

# Package ‘ordinalgmifs’

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**Title** Ordinal Regression for High-Dimensional Data

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**Depends** R (>= 2.10), survival

## Description

Provides a function for fitting cumulative link, adjacent category, forward and backward continuation ratio, and stereotype ordinal response models when the number of parameters exceeds the sample size, using the the generalized monotone incremental forward stagewise method.

**License** GPL (>= 2)

**Imports** methods

**BuildResaveData** best

**SystemRequirements** C++11

**NeedsCompilation** yes

**BuildVignettes** TRUE

**LazyData** true

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ordinalgmifs-package *Ordinal Response Regression for High-Dimensional Data*

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## Description

This package provides a function, `ordinalgmifs`, for fitting cumulative link, adjacent category, forward and backward continuation ratio, and stereotype ordinal response models when the number of parameters exceeds the sample size, using the the generalized monotone incremental forward stagewise method.

## Details

Package: ordinalgmifs  
 Version: 1.0.7  
 Date: 2022-04-06  
 Title: Ordinal Regression for High-Dimensional Data  
 Author: Kellie J. Archer, Jiayi Hou, Qing Zhou, Kyle Ferber, John G. Layne, Amanda Gentry  
 Maintainer: Kellie J. Archer <archer.43@osu.edu>  
 Depends: R (>= 2.10), survival  
 Description: Provides a function for fitting cumulative link, adjacent category, forward and backward continuation  
 License: GPL (>= 2)  
 Imports: methods  
 BuildResaveData: best  
 SystemRequirements: C++11  
 NeedsCompilation: yes  
 BuildVignettes: TRUE  
 LazyData: true

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<code>ordinalgmifs-package</code>	Ordinal Response Regression for High-Dimensional Data
<code>plot.ordinalgmifs</code>	Plot Solution Path for Ordinal GMIFS Fitted Model.
<code>predict.ordinalgmifs</code>	Predicted Probabilities and Class for Ordinal GMIFS Fit.
<code>print.ordinalgmifs</code>	Print the Contents of an Ordinal GMIFS Fitted Object.
<code>summary.ordinalgmifs</code>	Summarize an Ordinal GMIFS Object.

This package contains generic methods (coef, plot, predict, print, summary) that can be invoked for an object fitted using ordinalgmifs.

**Author(s)**

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**References**

Hastie T., Taylor J., Tibshirani R., and Walther G. (2007) Forward stagewise regression and the monotone lasso. *Electronic Journal of Statistics*, 1, 1-29.

**See Also**

See Also [ordinalgmifs](#). For models where no predictor is penalized see [vglm](#)

---

coef.ordinalgmifs      *Extract Model Coefficients*

---

**Description**

coef.ordinalgmifs is a generic function which extracts the model coefficients from a fitted model object fit using ordinalgmifs

**Usage**

```
## S3 method for class 'ordinalgmifs'  
coef(object, model.select = "AIC", ...)
```

**Arguments**

object	an ordinalgmifs object.
model.select	when x is specified any model along the solution path can be selected. The default is model.select="AIC" which extracts the coefficients from the model having the lowest AIC. Other options are model.select="BIC" or any numeric value from the solution path.
...	other arguments.

**Value**

Coefficients extracted from the model object.

**Author(s)**

Kellie J. Archer

**References**

Hastie T., Taylor J., Tibshirani R., and Walther G. (2007) Forward stagewise regression and the monotone lasso. *Electronic Journal of Statistics*, 1, 1-29.

**See Also**

See Also [ordinalgmifs](#), [summary.ordinalgmifs](#), [plot.ordinalgmifs](#), [predict.ordinalgmifs](#)

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eyedisease

*Eye Disease Risk Factors*

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**Description**

Eye Disease Risk Factors data from Section 9.1 of Agresti's Analysis of Ordinal Categorical Data. The primary data are from the Wisconsin Epidemiological Study of Diabetic Retinopathy. The primary outcome is severity of retinopathy which was measured in the left and right eye of every subject.

