# Package: RCT (via r-universe)

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**Title** Assign Treatments, Power Calculations, Balances, Impact Evaluation of Experiments

#### Version 1.2

**Description** Assists in the whole process of designing and evaluating Randomized Control Trials. Robust treatment assignment by strata/blocks, that handles misfits; Power calculations of the minimum detectable treatment effect or minimum populations; Balance tables of T-test of covariates; Balance Regression: (treatment ~ all x variables) with F-test of null model; Impact\_evaluation: Impact evaluation regressions. This function gives you the option to include control vars, fixed effect variables, cluster variables (for robust SE), multiple endogenous variables and multiple heterogeneous variables (to test treatment effect heterogeneity) summary\_statistics: Function that creates a summary statistics table with statistics rank observations in n groups: Creates a factor variable with n groups. Each group has a min and max label attach to each category. Athey, Susan, and Guido W. Imbens (2017) <arXiv:1607.00698>.

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**Encoding** UTF-8

**Imports** dplyr, purrr, glue, rlang, tidyr, stringr, MASS, pracma, estimatr, broom (>= 1.0.0), forcats, magrittr, ggplot2, utils, tidyselect (>= 1.0.0)

Suggests knitr, rmarkdown, testthat

RoxygenNote 7.3.1

VignetteBuilder knitr

Repository https://isidorogu.r-universe.dev

RemoteUrl https://github.com/isidorogu/rct

RemoteRef HEAD

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# Contents

balance_regression	2
balance_table	3
impact_eval	3
ntile_label	5
N_min	6
RCT	7
summary_statistics	8
tau_min	
tau_min_probability	10
treatment_assign	11
	13

# Index

balance\_regression() Runs a LPM of treatment status against all covariates (treatment~X'B).

# Description

balance\_regression() Runs a LPM of treatment status against all covariates (treatment~X'B).

#### Usage

balance\_regression(data, treatment)

#### Arguments

data	A data.frame, tibble or data.table
treatment	a string with treatment status column

#### Details

This functions runs a Linear Probability model of each treatment group & control on all the columns in data. For instance, if treatment column has values of (0,1,2), balance\_regression will run two models: 1) LPM(treatment(0,1)~X'b) and 2) LPM(treatment(0,2)~X'b). The value are the regression tables and details of the F\_test of these models.

#### Value

A list: regression\_tables = regression output of treatment against all covariates, F\_test = table with the F tests of each regression

#### Examples

data <-data.frame(x1 = rnorm(n = 100, mean = 100, sd = 15), x2= rnorm(n = 100, mean = 65), treat = rep(c(0,1,2,3,4), each = 20)) balance\_regression(data = data, treatment = "treat")

balance\_table

#### Description

Creates balance table for the X variables across treatment status

#### Usage

```
balance_table(data, treatment)
```

#### Arguments

data	A data.frame, tibble or data.table
treatment	a string with treatment status column

#### Details

balance\_table() performs t.test(X~treatment) for each X column in data. Every value of treatment i.e 1,2,3,...N is compared against control value (0) or the first value of the treatment column. For instance, If treatment column has values of (0,1,2,3), balance\_table will return: the mean value of each treatment (for all X's), and the p\_values of the t.test of (1,2,3) against treatment = 0.

## Value

A tibble with Mean\_value of each treatment status and p\_values

#### Examples

impact\_eval

Impact Evaluation of Treatment Effects

#### Description

Impact Evaluation of Treatment Effects

#### Usage

```
impact_eval(
   data,
   endogenous_vars,
   treatment,
   heterogenous_vars,
   fixed_effect_vars = NULL,
   control_vars,
   cluster_var
)
```

# Arguments

data	A data.frame, tibble or data.table	
endogenous_vars	5	
	Vector of Y's on which treatment effects will be evaluated	
treatment	Variable indicating the treatment status	
heterogenous_vars		
	Vector of variables for which you wish to assess treatment distributions/heterogeneities.	
fixed_effect_vars		
	Vector of variables to add as fixed effects. Default is without fixed effects	
control_vars	Vector of variables to control for in the evaluation. Default is without controls	
cluster_var	String of cluster variable the standard errors. Default is without clustered std errors	

# Details

This function carries out the evaluation of treatment effects on endogenous variables. It automatically runs the regressions of all the endogenous\_vars supplied & all the combinations of endogenous\_vars and heterogenous\_vars. Additionally, the function has the option of include fixed\_effects, controls and cluster variables for clustered std errors.

