

# Cheatsheets

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# Przykładowe cheatsheety



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## RStudio Cheat Sheets

The cheat sheets below make it easy to learn about and use some of our favorite packages. From time to time, we will add new cheat sheets to the gallery. If you'd like us to drop you an email when we do, let us know by clicking the button to the right.

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### Work with Strings Cheat Sheet

The `stringr` package provides an easy to use toolkit for working with strings, i.e. character data, in R. This cheatsheet guides you through `stringr`'s functions for manipulating strings. The back page provides a concise reference to *regular expressions*, a mini-language for describing, finding, and matching patterns in strings. Updated 10/17.

DOWNLOAD

The `stringr` package provides a set of internally consistent tools for working with character strings, i.e. sequences of characters surrounded by quotation marks.

#### Detect Matches

- `str_detect(string, pattern)` Detect the presence of a pattern match in a string. `str_detect_full(1, 8)` or `str_detect(2)`
- `str_which(string, pattern)` Find the indices of strings that contain a pattern match. `str_which_full(1, 8)` or `str_which(2)`
- `str_count(string, pattern)` Count the number of matches in a string. `str_count_full(1, 8)` or `str_count(2)`
- `str_locate(string, pattern)` Locate the positions of pattern matches in a string. Also `str_locate_all(1)` or `str_locate_full(1, 8)`

#### Subset Strings

- `str_sub(string, start = 1L, end = 1L)` Extract substrings from a character vector. `str_sub_full(1, 8)` or `str_sub(2)`
- `str_subset(string, pattern)` Return only the strings that contain a pattern match. `str_subset_full(1, 8)` or `str_subset(2)`
- `str_extract(string, pattern)` Return the first pattern match found in each string, as a vector. Also `str_extract_all(1)` or `str_extract_full(1, 8)`
- `str_match(string, pattern)` Return the first pattern match found in each string, as a matrix with a column for each (1) group in pattern. Also `str_match_all(1)` or `str_match_full(1, 8)`

#### Manage Lengths

- `str_length(string)` The width of strings (i.e. number of code points, which generally equals the number of characters). `str_length_full(1)`
- `str_pad(string, width, side = c("left", "right", "both"), pad = " ")` Pad strings to constant width. `str_pad_full(1, 8)`
- `str_trunc(string, width, side = c("left", "right", "center"), ellipsis = "...")` Truncate the width of strings, replacing content with ellipsis. `str_trunc_full(1, 8)`
- `str_trim(string, side = c("both", "left", "right"))` Trim whitespace from the start and/or end of a string. `str_trim_full(1)`

#### Mutate Strings

- `str_sub() <- value` Replace substrings by identifying the substrings with `str_sub()` and assigning into the results. `str_sub_full(1, 8) <- "foo"`
- `str_replace(string, pattern, replacement)` Replace the first matched pattern in each string. `str_replace_full(1, 8) <- "foo"`
- `str_replace_all(string, pattern, replacement)` Replace all matched patterns in each string. `str_replace_all_full(1, 8) <- "foo"`
- `str_to_lower(string, locale = "en")` Convert strings to lower case. `str_to_lower_full(1, 8)`
- `str_to_upper(string, locale = "en")` Convert strings to upper case. `str_to_upper_full(1, 8)`
- `str_to_title(string, locale = "en")` Convert strings to title case. `str_to_title_full(1, 8)`

#### Join and Split

- `str_c(..., sep = "", collapse = NULL)` Join multiple strings into a single string. `str_c_full(1, 8)` or `str_c(2)`
- `str_glue(..., sep = "", collapse = NULL)` Collapse a vector of strings into a single string. `str_glue_full(1, 8)` or `str_glue(2)`
- `str_split(string, pattern)` Split a string into a vector of substrings (splitting at occurrences of a pattern match). Also `str_split_all(1)` or `str_split_full(1, 8)`
- `str_split_full(string, pattern)` Split a vector of strings into a matrix of substrings (splitting at occurrences of a pattern match). Also `str_split_full_all(1)` or `str_split_full_full(1, 8)`