**Usage**

```
data(eyedisease)
```

**Format**

A data frame with 720 observations on the following 19 variables.

rme right eye macular oedema (absent = 0, present = 1)

lme left eye macular oedema (absent = 0, present = 1)

rre right eye refraction index

lre left eye refraction index

riop right eye intraocular eye pressure

liop left eye intraocular eye pressure

age age

diab duration of diabetes (in years)

gh glycosylated haemoglobin level

sbp systolic blood pressure

dbp diastolic blood pressure

bmi body mass index

pr pulse rate?

sex gender (male=1, female=2)

prot proteinuria (absent = 0, present = 1)

dose a numeric vector

rer1 right eye severity of retinopathy, an ordered factor with levels None < Mild < Moderate < Proliferative

ler1 left eye severity of retinopathy, an ordered factor with levels None < Mild < Moderate < Proliferative

id subject identifier

## References

R. Klein and B.E.K. Klein and S.E. Moss and M.D. Davis and D.L. DeMets. (1984) The Wisconsin Epidemiologic Study of Diabetic Retinopathy II. Prevalence and risk of diabetic retinopathy when age at diagnosis is less than 30 years. *Archives of Ophthalmology* 101, 520-526.

J. Williamson and K. Kim. (1996) A global odds ratio regression model for bivariate ordered categorical data from ophthalmologic studies. *Statistics in Medicine* 15: 1507-1518.

A. Agresti. (2010) *Analysis of Ordered Categorical Data*, Second Edition. Wiley. Hoboken, NJ.

## See Also

See Also as [ordinalgmifs](#)

## Examples

```
data(eyedisease)
```

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hccframe	<i>Liver Cancer Methylation Data</i>
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## Description

These data are a subset of subjects and CpG sites reported in the original paper where liver samples were assayed using the Illumina GoldenGate Methylation BeadArray Cancer Panel I. Technical replicate samples were removed to ensure all samples were independent. The matched cirrhotic samples from subjects with hepatocellular carcinoma (HCC, labeled Tumor) were also excluded. Therefore methylation levels in liver tissue are provided for independent subjects whose liver was Normal (N=20), cirrhotic but not having HCC (N=16, Cirrhosis non-HCC), and HCC (N=20, Tumor).

## Usage

```
data(hccframe)
```

## Format

A data frame with 56 observations on the following 46 variables.

group an ordered factor with levels Normal < Cirrhosis non-HCC < Tumor

CDKN2B\_seq\_50\_S294\_F a numeric vector representing a CpG site proportion methylation for CDKN2B

DDIT3\_P1313\_R a numeric vector representing a CpG site proportion methylation for DDIT3  
ERN1\_P809\_R a numeric vector representing a CpG site proportion methylation for ERN1  
GML\_E144\_F a numeric vector representing a CpG site proportion methylation for GML  
HDAC9\_P137\_R a numeric vector representing a CpG site proportion methylation for HDAC9  
HLA.DPA1\_P205\_R a numeric vector representing a CpG site proportion methylation for HLA.DPA1  
HOXB2\_P488\_R a numeric vector representing a CpG site proportion methylation for HOXB2  
IL16\_P226\_F a numeric vector representing a CpG site proportion methylation for IL16  
IL16\_P93\_R a numeric vector representing a CpG site proportion methylation for IL16  
IL8\_P83\_F a numeric vector representing a CpG site proportion methylation for IL8  
MPO\_E302\_R a numeric vector representing a CpG site proportion methylation for MPO  
MPO\_P883\_R a numeric vector representing a CpG site proportion methylation for MPO  
PADI4\_P1158\_R a numeric vector representing a CpG site proportion methylation for PADI4  
SOX17\_P287\_R a numeric vector representing a CpG site proportion methylation for SOX17  
TJP2\_P518\_F a numeric vector representing a CpG site proportion methylation for TJP2  
WRN\_E57\_F a numeric vector representing a CpG site proportion methylation for WRN  
CRIP1\_P874\_R a numeric vector representing a CpG site proportion methylation for CRIP1  
SLC22A3\_P634\_F a numeric vector representing a CpG site proportion methylation for SLC22A3  
CCNA1\_P216\_F a numeric vector representing a CpG site proportion methylation for CCNA1  
SEPT9\_P374\_F a numeric vector representing a CpG site proportion methylation for SEPT9  
ITGA2\_E120\_F a numeric vector representing a CpG site proportion methylation for ITGA2  
ITGA6\_P718\_R a numeric vector representing a CpG site proportion methylation for ITGA6  
HGF\_P1293\_R a numeric vector representing a CpG site proportion methylation for HGF  
DLG3\_E340\_F a numeric vector representing a CpG site proportion methylation for DLG3  
APP\_E8\_F a numeric vector representing a CpG site proportion methylation for APP  
SFTPB\_P689\_R a numeric vector representing a CpG site proportion methylation for SFTPB  
PENK\_P447\_R a numeric vector representing a CpG site proportion methylation for PENK  
COMT\_E401\_F a numeric vector representing a CpG site proportion methylation for COMT  
NOTCH1\_E452\_R a numeric vector representing a CpG site proportion methylation for NOTCH1  
EPHA8\_P456\_R a numeric vector representing a CpG site proportion methylation for EPHA8  
WT1\_P853\_F a numeric vector representing a CpG site proportion methylation for WT1  
KLK10\_P268\_R a numeric vector representing a CpG site proportion methylation for KLK10  
PCDH1\_P264\_F a numeric vector representing a CpG site proportion methylation for PCDH1  
TDGF1\_P428\_R a numeric vector representing a CpG site proportion methylation for TDGF1  
EFNB3\_P442\_R a numeric vector representing a CpG site proportion methylation for EFNB3  
MMP19\_P306\_F a numeric vector representing a CpG site proportion methylation for MMP19  
FGFR2\_P460\_R a numeric vector representing a CpG site proportion methylation for FGFR2  
RAF1\_P330\_F a numeric vector representing a CpG site proportion methylation for RAF1