# Value

impact\_eval() returns a list of regression tables. The names of the list are the same as the endogenous variables. for heterogeneities the names are endogenous\_var\_heterogenous\_var

#### Examples

evaluation<-impact\_eval(data = data,</pre>

4

ntile\_label

ntile\_label() ranks observations in n groups, with labels

# Description

ntile\_label() ranks observations in n groups, with labels

#### Usage

ntile\_label(var, n, digits = 0)

#### Arguments

var	The variable wished to be ntile_label
n	rank the variable in n groups
digits	How many digits to include in the label

#### Details

 $n_{tile_label}$  is very similar to ntile from dplyr. But  $n_{tile_label}$  creates the n groups and then labels them. For each group i, the value of the ntile\_label is [min(i) - max(i)].

#### Value

A ordered factor vector of each n group. The value has the form of  $[\min(n_i) - \max(n_i)]$ 

N\_min

 $N_{min}()$  computes the minimum population needed to detect difference between control group and each treatment, given a target minimum detectable effect

# Description

N\_min() computes the minimum population needed to detect difference between control group and each treatment, given a target minimum detectable effect

#### Usage

```
N_min(
   outcome_var,
   tau_min,
   power = 0.8,
   significance = 0.05,
   share_control,
   n_groups = 2
)
```

#### Arguments

outcome_var	the variable for which you wish to test the impact of treatment
tau_min	the target detectable effect (in outcome_var units)
power	The level of power of the test (1 - $Pr(Reject H_0   H_0 True)$ ). Default is 0.8
significance	The level of significance of the test $Pr(Reject H_0   H_0 False)$ . Default is 0.05
share_control	The share of observations in N assigned to control. This argument allows for sequences (i.e. $seq(0,1,0.1)$ )
n_groups	Number of groups (control + # treatment groups)

#### Details

This function calculates the minimum experiment's population needed in order to detect at least a difference of tau\_min statistically significantly. This is between any two given groups (e.g. control vs each treatment), given the outcome variable, power and significance

#### Value

A tibble with the share\_control and N observations in control group (N\_control), the share and N of each treatment c(share\_ti, N\_ti), total share of treatment rows and N treated (share\_treat, N\_treat), N, the minimum detectable difference between control and all treatments together (tau\_min\_global), the minimum detectable difference between control and each treatment (tau\_min\_each\_treat)

#### RCT

#### Examples

RCT	Designing, random assigning and evaluating Randomized Control Tri-
	als

#### Description

RCT provides three important group of functions: a) functions for pre-processing the design of the RCT b) Functions for assigning treatment status and checking for balances c) Function for evaluating the impact of the RCT

#### Details

RCT helps you focus on the statistics of the randomized control trials, rather than the heavy programming lifting. RCT helps you in the whole process of designing and evaluating a RCT. 1. Clean and summarise the data in which you want to randomly assign treatment 2. Decide the share of observations that will go to control group 3. Decide which variables to use for strata building 4. Robust Random Assignment by strata/blocks 5 Impact evaluation of all y's and heterogeneities To lean more about RCT, start with the vignette: browseVignettes(package = "RCT")

#### **RCT** functions

treatment\_assign: Robust treatment assign by strata/blocks

impact\_eval: Automatized impact evaluation with heterogeneous treatment effects

balance\_table: Balance tables for any length of covariates

balance\_regression: LPM of treatment status against covariates with F-test

tau\_min: Computation of the minimum detectable effect between control & treatment units

tau\_min\_probability: Computation of the minimum detectable effect between control & treatment units for dichotomous y-vars

summary\_statistics: Summary statistics of all numeric columns in your data

ntile\_label: Rank and divide observations in n groups, with label

#### Author(s)

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#### References

Athey, Susan, and Guido W. Imbens (2017) "The Econometrics Randomized Experiments". Handbook of economic field experiments. https://arxiv.org/abs/1607.00698

## See Also

Useful links: https://github.com/isidorogu/RCT Report bugs at https://github.com/isidorogu/RCT/issues

summary\_statistics summary\_statistics() Creates summary statistics table of all numeric variables in data

# Description

summary\_statistics() Creates summary statistics table of all numeric variables in data

#### Usage

```
summary_statistics(
    data,
    probs = c(0, 0.05, 0.1, 0.25, 0.5, 0.75, 0.9, 0.95, 1),
    na.rm = T
)
```

# Arguments

data	A data.frame, tibble or data.table
probs	The quantiles to compute. Default is c(0, 0.05, 0.1, 0.25, 0.5, 0.75, 0.9, 0.95, 1)
na.rm	whether to exclude NA's from calculations

# Details

This function computes the selected quantiles, mean and N values of all the numeric columns of data.

