# Package 'PropScrRand'

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Title Propensity Score Methods for Assigning Treatment in Randomized

Type Package

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<b>Description</b> Contains functions to run propensity-biased allocation to balance covariate distributions in sequential trials and propensity-constrained randomization to balance covariate distributions in trials with known baseline covariates at time of randomization. Currently only supports trials comparing two groups.
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PropScrRand-package

Propensity score methods for assigning treatment in randomized trials

### Description

Provides propensity score-based methods for allocating units to treatment experiments with two treatment levels (e.g., treatment and control).

#### **Details**

Package: PropScrRand
Type: Package
Version: 1.1
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License: GPL-3

For sequential allocation, the functions the user will interact with directly are pba and pbaAgain. Both of these functions perform propensity-biased allocation, producing a treatment assignment for the current unit, among other information. The function plotpi can be used to investigate the strength of balance forced by various values of the tuning parameter k, with curves for new values of k added to the plot via addpi. For randomization when all baseline covariates are known, use pcr, which will conduct propensity-constrained randomization. The remaining functions are called from these internally.

### Author(s)

Travis M. Loux

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genPerms

Generate Treatment Permutations

#### **Description**

Used within calls to pcr to generate a set of unique treatment permutations for randomization.

#### Usage

```
genPerms(n, n1, nPerms)
```

### Arguments

n Total number of units to be randomized.

n1 Number of units to receive treatment.

nPerms Number of permutations to generate.

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#### **Details**

This function randomly samples nPerms of the choose(n,n1) possible treatment permutations. If nPerms > choose(n,n1), then all choose(n,n1) permutations are generated systematically. Also, in the case of 1-to-1 allocation, the complement treatment vectors are also produced, so the returned matrix has 2\*nPerms permutations. Uniqueness is checked throughout and duplicate permutations disgarded.

#### Value

The result is an  $n1 \times nPerms$  (or  $n1 \times choose(n,n1)$ ) or  $n1 \times 2*nPerms$ ) matrix. Each column represents one treatment permutation, with the values in the column giving the index of the treated units.

#### Author(s)

Travis M. Loux

## **Examples**

```
genPerms(n=50, n1=25, nPerms=500)
genPerms(n=50, n1=35, nPerms=500)
```

getVar

Compute Propensity Score Variance

## Description

Given a data set and vector of indices for treated units, computes the variance of the propensity score fitted via logistic regression.

## Usage

```
getVar(covs, tIndex)
```

#### **Arguments**

covs A data frame of baseline covariates.

tIndex A vector indicating which units are to receive treatment.

#### Value

Returns the variance of the fitted propensity scores.

## Author(s)

Travis M. Loux

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#### **Description**

Performs propensity-biased allocation for assigning a new unit to treatment in a sequential design with two treatment levels (i.e., treatment and control).

## Usage

```
pba(x, tr, newx, k = 1, global = 0.5)
pbaAgain(previous, newx, k = NA)
```

#### **Arguments**

x A data frame of the covariate values of previously assigned units.

tr A vector of treatment assignments (0 or 1) for previously assigned units.

newx Data frame of covariate values of the new unit.

k Balancing parameter.

global Global target proportion to be treated.

previous The output of a previous call to pba or pbaAgain

#### **Details**

The function pba generates a treatment assignment for a new unit. The steps of the process include regressesing tr on x by logistic regression, computeing the fitted value of the new unit using covarate values in newx, and transforming the fitted propensity score into the probability of treatment by a call to piFunction using k and global as parameters. The balancing parameter k must be one of 0, Inf, or the ratio of two positive odd integers. Small values of k result in less restrictive randomization while larger values of k result in more forced balance. In particular, k = 0 is equivalent to pure randomization and k = 1nf results in deterministic allocation. Finally, a treatment assignment for the new unit is generated via a Bernoulli trial with probability from piFunction.

The function pbaAgain takes as input the output from a previous call to pba or pbaAgain and runs pba for the new unit using the values of newx. If k = NA (the default), the value of k from previous is used; otherwise, the provided value of k is used. The parameter global is assumed to stay the same throughout the trial. The output of pbaAgain contains the same information as pba.

#### Value

results	A list of results from the PBA procedure.
phat	The fitted propensity score for the new unit.
ptreat	The probability of assignment to the treatment group for the new unit.
newtr	Result of random assignment using ptreat.
input	A list of inputs to PBA procdure. Used in future calls to pbaAgain.

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x Input x.
tr Input tr.
newx Input newx.
k Input k.
global Input global.