BMP2\_E435\_F a numeric vector representing a CpG site proportion methylation for BMP2  
GRB10\_P496\_R a numeric vector representing a CpG site proportion methylation for GRB10  
CTSH\_P238\_F a numeric vector representing a CpG site proportion methylation for CTSH  
SLC6A8\_seq\_28\_S227\_F a numeric vector representing a CpG site proportion methylation for SLC6A8  
PLXDC1\_P236\_F a numeric vector representing a CpG site proportion methylation for PLXDC1  
TFE3\_P421\_F a numeric vector representing a CpG site proportion methylation for TFE3  
TSG101\_P139\_R a numeric vector representing a CpG site proportion methylation for TSG101

### Source

The full dataset is available as GSE18081 from Gene Expression Omnibus at <https://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE18081>

### References

Archer KJ, Mas VR, Maluf DG, Fisher RA. High-throughput assessment of CpG site methylation for distinguishing between HCV-cirrhosis and HCV-associated hepatocellular carcinoma. *Molecular Genetics and Genomics*, 283(4): 341-349, 2010.

### See Also

See Also as [ordinalgmifs](#)

### Examples

```
data(hccframe)
```

---

ordinalgmifs	<i>Ordinal Generalized Monotone Incremental Forward Stagewise Regression</i>
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### Description

This function can fit a cumulative link, adjacent category, forward and backward continuation ratio, and stereotype ordinal response model when the number of parameters exceeds the sample size, using the the generalized monotone incremental forward stagewise method.

### Usage

```
ordinalgmifs(formula, data, x = NULL, subset, epsilon = 0.001, tol = 1e-05,  
scale = TRUE, probability.model = "Cumulative", link = "logit",  
verbose=FALSE, assumption=NULL, ...)
```

