

# Package ‘bacr’

October 23, 2016

**Type** Package

**Title** Bayesian Adjustment for Confounding

**Version** 1.0.1

**Depends** R(>= 2.13.0), graphics, stats, MCMCpack

**Date** 2016-10-23

**Author** Chi Wang <chi.wang@uky.edu>

**Maintainer** Chi Wang <chi.wang@uky.edu>

**Description** Estimating the average causal effect based on the Bayesian Adjustment for Confounding (BAC) algorithm.

**License** GPL-2

**NeedsCompilation** no

**Repository** CRAN

**Date/Publication** 2016-10-23 22:58:52

## R topics documented:

bacr-package . . . . .	1
bac . . . . .	3
plot.bacr . . . . .	5
summary.bacr . . . . .	5

<b>Index</b>	<b>7</b>
--------------	----------

---

bacr-package	<i>Bayesian Adjustment for Confounding</i>
--------------	--

---

## Description

This package implements the Bayesian Adjustment for Confounding (BAC) algorithm for estimating the Average Causal Effect (ACE) in Generalized Linear Models. It accounts for uncertainty in confounder and effect modifier selection and allows estimation of the ACE for the whole population or for a specific subpopulation.

## Details

```

Package:   bacr
Type:     Package
Version:  1.0.1
Depends:  R(>= 2.13.0), graphics, stats, MCMCpack
Date:     2016-10-23
License:  GPL-2

```

## Author(s)

Chi Wang

Maintainer: Chi Wang <chi.wang@uky.edu>

## References

Wang C, Dominici F, Parmigiani G, Zigler CW. Accounting for Uncertainty in Confounder and Effect Modifier Selection When Estimating Average Causal Effects in Generalized Linear Models. *Biometrics*, 71(3): 654-665, 2015.

## Examples

```

##### Note that the example below is for illustration purpose only. ###
##### In practice, larger number of iterations will be needed.#####
##### simulate data #####
n = 200; m = 4
V = matrix(rnorm(n*m),ncol=m)
X = rbinom(n, size=1, prob=exp(V[,1])/(1+exp(V[,1])))
beta = c(1,1,1,0.5)
temp0 = cbind(rep(0,n), V[,1:3])
temp1 = cbind(rep(1,n), V[,1:3])
Y0 = rbinom(n, size=1, prob=exp(temp0)/(1+exp(temp0)))
Y1 = rbinom(n, size=1, prob=exp(temp1)/(1+exp(temp1)))
Y = Y0
Y[X==1] = Y1[X==1]
Z = as.data.frame(cbind(Y, X, V))
names(Z) = c("Y", "X", paste("V", 1:m, sep=""))
##### run BAC #####
result = bac(data=Z, exposure="X", outcome="Y", confounders=paste("V", 1:m, sep=""),
             interactors=NULL, familyX="binomial", familyY="binomial", omega=Inf,
             num_its=5, burnM=1, burnB=1, thin=1)
##### summarize results #####
summary(result)
plot(result)

### Adding interaction terms #####
beta = c(1,1,1,1,1)
temp0 = cbind(rep(0,n), V[,1:3], rep(0,n)*V[,3])
temp1 = cbind(rep(1,n), V[,1:3], rep(1,n)*V[,3])

```

```

Y0 = rbinom(n, size=1, prob=exp(temp0)/(1+exp(temp0)))
Y1 = rbinom(n, size=1, prob=exp(temp1)/(1+exp(temp1)))
Y = Y0
Y[X==1] = Y1[X==1]
Z = as.data.frame(cbind(Y, X, V))
names(Z) = c("Y", "X", paste("V", 1:m, sep=""))
result = bac(data=Z, exposure="X", outcome="Y", confounders=paste("V", 1:m, sep=""),
             interactors=paste("V", 1:m, sep=""), familyX="binomial", familyY="binomial",
             omega=Inf, num_its=5, burnM=1, burnB=1, thin=1)
summary(result)
plot(result)

##### Estimate the ACE in the exposed subgroup #####
result = bac(data=Z, exposure="X", outcome="Y", confounders=paste("V", 1:m, sep=""),
             interactors=paste("V", 1:m, sep=""), familyX="binomial", familyY="binomial",
             omega=Inf, num_its=5, burnM=1, burnB=1, thin=1, population=(X==1))
summary(result)
plot(result)

