Package: endogenous (via r-universe)

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Type Package
Title Classical Simultaneous Equation Models
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Description Likelihood-based approaches to estimate linear regression parameters and treatment effects in the presence of endogeneity. Specifically, this package includes James Heckman's classical simultaneous equation models-the sample selection model for outcome selection bias and hybrid model with structural shift for endogenous treatment. For more information, see the seminal paper of Heckman (1978) <doi:10.3386 w0177=""> in which the details of these models are provided. This package accommodates repeated measures on subjects with a working independence approach. The hybrid model further accommodates treatment effect modification.</doi:10.3386>
Imports mytnorm
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NeedsCompilation no
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hybrid	Hybrid model with structural shift (permits covariate-specific treatment effects)

Description

James Heckman's Hybrid Model with Structural Shift (also known as the Treatment Effects Model). Jointly models outcome regression model and endogenous variable probit model (e.g., outcome associations in the presence of endogenous treatment in observational data). Can handle clustered data. Accommodates treatment effect modification based on observable covariates.

Usage

```
## S3 method for class "hybrid"
hybrid(outcome, probit, modifiers = NULL, init = NULL, id = NULL, se = "R")
```

Arguments

outcome	an object of class "formula" with a numeric vector on the left hand side, and
	predictors of interest on the right hand side.
probit	an object of class "formula" with a binary (0/1) numeric vector on the left hand

side (1 indicating medication use), and predictors of medication use on the right hand side (right hand side permitted to contain variables on the right hand side

of the outcome equation).

an object of class "formula" with a binary numeric vector indicating medication use on the left hand side, and treatment effect modifiers on the right hand side. If effect modifiers are treatment group specific (e.g., medication dose), set the effect modifier variables to zero for the untreated observations. If any other numeric values are used, they will ultimately be zet to zero. If NULL, the average treatment effect will be estimated under the assumption of no effect modifica-

a vector of initial values. The ordering of subparameters is: alpha (probit model parameters), beta (outcome model parameters), eta (an intercept, with or without effect mofidier paramters), sigmay (outcome error standard deviation), rho (error correlation). If NULL, an initial value will be chosen through OLS linear

regression and probit-link GLM without regard to endogeneity.

a numeric vector indicating subject IDs if data are clustered. In the absence of

clustered data, this can be left blank (defaults to NULL).

a string, either "M" for model-based standard errors (based on inverse observed Fisher information), or "R" for robust standard errors (based on methods of Huber and White). Defaults to "R". If id is provided for clustered data, the clusterrobust variance estimator (with working independence) will be used even if the

user specifies type "M".

modifiers

init

id

se

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Details

The model is evaluated with numerical minimization of the negative log-likelihood (the BFGS is used). The probit model and error correlation parameters are weakly identified and hence the error variance is set at unity. The data must be complete (no missing values) and numeric, with the exception of factors, which may be used on the right hand side of equations.

Value

hybrid prints a summary of the coefficient estimates, standard errors, Wald-based confidence intervals, and p-values for the outcome model, the treatment effects (and potentially effect modifiers), and the medication use probit model.

alpha estimate of the medication use probit model parameters.

beta estimate of the outcome model parameters.

eta estimate of the treatment effect, with or without effect modifier parameters.

sigma estimate of the standard deviation of the outcome error.

rho estimate of the correlation between the errors.

vcov entire estimated variance-covariance matrix, provided if the user wishes to per-

form any more specific hypothesis tests.

init initial value ultimately used, whether specified by the user or generated through

the default approach.

fitted vector of fitted outcome values.

call the matched call.

out.form the formula used for the outcome model.

prob. form the formula used for the medication use probit model.

tx. form the formula used for the treatment effects model (potentially with effect modi-

fiers).

sterr the choice of the variance estimate procedure (either model-based or robust).

labels labels for predictors to be passed into output.

Author(s)

Andrew J. Spieker, Ph.D.

References

Heckman JJ. Dummy endogenous variables in a simultaneous equation system. *Econometrica* 46(4), 931-959.

Maddala GS. *Limited-dependent and qualitative variables in econometrics*. Cambridgeshire: Cambridge University Press; 1983.

Spieker AJ, Delaney JAC, and McClelland RL. Evaluating the treatment effects model for estimation of cross-sectional associations between risk factors and cardiovascular biomarkers influenced by medication use. *Pharmacoepidemiology and Drug Safety* 24(12), 1286-1296.