**Arguments**

<code>formula</code>	an object of class "formula" (or one that can be coerced to that class): a symbolic description of the model to be fitted. The left side of the formula is the ordinal outcome while the variables on the right side of the formula are the covariates that are not included in the penalization process. Note that if all variables in the model are to be penalized, an intercept only model formula should be specified.
<code>data</code>	an optional data frame, list or environment (or object coercible by <code>as.data.frame</code> to a data frame) containing the variables in the model.
<code>x</code>	an optional matrix of predictors that are to be penalized in the model fitting process.
<code>subset</code>	an optional vector specifying a subset of observations to be used in the fitting process.
<code>epsilon</code>	small incremental amount used to update a coefficient at a given step.
<code>tol</code>	the iterative process stops when the difference between successive log-likelihoods is less than this specified level of tolerance.
<code>scale</code>	logical, if TRUE the penalized predictors are centered and scaled.
<code>probability.model</code>	the type of ordinal response model to be fit. Can be "Cumulative", "AdjCategory", "ForwardCR", "BackwardCR", or "Stereotype"
<code>link</code>	the link function used. Allowable links for "Cumulative", "ForwardCR", and "BackwardCR" are "logit", "probit", and "cloglog". For an "AdjCategory" model only a "loge" link is allowed; for a "Stereotype" model only a "logit" link is allowed.
<code>verbose</code>	logical, if TRUE the step number is printed to the console (default is FALSE).
<code>assumption</code>	integer, only use with <code>probability.model = "ForwardCR"</code> and <code>link = "cloglog"</code> to denote the assumption to use for discrete censored survival modeling. If <code>assumption = 1</code> , assume the observation was censored at the end of the discrete time interval in which the censoring occurred; if <code>assumption = 2</code> , assume the observation was censored at the beginning of the interval in which censoring occurred; if <code>assumption = 3</code> , assume constant hazard rate within the interval in which the censoring occurred; if no censoring occurs, do not specify a value for <code>assumption</code> .
<code>...</code>	additional arguments

**Details**

A model specified as `response~terms,x=penalized.terms` where `response` is the ordinal response vector and `terms` is the series of variables in the model that are not to be penalized and `x` is a matrix of variables that are to be penalized. For example, `terms` may include the variables `age` and `gender` while `x` includes hundreds to thousands of features from a high-throughput genomic experiment. In the event that no baseline demographic/clinical characteristics/subject level variables are available or needed in `terms` (all variables are to be penalized) then the model is specified as `response~1,x=penalized.terms`.



**Value**

AIC	a vector of AIC values for each step (if x is specified).
BIC	a vector of BIC values for each step (if x is specified).
alpha	the ordinal threshold estimates for the fitted model.
theta	the coefficient estimates for the unpenalized variables (if terms are specified on the right hand side of the model formula).
beta	the coefficient estimates for the penalized variables (if x is specified in the model).
phi	the scaling coefficient estimates (if a "Stereotype" logit model is fit).
logLik	a vector of log-likelihood values for each step (if terms are specified on the right hand side of the model formula).
link	the link function used in the model fit.
model.select	the step at which the minimum AIC was observed (if terms are specified on the right hand side of the model formula).
probability.model	the model fit.
scale	logical indicating whether penalized variables were centered and scaled.
w	the unpenalized variables in the model (if any).
x	the penalized variables in the model (if any).
y	the ordinal response.

**Author(s)**

Kellie J. Archer, Jiayi Hou, Qing Zhou, Kyle Ferber, John G. Layne, Amanda Gentry

**References**

Hastie T., Taylor J., Tibshirani R., and Walther G. (2007) Forward stagewise regression and the monotone lasso. *Electronic Journal of Statistics*, 1, 1-29.

**See Also**

See Also [coef.ordinalgmifs](#), [summary.ordinalgmifs](#), [plot.ordinalgmifs](#), [predict.ordinalgmifs](#)

**Examples**

```
data(hccframe)
# To minimize processing time, MPO_E302_R is coerced into the model and only a subset of
# two CpG sites (DDIT3_P1313_R and HDAC9_P137_R) are included as penalized covariates
# in this demonstration, and epsilon is set to 0.01
hcc.fit <- ordinalgmifs(group ~ MPO_E302_R, x = c("DDIT3_P1313_R", "HDAC9_P137_R"),
data = hccframe, epsilon = 0.01)
coef(hcc.fit)
summary(hcc.fit)
phat <- predict(hcc.fit)
head(phat$predicted)
table(phat$class, hccframe$group)
```

---

plot.ordinalgmifs      *Plot Solution Path for Ordinal GMIFS Fitted Model.*

---

### Description

This function plots either the coefficient path, the AIC, or the log-likelihood for a fitted ordinalgmifs object.

