**References**

Robert, C.P. (1995) *Statistics & Computing* 5 121-125

**See Also**

[rtnorm](#)

**Examples**

```
hist(rtnorm(100, lower=-1, upper=1))
```

---

simulate.MCMCglmm      *Simulate method for GLMMs fitted with MCMCglmm*

---

**Description**

Simulated response vectors for GLMMs fitted with MCMCglmm

**Usage**

```
## S3 method for class 'MCMCglmm'  
simulate(object, nsim = 1, seed = NULL, newdata=NULL, marginal = object$Random$formula,  
         type = "response", it=NULL, posterior = "all", verbose=FALSE, ...)
```

**Arguments**

object	an object of class "MCMCg1mm"
nsim	number of response vectors to simulate. Defaults to 1.
seed	Either NULL or an integer that will be used in a call to <code>set.seed</code> before simulating the response vectors. The default, NULL will not change the random generator state.
newdata	An optional data frame for which to simulate new observations
marginal	formula defining random effects to be marginalised
type	character; either "terms" (link scale) or "response" (data scale)
it	integer; optional, MCMC iteration on which predictions should be based
posterior	character; if it is NULL should the response vector be simulated using the marginal posterior means ("mean") of the parameters, or the posterior modes ("mode"), random draws from the posterior with replacement ("distribution") or without replacement ("all")
verbose	logical; if TRUE, warnings are issued with newdata when the original model has fixed effects that do not appear in newdata and/or newdata has random effects not present in the original model.
...	Further arguments to be passed

**Value**

A matrix (with nsim columns) of simulated response vectors

**Author(s)**

Jarrod Hadfield <j.hadfield@ed.ac.uk>

**See Also**

[MCMCg1mm](#)

---

sir	<i>Design Matrix for Simultaneous and Recursive Relationships between Responses</i>
-----	---

---

**Description**

Forms design matrix for simultaneous and recursive relationships between responses

**Usage**

```
sir(formula1=NULL, formula2=NULL, diag0=FALSE)
```

**Arguments**

formula1	formula
formula2	formula
diag0	logical: should the design matrix have zero's along the diagonal

**Value**

design matrix

**Author(s)**

Jarrold Hadfield <j.hadfield@ed.ac.uk>

**Examples**

```
fac1<-factor(sample(letters[1:3], 5, TRUE), levels=letters[1:3])
fac2<-factor(sample(letters[1:3], 5, TRUE), levels=letters[1:3])
cbind(fac1, fac2)
sir(~fac1, ~fac2)
```

---

sm2asreml

*Converts sparseMatrix to asreml's giv format*


---

**Description**

Converts sparseMatrix to asreml's giv format: row-ordered, upper triangle sparse matrix.

**Usage**

```
sm2asreml(A=NULL, rownames=NULL)
```

**Arguments**

A	sparseMatrix
rownames	rownames of A

**Value**

data.frame: if A was formed from a pedigree equivalent to giv format returned by asreml. Ainverse

**Author(s)**

Jarrold Hadfield <j.hadfield@ed.ac.uk>

**See Also**

inverseA

**Examples**

```
data(bird.families)
A<-inverseA(bird.families)
Aasreml<-sm2asreml(A$Ainv, A$node.names)
```

---

spl	<i>Orthogonal Spline Design Matrix</i>
-----	--

---

**Description**

Orthogonal Spline Design Matrix

**Usage**

```
spl(x, k=10, knots=NULL, type="LRTP")
```

**Arguments**

x	a numeric covariate
k	integer, defines knot points at the 1:k/(k+1) quantiles of x
knots	vector of knot points
type	type of spline - currently only low-rank thin-plate ("LRTP") are implemented