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Examples

```
#- Generate Data -#
require(mvtnorm)
set.seed(1)
N <- 2000
X1 <- rnorm(N, 0, 1);</pre>
X2 <- rnorm(N, 0, 1);
X3 \leftarrow rnorm(N, 0, 1);
errors <- rmvnorm(N, sigma = 50*matrix(c(1, 0.5, 0.5, 1), nrow = 2))
Y < -50 + X1 + X2 + errors[,1]
Z \leftarrow rep(0, N)
Z[(-5 + X1 + X3 + errors[,2]) > 0] < -1
Y[Z == 1] \leftarrow Y[Z == 1] - 0.5*X1[Z == 1]
#- Estimate Model with No Effect Modification -#
hybrid(Y \sim X1 + X2, probit = Z \sim X1 + X3)
#- Estimate Model with Effect Modification -#
hybrid(Y \sim X1 + X2, probit = Z \sim X1 + X3, modifiers = Z \sim X1)
#- Estimate Model with Effect Modification and Model-Based Variance -#
hybrid(Y \sim X1 + X2, probit = Z \sim X1 + X3, modifiers = Z \sim X1, se = "M")
```

sampselect

Sample selection model (endogenous probit).

Description

James Heckman's Classical Simultaneous Equation Model (also known as the Sample Selection Model). Used to account for endogenous sample selection. Jointly models outcome model with propensity of selection, in which some of the outcomes are unobserved. Can also handle clustered data.

Usage

```
## S3 method for class "sampselect"
sampselect(outcome, probit, init = NULL, id = NULL, se = "R")
```

Arguments

outcome

an object of class "formula" with a numeric vector on the left hand side, and predictors of interest on the right hand side. Values on the left hand side that correspond to unobserved outcomes should be set to numeric values (to zero, for example, although they can be set to any numeric values).

probit

an object of class "formula" with a binary (0/1) numeric vector on the left hand side (1 indicating unobserved outcome), and predictors of selection on the right hand side (right hand side permitted to contain variables on the right hand side of the outcome equation).

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init	a vector of initial values. The ordering of subparameters is: alpha (probit model parameters), beta (outcome model parameters), sigmay (outcome error standard deviation), rho (error correlation). If NULL, an initial value will be chosen through OLS linear regression and probit-link GLM without regard to endogeneity.
id	a numeric vector indicating subject IDs if data are clustered. In the absence of clustered data, this can be left blank (defaults to NULL).
se	a string, either "M" for model-based standard errors (based on inverse observed Fisher information), or "R" for robust standard errors (based on methods of Huber and White). Defaults to "R". If id is provided for clustered data, the cluster-robust variance estimator (with working independence) will be used even if the user specifies type "M".

Details

The model is evaluated with numerical minimization of the negative log-likelihood (the BFGS is used). The probit model and error correlation parameters are weakly identified and hence the error variance is set at unity. The data must be complete (no missing values) and numeric, with the exception of factors, which may be used on the right hand side of equations.

Value

sampselect prints a summary of the coefficient estimates, standard errors, Wald-based confidence intervals, and p-values for the outcome model and the selection use probit model.

alpha	estimate of the selection probit model parameters.
beta	estimate of the outcome model parameters.
sigma	estimate of the standard deviation of the outcome error.
rho	estimate of the correlation between the errors.
vcov	entire estimated variance-covariance matrix, provided if the user wishes to perform any more specific hypothesis tests.
init	initial value ultimately used, whether specified by the user or generated through the default approach. $$
fitted	vector of fitted outcome values.
call	the matched call.
out.form	the formula used for the outcome model.
prob.form	the formula used for the selection probit model.
sterr	the choice of the variance estimate procedure (either model-based or robust).
labels	labels for predictors to be passed into output.

Author(s)

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References

Heckman JJ. Dummy endogenous variables in a simultaneous equation system. *Econometrica* 46(4), 931-959.

Maddala GS. *Limited-dependent and qualitative variables in econometrics*. Cambridgeshire: Cambridge University Press; 1983.

Examples

```
#- Generate Data -#
require(mvtnorm)
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N <- 2000
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X2 <- rnorm(N, 0, 1);
X3 <- rnorm(N, 0, 1);
errors <- rmvnorm(N, sigma = 50*matrix(c(1, 0.5, 0.5, 1), nrow = 2))
Y <- 50 + X1 + X2 + errors[,1]
Z \leftarrow rep(0, N)
Z[(-5 + X1 + X3 + errors[,2]) > 0] < -1
Y[Z == 1] <- 0
#- Estimate Model -#
sampselect(Y \sim X1 + X2, probit = Z \sim X1 + X3)
#- Estimate Model with Model-Based Variance -#
sampselect(Y \sim X1 + X2, probit = Z \sim X1 + X3, se = "M")
```

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