#### Order Strings

- `str_sort(string, decreasing = FALSE, na.last = "first", locale = "en", na.rm = FALSE)` Sort a character vector. `str_sort_full(1, 8)`
- `str_order(string, decreasing = FALSE, na.last = "first", locale = "en", na.rm = FALSE)` Sort a character vector. `str_order_full(1, 8)`

#### Helpers

- `str_conv(string, encoding)` Override the encoding of a string. `str_conv_full(1, 8)` or `str_conv(2)`
- `str_detect(string, pattern, match = NA)` View `NA`, encoding of first regex match in each string. `str_detect_full(1, 8)`
- `str_view(string, pattern, match = NA)` View `NA`, encoding of all regex matches. `str_view_full(1, 8)`
- `str_wrap(string, width = 80, indent = 0, exdent = 0)` Wrap strings into nicely formatted paragraphs. `str_wrap_full(1, 8)`

RStudio is a trademark of RStudio, Inc. © 2017 RStudio. All rights reserved. Learn more at <https://www.rstudio.com>. Cheatsheets from <https://www.rstudio.com/cheatsheets/>. Stringr 1.2.0, updated 2017-10-17.

<https://www.rstudio.com/resources/cheatsheets/>

MI

**Dlaczego warto je tworzyć, po co są  
nam one potrzebne w naszej  
grupie?**

# Nasze cheatsheety

## factorMerger Cheat Sheet

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Przemysław Biecek [aut, ths]  
University of Warsaw



### Introduction

How to check if averages are different among k groups? Use ANOVA!

How to visualise how these groups are different? Use factorMerger!

The aim of factorMerger is to provide informative and easy to understand visualisations of post-hoc comparisons. It gives consistent and non-overlapping adaptive fusing of groups based on likelihood ratio test (LRT). The package factorMerger works for wide spectrum of families like Gaussian, binomial or survival.

Results from the adaptive fusing are presented with the Merging Paths Plots - a hierarchical representation of LRT-based distances among groups.

In addition, the Generalized Information Criterion (GIC) is presented for fused models. This criterion may be used to choose the optimal segmentation of groups.

Graphical summary of the variable of interest in each group is presented in the right panel.

Find more in <https://arxiv.org/abs/1709.04412>

### Example

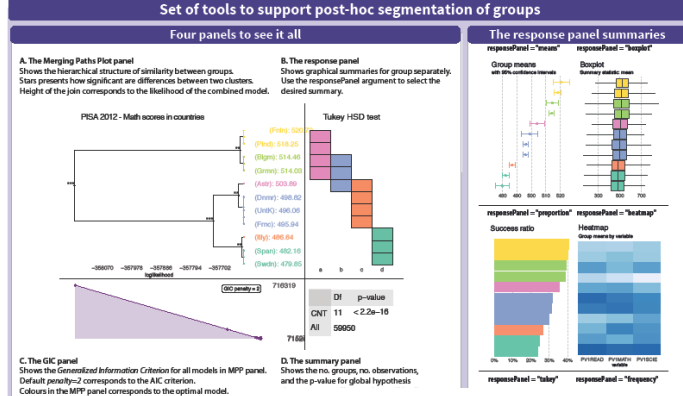
library(factorMerger)

```
fmAll <- mergeFactors(
  response = pisaEuromath,
  factor = pisaEurocountry,
  method = "fast-adaptive",
  family = "gaussian")
```

```
print(fmAll)
```

```
plot(fmAll,
  panel = "all",
  responsePanel = "tukey")
```

factorMerger in version 0.3.2 (2017) Agnieszka Sikko, Przemysław Biecek <https://cran.r-project.org/package=randomForestExplainer> - CC BY



## MLExpResso

### Cheat Sheet

Aleksandra Dąbrowska [aut, cre]  
Alię Gosiewska [aut]  
Przemysław Biecek [aut, ths]



### Introduction

MLExpResso is an R package for integrative analyses and visualization of gene expression and DNA methylation data.