```

---

bac

*Bayesian Adjustment for Confounding (BAC)*


---

## Description

Estimating the Average Causal Effect (ACE) based on the BAC algorithm

## Usage

```

bac(data, exposure, outcome, confounders, interactors,
    familyX, familyY, omega = Inf, num_its, burnM, burnB,
    thin, population = NULL)

```

## Arguments

data	a data from containing the input data.
exposure	the exposure variable
outcome	the outcome variable
confounders	a vector of potential confounder variable names
interactors	a vector of the names of potential confounders that may interact with the exposure
familyX	the family of the exposure model. Currently, it allows gaussian, binomial, and poisson.
familyY	the family of the outcome model. Currently, it allows gaussian, binomial, and poisson.
omega	a dependent parameter, which is the prior odds of including a predictor in the outcome model, given that the same predictor is in the exposure model. The default value is Inf, which forces predictors in the exposure model to be included in the outcome model.

num_its	number of MCMC iterations excluding the burn-in iterations.
burnM	number of burn-in iterations when sampling the exposure and outcome models.
burnB	number of burn-in iterations when sampling model coefficients based on a given outcome model.
thin	the thinning parameter when sampling model coefficients based on a given outcome model.
population	the population for which the ACE is based on. It can be either unspecified or a vector of TRUE and FALSE. If unspecified, the function will estimate the ACE for the whole population. If specified, the function will estimate the ACE for the subpopulation defined by the individuals indicated by TRUE.

### Details

The function may run slowly for data with large sample size or many potential confounders. The users are suggested to choose small number of iterations first, evaluate the computational speed, then increase the number of iterations. Note that this function assumes a non-informative prior for outcome model coefficients and does not handle informative priors.

### Value

a list variable, which contains

data	a data from containing the input data.
MX	a matrix of MCMC samples of exposure models.
MY	a matrix of MCMC samples of outcome models.
models	a list variable containing unique outcome models in the MCMC sample and the appearance frequencies.
exposure	the exposure variable.
outcome	the outcome variable.
confounders	a vector of potential confounder variable names.
interactors	a vector of the names of potential confounders that may interact with the exposure.
predictorsY	All the possible predictors of the outcome.
ACE	a vector of MCMC samples of the ACE.
para	a list of model parameters

### Author(s)

Chi Wang

### References

Wang C, Dominici F, Parmigiani G, Zigler CW. Accounting for Uncertainty in Confounder and Effect Modifier Selection When Estimating Average Causal Effects in Generalized Linear Models. *Biometrics*, 71(3): 654-665, 2015.

**See Also**

[plot.bacr](#), [summary.bacr](#)

---

plot.bacr	<i>Plot Method for Class 'bacr'</i>
-----------	-------------------------------------

---

**Description**

Plot the posterior inclusion probabilities of potential confounders for a bacr object

**Usage**

```
## S3 method for class 'bacr'  
plot(x, ...)
```

**Arguments**

x	An object returned by the bac function
...	arguments passed to or from other methods.

**Author(s)**

Chi Wang

**See Also**

[bac](#), [summary.bacr](#)

---

summary.bacr	<i>Summary Method for Class 'bacr'</i>
--------------	--

---

**Description**

Summarize results from running the bac function

**Usage**

```
## S3 method for class 'bacr'  
summary(object, ...)
```

**Arguments**

object	A bacr object
...	arguments passed to or from other methods.

**Value**

a list variable, which contains

posterior.mean Posterior mean of the ACE

CI 95% posterior interval of the ACE

PIP A vector of posterior inclusion probabilities for potential confounders

**Author(s)**

Chi Wang

**See Also**

[bac](#), [plot.bacr](#)

# Index

\*Topic **BAC**

    bacr-package, 1

    bac, 3, 5, 6

    bacr (bacr-package), 1

    bacr-package, 1

    plot.bacr, 5, 5, 6

    summary.bacr, 5, 5