### Usage

```
## S3 method for class 'ordinalgmifs'
plot(x, type = "trace", xlab=NULL, ylab=NULL, main=NULL, ...)
```

### Arguments

x	an ordinalgmifs object.
type	default is "trace" which plots the coefficient path for the fitted object. Also available are "AIC", "BIC", and "logLik".
xlab	a default x-axis label will be used which can be changed by specifying a user-defined x-axis label.
ylab	a default y-axis label will be used which can be changed by specifying a user-defined y-axis label.
main	a default main title will be used which can be changed by specifying a user-defined main title.
...	other arguments.

### Author(s)

Kellie J. Archer

### See Also

See Also [ordinalgmifs](#), [coef.ordinalgmifs](#), [summary.ordinalgmifs](#), [predict.ordinalgmifs](#)

---

predict.ordinalgmifs      *Predicted Probabilities and Class for Ordinal GMIFS Fit.*

---

### Description

This function returns a list that includes the predicted probabilities as well as the predicted class for an ordinalgmifs fitted object.

**Usage**

```
## S3 method for class 'ordinalgmifs'
predict(object, neww = NULL, newdata, newx = NULL, model.select = "AIC", ...)
```

**Arguments**

object	an ordinalgmifs fitted object.
neww	an optional formula that includes the unpenalized variables to use for predicting the response. If omitted, the training data are used.
newdata	an optional data.frame that minimally includes the unpenalized variables to use for predicting the response. If omitted, the training data are used.
newx	an optional matrix of penalized variables to use for predicting the response. If omitted, the training data are used.
model.select	when x is specified any model along the solution path can be selected. The default is model.select="AIC" which calculates the predicted values using the coefficients from the model having the lowest AIC. Other options are model.select="BIC" or any numeric value from the solution path.
...	other arguments.

**Value**

predicted	a matrix of predicted probabilities from the fitted model.
class	a vector containing the predicted class taken as that class having the largest predicted probability.
...	other arguments.

**Author(s)**

Kellie J. Archer, Jiayi Hou, Qing Zhou, Kyle Ferber, John G. Layne, Amanda Gentry

**See Also**

See Also [ordinalgmifs](#), [coef.ordinalgmifs](#), [summary.ordinalgmifs](#), [plot.ordinalgmifs](#)

---

print.ordinalgmifs      *Print the Contents of an Ordinal GMIFS Fitted Object.*

---

**Description**

This function prints the names of the list objects from an ordinalgmifs fitted model.

**Usage**

```
## S3 method for class 'ordinalgmifs'
print(x, ...)
```

**Arguments**

x                    an ordinalgmifs object.  
 ...                other arguments.

**Note**

The contents of an ordinalgmifs fitted object differ depending upon whether x is specified in the ordinalgmifs model (i.e., penalized variables are included in the model fit hence a solution path is returned) or only terms on the right hand side of the equation are included (unpenalized variables). In the latter case, we recommend using the VGAM package.

**Author(s)**

Kellie J. Archer

**See Also**

See Also [ordinalgmifs](#), [coef.ordinalgmifs](#), [summary.ordinalgmifs](#), [plot.ordinalgmifs](#), [predict.ordinalgmifs](#)

---

summary.ordinalgmifs    *Summarize an Ordinal GMIFS Object.*

---

**Description**

summary method for class ordinalgmifs.

**Usage**

```
## S3 method for class 'ordinalgmifs'
summary(object, model.select = "AIC", ...)
```

**Arguments**

object            an ordinalgmifs object.  
 model.select    when x is specified any model along the solution path can be selected. The default is model.select="AIC" which extracts the model having the lowest AIC. Other options are model.select="BIC" or any numeric value from the solution path.  
 ...                other arguments.

**Details**

Prints the following items extracted from the fitted ordinalgmifs object: the probability model and link used and model parameter estimates. For models that include x, the parameter estimates, AIC, BIC, and log-likelihood are printed for indicated model.select step or if model.select is not supplied the step at which the minimum AIC was observed.

**Author(s)**

Kellie J. Archer

**See Also**

See Also [ordinalgmifs](#), [coef.ordinalgmifs](#), [plot.ordinalgmifs](#), [predict.ordinalgmifs](#)

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