**Value**

Design matrix post-multiplied by the inverse square root of the penalty matrix

**Author(s)**

Jarrod Hadfield <j.hadfield@ed.ac.uk>

**Examples**

```
## Not run:
x<-rnorm(100)
y<-x^2+cos(x)-x+0.2*x^3+rnorm(100)
plot(y~x)
lines((x^2+cos(x)-x+0.2*x^3)[order(x)]~sort(x))

dat<-data.frame(y=y, x=x)

m1<-MCMCg1mm(y~x, random=~idv(spl(x)), data=dat, pr=TRUE, verbose=FALSE) # penalised smoother
m2<-MCMCg1mm(y~x+spl(x), data=dat, verbose=FALSE) # non-penalised

pred1<-(cbind(m1$X, m1$Z)%*%colMeans(m1$Sol))@x
pred2<-(cbind(m2$X)%*%colMeans(m2$Sol))@x

lines(pred1[order(x)]~sort(x), col="red")
```

```

lines(pred2[order(x)]~sort(x), col="green")

m1$DIC-mean(m1$Deviance) # effective number of parameters < 13
m2$DIC-mean(m2$Deviance) # effective number of parameters ~ 13

## End(Not run)

```

---

SShorns

*Horn type and genders of Soay Sheep*


---

### Description

Horn type and genders of Soay Sheep *Ovis aires*

### Usage

```
data(SShorns)
```

### Format

a data frame with 666 rows and 3 columns, with individual identifier (id), horn type (horn) and gender (sex).

### References

Clutton-Brock T., Pemberton, J. Eds. 2004 Soay Sheep: Dynamics and Selection in an Island Population

---

summary.MCMCglmm

*Summarising GLMM Fits from MCMCglmm*


---

### Description

summary method for class "MCMCglmm". The returned object is suitable for printing with the print.summary.MCMCglmm method.

### Usage

```
## S3 method for class 'MCMCglmm'
summary(object, random=FALSE, ...)
```

### Arguments

object	an object of class "MCMCglmm"
random	logical: should the random effects be summarised
...	Further arguments to be passed

**Value**

DIC	Deviance Information Criterion
fixed.formula	model formula for the fixed terms
random.formula	model formula for the random terms
residual.formula	model formula for the residual terms
solutions	posterior mean, 95% HPD interval, MCMC p-values and effective sample size of fixed (and random) effects
Gcovariances	posterior mean, 95% HPD interval and effective sample size of random effect (co)variance components
Gterms	indexes random effect (co)variances by the component terms defined in the random formula
Rcovariances	posterior mean, 95% HPD interval and effective sample size of residual (co)variance components
Rterms	indexes residuals (co)variances by the component terms defined in the rcov formula
csats	chain length, burn-in and thinning interval
cutpoints	posterior mean, 95% HPD interval and effective sample size of cut-points from an ordinal model
theta_scale	posterior mean, 95% HPD interval, MCMC p-values and effective sample size of scaling parameter in theta_scale models.

**Author(s)**

Jarrold Hadfield <j.hadfield@ed.ac.uk>

**See Also**

[MCMCglmm](#)

---

Tri2M

*Lower/Upper Triangle Elements of a Matrix*

---

**Description**

Lower/Upper triangle elements of a matrix or forms a matrix from a vector of lower/upper triangle elements

**Usage**

Tri2M(x, lower.tri = TRUE, reverse = TRUE, diag = TRUE)

**Arguments**

x	Matrix or vector
lower.tri	If x is a matrix then the lower triangle (TRUE) or upper triangle FALSE elements (including diagonal elements) are returned. If x is a vector a matrix is formed under the assumption that x are the lower triangle (TRUE) or upper triangle (FALSE) elements.
reverse	logical: if TRUE a symmetric matrix is formed, if FALSE the remaining triangle is left as zeros.
diag	logical: if TRUE diagonal elements are included.

**Value**

numeric or matrix

**Author(s)**

Jarrod Hadfield <j.hadfield@ed.ac.uk>

**Examples**

```
M<-rIW(diag(3), 10)
x<-Tri2M(M)
x
Tri2M(x, reverse=TRUE)
Tri2M(x, reverse=FALSE)
```

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