## Author(s)

Travis Loux

#### References

Loux, T.M. (2013) A simple, flexible, and effective covariate-adaptive treatment allocation procedure. Statistics in Medicine 32(22), 3775-3787. DOI: 10.1002/sim.5837

## **Examples**

```
x0 = data.frame(matrix(rnorm(60), ncol=3))
t0 = rbinom(nrow(x0), size=1, prob=0.5)

x1 = data.frame(matrix(rnorm(3), ncol=3))
trial1 = pba(x=x0, tr=t0, newx=x1, k=Inf)

x2 = data.frame(matrix(rnorm(3), ncol=3))
trial2 = pbaAgain(previous=trial1, newx=x2)

x3 = data.frame(matrix(rnorm(3), ncol=3))
trial3 = pbaAgain(previous=trial2, newx=x3, k=5/3)
```

pcr

Propensity-Constrained Randomization

## **Description**

Performs propensity-contstrained randomization on a given data set with measured covariates on all units.

## Usage

```
pcr(x, nTreat, M, m)
```

## **Arguments**

X	Data frame of covariates.
nTreat	Number of units to be treated.
М	Number of candidate permutations to create.
m	Number of permutations to keep.

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#### **Details**

Given the parameters, pcr generates M unique permutations by calling genPerms. For each permutation, the empirical propensity scores are computed and the variance returned by getVar. Finally, the m permutations with the smallest propensity score variance are found. The m permutations returned in best can then be used to perform randomization and randomization inference. We suggest M >= 10000 and m/M <= 0.10.

#### Value

treatments The (approximately) M permutations generated by genPerms.

 $\label{eq:approx} \textbf{A vector of the propensity score variances for all M permutations in treatments}.$ 

cutoff The calculated m/M quantile of propensity score variances.

best The column indices of the permutations in treatments with propensity score

variance below cutoff.

## Author(s)

Travis Loux

#### References

Loux, T.M. (2015) Randomization, matching, and propensity scores in the design and analysis of experimental studies with known covariates. Statistics in Medicine. 34(4), 558-570. DOI: 10.1002/sim.6361

## **Examples**

```
# 1:1 allocation, M small
dat1 = data.frame(x1=rnorm(50),
                  x2=rnorm(50),
                  x3=sample(c('a','b','c'), size=50, replace=TRUE))
trial1 = pcr(x=dat1, nTreat=25, M=500, m=50)
# 1:1 allocation, M large
dat2 = data.frame(x1=rnorm(10),
                  x2=rnorm(10),
                  x3=sample(c('a','b','c'), size=10, replace=TRUE))
trial2 = pcr(x=dat2, nTreat=5, M=200, m=20)
# non-1:1 allocation, M small
dat3 = data.frame(x1=rnorm(50),
                  x2=rnorm(50),
                  x3=sample(c('a','b','c'), size=50, replace=TRUE))
trial3 = pcr(x=dat3, nTreat=35, M=200, m=20)
# non-1:1 allocation, M large
dat4 = data.frame(x1=rnorm(10),
                  x2=rnorm(10),
                  x3=sample(c('a','b','c'), size=10, replace=TRUE))
trial4 = pcr(x=dat4, nTreat=6, M=300, m=30)
```

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piFunction

Get PBA Treatment Probability

## **Description**

Used within calls to pba and pbaAgain to obtain the probability a unit is assigned treatment given its fitted propensity score.

## Usage

```
piFunction(fit, kparam, qparam)
```

## **Arguments**

fit Fitted propensity score.

kparam Balancing parameter.

qparam Global target for proportion of units treated.

#### **Details**

The input kparam must be one of 0, Inf, or the ratio of two positive odd integers. Both fit and qparam must be between 0 and 1.

#### Value

A numeric object. In the conext of PBA, the probability of assignment to treatment for the current unit.

## Author(s)

Travis M. Loux

## **Examples**

```
piFunction(fit=0.6, kparam=1, qparam=0.5)
piFunction(fit=0.6, kparam=5, qparam=0.5)
piFunction(fit=0.6, kparam=1/5, qparam=0.5)
piFunction(fit=0.6, kparam=1, qparam=2/3)
piFunction(fit=0.6, kparam=5, qparam=2/3)
piFunction(fit=0.6, kparam=1/5, qparam=2/3)
```

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plotpi

Plots of piFunction

## Description

Can be used to investigate the strength of balance forced by various values of the tuning parameter k.

## Usage

```
plotpi(k, global = 0.5)
addpi(k, global = 0.5, ...)
```

## **Arguments**

k Balancing parameter.

global Global target for proportion of units treated.
... Parameters in addpi passed to lines function.

## **Details**

The function plotpi creates a plot of treatment probability against fitted propensity score for values of k and global. The function addpi adds a curve for a new combination of k and global to an existing plot.

## Author(s)

Travis M. Loux

## Examples

```
plotpi(k=3, global=0.5)
addpi(k=5/3, lty=2, col='red')
plotpi(k=1/3, global=2/3)
```

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