Key functions of this package are:

- identification of genes with affected expression - calculate\_test() function,
- identification of DMR - differentially methylated regions - calculate\_test() function,
- identification of regions with changes in expression and methylation - calculate\_comparison\_table() function,
- visualization of identified regions - plot\_gene() and plot\_volcano() functions.

The joint modeling and visualization of genes expression and methylation improve interpretability of identified signals.

### MLExpRessoData

The methodology is supplemented with sample applications to The Cancer Genome Atlas data.

MLExpRessoData is an R package which contains information from The Cancer Genome Atlas (TCGA) Data Portal. Data sets in this package are based on Bioconductor package RTCGA. In examples, we use both, methylation and expression data.

• BRCA\_exp - it contains information about gene expression: read counts per gene, computed for genes for 756 patients with breast cancer. Rows of this data set correspond to samples taken from patients. First column SUBTYPE corresponds to a subtype of BRCA cancer, next columns correspond to genes.

• BRCA\_met - it contains information about methylation of CpG probes for patients with breast cancer. Rows of this data set correspond to patients, more precisely, to samples taken from patients. First column SUBTYPE corresponds to a subtype of BRCA cancer, next columns correspond to CpG probes. Values inside the table indicate the percentage methylation level of CpG probe for a specified sample.

For aggregation CpG probes to correspond genes we use the Illumina Human Methylation data set from TCGA. Haplotype UCSB has knownGene Bioconductor package.

MLExpResso in version 0.2.0(2017) Aleksandra Dąbrowska, Alię Gosiewska, Przemysław Biecek - CC BY

## randomForestExplainer What's in the forest?

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Przemysław Biecek [aut, ths]  
University of Warsaw



### Basics

The aim of the randomForestExplainer package is to support structure exploration and visualization for a random forest model.

Once you have a model created with the randomForest package, use following functions to examine its structure.

```
library(randomForest)
library(randomForestExplainer)

forest <- randomForest(PYJMATH~X,
  data = pisa2015, localImp = TRUE)
```

### Variable Importance

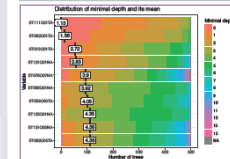
The measure\_importance() function calculates different measures of importance for variables presented in the forest. Note that different variables are available for classification forests and regression forests.

## randomForestExplainer - Structure mining and visualisation for Random Forests

### Variable Depth

The min\_depth\_distribution() function calculates distribution of minimal depth of given variable in all trees. Use the plot\_min\_depth\_distribution() function to plot this distribution along with mean depths for variables. In general, the higher are variables the more influential they are.

```
forest_frame <-
min_depth_distribution(forest)
plot_min_depth_distribution(forest_frame)
```

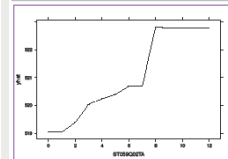


The min\_depth\_distribution() function

### Partial Dependence Plots

The partial() function from the pdp package calculates marginal relation between target variable and selected one or two independent variables. The relation can be noted with lattice graphical system with the plotPartial() function or with ggplot2 system with the autoplot() function.

```
library(pdp)
pdp1 <- partial(forest, c("ST059082TA"))
plotPartial(pdp1)
```



Variable depth and variable importance are useful in identification which ble/variables are worth watching, while the plots are useful to understand the nature of relation between target variable and ble/s of interest.

## Identification of genes with affected expression

MLExpResso: calculate\_test(data, condition, test)

Function calculate\_test() computes log folds, p-values and means for chosen test for both, methylation and expression data.

```
> BRCA_exp[1:3, 1:5]
```

```
TCGA AL A05B 01A ILR AL44 07 SUBTYPE AANAL A05D01 AAIL A1X
TCGA AL A05D 01A ILR AL15 07 LUM1 2 2354 2670 317
TCGA AL A05D 01A ILR AL15 07 LUM1 2 1840 5076 312
TCGA AL A05D 01A ILR AL15 07 LUM1 11 3391 9527 736
```