#### Value

A tibble with the Mean, N (not NA) and probs selects for each numeric column

```
data <-data.frame(x = c(1:5), y = c(100, 200, 300, 410, 540), z = rep("c", 5))
summary_statistics(data)</pre>
```

tau\_min

*tau\_min() computes the minimum detectable difference between control group and each treatment* 

#### Description

tau\_min() computes the minimum detectable difference between control group and each treatment

#### Usage

```
tau_min(
   outcome_var,
   N,
   power = 0.8,
   significance = 0.05,
   share_control,
   n_groups = 2
)
```

#### Arguments

outcome_var	the variable for which you wish to test the impact of treatment
Ν	number of observations in the RCT, usually nrow(data)
power	The level of power of the test (1 - $Pr(Reject H_0   H_0 True)$ ). Default is 0.8
significance	The level of significance of the test Pr(Reject H_0   H_0 False). Default is 0.05
share_control	The share of observations in N assigned to control. This argument allows for sequences (i.e. $seq(0,1,0.1)$ )
n_groups	Number of groups (control + # treatment groups)

#### Details

This function calculates the minimum difference that could show significant E[Y(1)-Y(0)] = tau, between any two given groups (e.g. control vs each treatment), given the population size (N), the outcome variable, power and significance

#### Value

A tibble with the share\_control and N observations in control group (N\_control), the share and N of each treatment c(share\_ti, N\_ti), total share of treatment rows and N treated (share\_treat, N\_treat), N, the minimum detectable difference between control and all treatments together (tau\_min\_global), the minimum detectable difference between control and each treatment (tau\_min\_each\_treat)

tau\_min\_probability tau\_min\_probability() computes the minimum detectable difference between control group and each treatment for a dichotomous variable

# Description

tau\_min\_probability() computes the minimum detectable difference between control group and each treatment for a dichotomous variable

#### Usage

```
tau_min_probability(
   prior,
   N,
   power = 0.8,
   significance = 0.05,
   share_control,
   n_groups = 2
)
```

#### Arguments

prior	Pr(Y=1).
Ν	number of observations in the RCT, usually nrow(data)
power	The level of power of the test (1 - Pr(Reject H_0   H_0 True) ). Default is 0.8
significance	The level of significance of the test Pr(Reject H_0   H_0 False). Default is 0.05
share_control	The share of observations in N assigned to control. This argument allows for sequences (i.e. $seq(0,1,0.1)$ )
n_groups	Number of groups (control + # treatment groups)

#### Details

This function calculates the minimum difference that could show significant Pr[Y(1)-Y(0)] = tau, between any two given groups (e.g. control vs each treatment), given the population size (N), the outcome variable, power and significance

# Value

A tibble with the share\_control and N observations in control group (N\_control), the share and N of each treatment c(share\_ti, N\_ti), total share of treatment rows and N treated (share\_treat, N\_treat), N, the minimum detectable difference between control and all treatments together (tau\_min\_global), the minimum detectable difference between control and each treatment (tau\_min\_each\_treat)

```
tau_min_probability(0.4, N = 1000, share_control = seq(0,1,0.1), n_groups = 3)
```

treatment\_assign treatment\_assign() carries out robust treatment assignment by strata/blocks

## Description

treatment\_assign() carries out robust treatment assignment by strata/blocks

#### Usage

```
treatment_assign(
   data,
   share_control,
   n_t = 2,
   strata_varlist,
   missfits = c("global", "NA", "strata"),
   seed = 1990,
   share_ti = rep(1/n_t - share_control/n_t, times = n_t),
   key
)
```

#### Arguments

data	A data.frame, tibble or data.table
share_control	share of the observations assigned to control group
n_t	Number of treatments groups
strata_varlist	vector of categorical variables to form the strata/blocks for random assignment. Should be in the form of vars(var1, var2,)
missfits	How to handle the misfits. Default is "global". See Carril (2016) for details.
seed	A number used to set.seed().
share_ti	The share of each treatment group. If NULL (Default), each treatment group will have equal share.
key	The key identifier column of data.

#### Details

This function creates a variable that indicates the treatment status. The random assignment is made by strata/blocks. It can handle equal or unequal treatment shares. Finally, it has three methods available to handle misfits (same as randtreat in STATA): "global": assigning the observations that couldn't be randomly assigned globally, "strata": assigning the observations that couldn't be randomly assigned by strata, "NA": set the the treat observations that couldn't be randomly assigned to NA.

#### Value

A list: "data" = the data with key, treat, strata, misfit column., "summary\_strata" = A summary tibble with the membership of each strata and its size.

# Index

balance\_regression, 2
balance\_table, 3

 $impact_eval, 3$ 

N\_min,6 ntile\_label,5

RCT, 7 RCT-package (RCT), 7

summary\_statistics, 8

tau\_min,9
tau\_min\_probability,10
treatment\_assign,11