### Example

```
library("MLExpResso")
library("MLExpRessoData")
exp <- BRCA_exp[, 1:]
gr_exp <- BRCA_exp$SUBTYPE
gr_exp <- ifelse(gr_exp == 'LumA', 'LumA', 'other')
res_exp <- calculate_test(exp, gr_exp, 'lrr')
```

```
> head(res_exp)
  id log2_fold pval mean_LumA mean_other mean
1 A05D01 2.3 3.2e-32 539 2324 2485
2 CEN2 2.9 2.4e-26 633 4297 2574
3 KPN1 1.4 8.6e-24 1154 7647 1644
4 PKN1 3.8 2.3e-22 396 3480 2031
5 BIRC5 2.0 2.0e-21 1957 6658 1419
6 C57 1.4 5.3e-21 778 6279 484
```

## Identification of DMR - differentially methylated regions

MLExpResso: calculate\_test(data, condition, test)

Argument test allows using two different statistic tests for finding differences in methylation levels. All available values are in the table on the right.

MLExpResso: aggregate\_probes(data)

Function aggregate\_probes() aggregates CpG probes to corresponding genes using, by default, the Illumina human methylation data.

```
> BRCA_met[1:3, 1:5]
```

```
HUGO AL A05B 01A ILR AL15 07 SUBTYPE CEN2 CEN201102 CEN201102 CEN201102
TCGA AL A05D 01A ILR AL15 07 LUM1 0 0.00 0.24 0.308 0.028
TCGA AL A05D 01A ILR AL15 07 KPN1 0.014 0.70 0.308 0.014
```

### Example

```
met <- aggregate_probes(BRCA_met)
gr_met <- BRCA_met$SUBTYPE
gr_met <- ifelse(gr_met == 'LumA', 'LumA', 'other')
res_met <- calculate_test(met, gr_met, 'ttest')
> head(res_met)
  id log2_fold pval mean_LumA mean_other mean
1 LCAV2 -0.152 3.8e-17 0.25 0.41 0.33
2 R2L1 0.001 2.6e-13 0.31 0.36 0.33
3 PTPRX 0.115 5.4e-12 0.47 0.31 0.36
4 INH32 -0.134 5.9e-12 0.18 0.31 0.25
5 CD7 0.098 1.6e-11 0.86 0.77 0.81
6 KSR1 0.700 7.1e-11 0.66 0.46 0.55
```

Argument test allows using many different statistic tests for finding differences in expression. All available values are in the table below.

```
Value Test
test student's t-test
Yajman2 negative binomial test
Yajman2 likelihood-ratio test
Yajman2 quasi-likelihood F-test
```

MLExpResso: plot\_diff\_boxplot(data, condition, gene)

Function plot\_diff\_boxplot() generates a boxplot of values from chosen data frame column with division in groups (two or more).

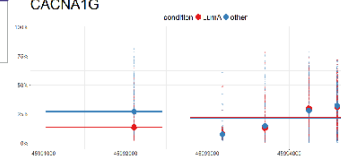
```
plot_diff_boxplot(data = exp,
  condition = gr_exp,
  gene = 'CACNA1G')
```

### Example

MLExpResso: plot\_methylation\_path(data, condition, gene)

Function plot\_methylation\_path() visualizes a chosen gene with marked CpG probes. Y axis describes methylation level. X axis describes a location of the probe on the chromosome. Horizontal lines show the mean methylation level for each island in a division to groups. Groups are defined by colors. Large dots symbolize means of methylation level for CpG probes, small dots symbolize methylation levels for each observation.

```
plot_methylation_path(data = BRCA_met,
  condition = gr_met,
  gene = 'CACNA1G',
  observ = T)
```



### Literature

ggRandomForests: Random Forests for Regression. John Eltinger (2016)

pdp: An R Package for Constructing Partial Dependence Plots. Brandon M. Greenwell (2017)

forestFloor: Forest Floor Visualizations of Random Forests. Soeren Welling, Hanne Rejsgard, Per Brockhoff, Line Clemmensen (2016)



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# Jak je zrobić?



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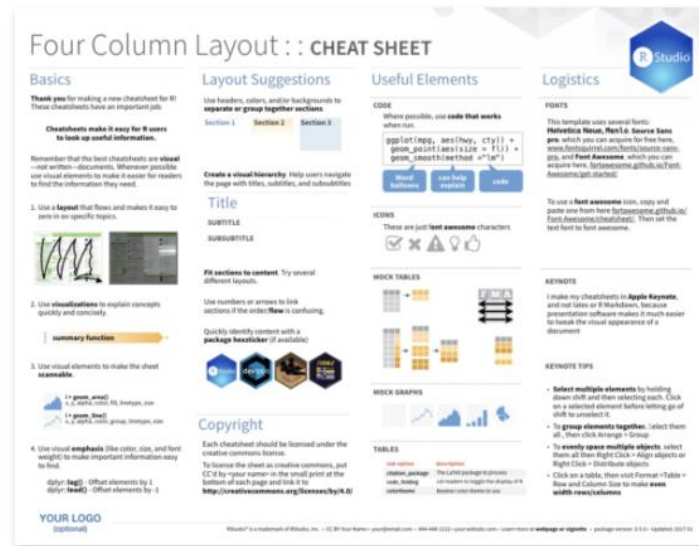
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Want to contribute a cheatsheet of your own?

We'd like to help you make and share high quality cheatsheets on R topics. The template below provides a useful starting place. It contains tips for designing a three or four column cheatsheet, as well as reusable elements to build your sheet with.



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<https://www.rstudio.com/resources/cheatsheets/how-to-contribute-a-cheatsheet/>

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# Stary szablon

## Four Column layout Cheat Sheet

Your LOGO

### Basics

Thank you for making a new cheatsheet for R! These cheatsheets have an important job: Cheatsheets make it easy for R users

to look up useful information.

Remember that the best cheatsheets are **visual**—not written—documents. Whenever possible use visual elements to make it easier for readers to find the information they need.

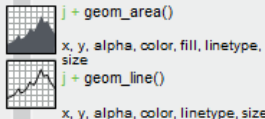
1. Use a **layout** that flows from top to bottom and left to right to make it easy to zero in on specific topics.



2. Use **visualizations** to explain concepts quickly and concisely.

summary function

3. Use **visual elements** to make the sheet scannable.



4. Use **visual emphasis** (like color, size, and weight) to make important information easy to find.

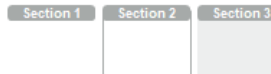
```
dplyr::bind_rows(y, z)
```

Append z to y as new rows.

Title - Group sections with titles, subtitles, and subsubtitles to create a visual hierarchy

### Layout suggestions

Use headers, outlines, and/or backgrounds to **separate or group together sections**.



Use titles, subtitles, and subsubtitles to **create a visual hierarchy** that will help users navigate the page.

### Title

#### Subtitle

Fit **sections to content**. Try several different layouts.

Use numbers or arrows to link sections if the **order/flow** is confusing.

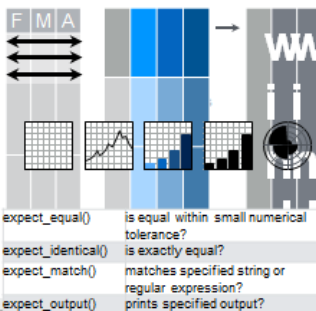
### Useful elements

#### icons



These are just font awesome characters

#### Mock tables



### Copyright

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### Subtitle

#### Example code

Where possible, use **code that works** when run.

```
dplyr::lead
Copy with values shifted by 1.
dplyr::lag
Copy with values lagged by 1.
dplyr::dense_rank
Ranks with no gaps.
dplyr::min_rank
Ranks. Ties get min rank.
dplyr::percent_rank
Ranks rescaled to [0, 1].
dplyr::row_number
Ranks. Ties got to first value.
dplyr::ntile
Bin vector into n buckets.
dplyr::between
Are values between a and b?
dplyr::cume_dist
Cumulative distribution.
```

#### Color Scheme

Please use the following **color scheme** when designing new cheatsheets to be distributed through <http://www.rstudio.com/resources/cheatsheets/>



#### Keynote

I make my cheatsheets in **Apple Keynote**, and not latex or R Markdown, because presentation software makes it much easier to tweak the visual appearance of a document

#### Keynote tips

- **Select multiple elements** by holding down shift and then selecting each. Click on a selected element before letting go of shift to unselect it.
- **To group elements together**, select them all, then click Arrange > Group
- **To evenly space multiple objects**, select them all then Right Click > Align objects or Right Click > Distribute objects
- Click on a table, then visit Format > Table > Row and Column Size to make **even width rows/columns**.

#### Code snippets

```
ggplot(mpg, aes(hwy, cty)) +
  geom_point(aes(color = cty)) +
  geom_smooth(method = "lm") +
  coord_cartesian() +
  scale_color_gradient() +
  theme_bw()
```

Word balloons

can be useful for

explaining code

### Fonts

This template uses several fonts: **Helvetica Neue**, **Menlo**, **Source Sans pro**, which you can acquire for free here, <http://www.fontsquirrel.com/fonts/source-sans-pro>, and **Font Awesome**, which you can acquire here, <http://fontawesome.github.io/Font-Awesome/get-started/>. To use a font awesome icon, copy and paste one from here <http://fontawesome.github.io/Font-Awesome/cheatsheet/>. Then set the text font to font awesome.



# Składowe R-owej ściągawki

## Four Column Layout : : CHEAT SHEET



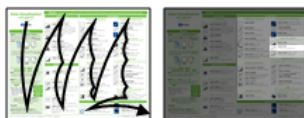
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2. Use **visualizations** to explain concepts quickly and concisely.

summary function

3. Use visual elements to make the sheet **scannable**.

```
i + geom_area()
x, y, alpha, color, fill, linetype, size

i + geom_line()
x, y, alpha, color, group, linetype, size
```

4. Use visual **emphasis** (like color, size, and font weight) to make important information easy to find.

```
dplyr::lag() - Offset elements by 1
dplyr::lead() - Offset elements by -1
```

YOUR LOGO  
(optional)

### Layout Suggestions

Use headers, colors, and/or backgrounds to **separate or group together** sections.



Create a **visual hierarchy**. Help users navigate the page with titles, subtitles, and subtitles.

Title

SUBTITLE

SUBSUBTITLE

Fit **sections to content**. Try several different layouts.

Use numbers or arrows to link sections if the **order/flow** is confusing.

Quickly identify content with a **package hexsticker** (if available)



### Copyright

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### Useful Elements

#### CODE

Where possible, use **code that works** when run.

```
ggplot(mpg, aes(hwy, cty)) +
  geom_point(aes(size = fl)) +
  geom_smooth(method = "lm")
```

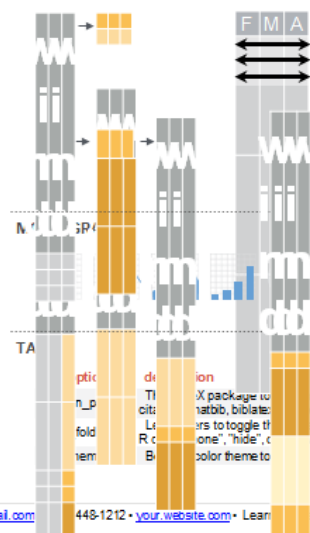
Word balloons can help explain code

#### ICONS

These are just **fontawesome** characters



#### MOCK TABLES



### Logistics

#### FONTS

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To use a **font awesome** icon, copy and paste one from here [fontawesome.github.io/Font-Awesome/cheatsheet/](https://fontawesome.github.io/Font-Awesome/cheatsheet/). Then set the text font to font awesome.

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- To **evenly space multiple objects**, select them all then Right Click > Align objects or Right Click > Distribute objects
- Click on a table, then visit Format > Table > Row and Column Size to make **even width rows/columns**.

# Praca w grupach

<https://github.com/kapelner/ICEbox>

<https://github.com/AppliedDataSciencePartners/xgboostExplainer>

<https://github.com/cran/nlstools>