talk04 练习与作业

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0.1 练习和作业说明

将相关代码填写入以"'{r}" 标志的代码框中,运行并看到正确的结果; 完成后,用工具栏里的"Knit" 按键生成 PDF 文档;

将 PDF 文档改为: 姓名-学号-talk04 作业.pdf,并提交到老师指定的平台/钉群。

0.2 Talk04 内容回顾

待写 ...

0.3 练习与作业: 用户验证

请运行以下命令,验证你的用户名。

如你当前用户名不能体现你的真实姓名,请改为拼音后再运行本作业!

```
Sys.info()[["user"]]
```

[1] "mingyuwang"

```
Sys.getenv("HOME")
```

[1] "C:/Users/rhong/Documents"

0.4 练习与作业 1: R session 管理

0.4.1 完成以下操作

- 定义一些变量(比如 x, y, z 并赋值; 内容随意)
- 从外部文件装入一些数据(可自行创建一个 4 行 5 列的数据,内容随意)
- 保存 workspace 到.RData
- 列出当前工作空间内的所有变量
- 删除当前工作空间内所有变量
- 从.RData 文件恢复保存的数据
- 再次列出当前工作空间内的所有变量,以确认变量已恢复
- 随机删除两个变量
- 再次列出当前工作空间内的所有变量

```
## 代码写这里,并运行;
x <- 1
y <- 2
z <- 3
data <- read.table("data/Table1.txt", header = TRUE)
# save.image(file = ".RData")
```

```
rm(list = ls())
ls()

## character(0)

load(file = ".RData")
ls()

## [1] "data" "x" "y" "z"

rm(list = c("x", "y"))
ls()

## [1] "data" "z"

0.5 练习与作业 2: Factor 基础
```

0.5.1 factors 增加

• 创建一个变量:

x <- c("single", "married", "married", "single");</pre>

- 为 x 增加两个 levels, single, married;
- 以下操作能成功吗?

 $x[3] \leftarrow "widowed";$

• 如果不,请提供解决方案;

```
## 代码写这里,并运行;
x <- c("single", "married", "single")
x <- factor(x, levels = c("single", "married"))
try(x[3] <- "widowed")

## Warning in `[<-.factor`(`*tmp*`, 3, value = "widowed"): 因子层次有错,产生了NA

# 解决方案
x <- factor(x, levels = c("single", "married", "widowed"))
try(x[3] <- "widowed")
x

## [1] single married widowed single
## Levels: single married widowed</pre>
```

0.5.2 factors 改变

• 创建一个变量:

```
v = c("a", "b", "a", "c", "b")
```

- 将其转化为 factor, 查看变量内容
- 将其第一个 levels 的值改为任意字符,再次查看变量内容

```
## 代码写这里,并运行;
v <- c("a", "b", "a", "c", "b")
v <- factor(v)
v
```

[1] a b a c b
Levels: a b c

```
levels(v)[1] <- "d"
## [1] d b d c b
## Levels: d b c
  • 比较改变前后的 v 的内容, 改变 levels 的操作使 v 发生了什么变化?
答: 改变 levels 的操作使 v 的内容发生了变化,将原来的"a" 改为了"d"。
0.5.3 factors 合并
  • 创建两个由随机大写字母组成的 factors
  • 合并两个变量, 使其 factors 得以在合并后保留
## 代码写这里,并运行;
x <- factor(sample(LETTERS, 10, replace = TRUE))</pre>
y <- factor(sample(LETTERS, 10, replace = TRUE))
## [1] B C I Z F Y W A U E
## Levels: A B C E F I U W Y Z
## [1] T X Z Z U I R C L X
## Levels: C I L R T U X Z
c(x, y)
## [1] BCIZFYWAUETXZZUIRCLX
```

Levels: A B C E F I U W Y Z L R T X

0.5.4 利用 factor 排序

以下变量包含了几个月份,请使用 factor,使其能按月份,而不是英文字符串排序:

mon <- c("Mar","Nov","Mar","Aug","Sep","Jun","Nov","Nov","Oct","Jun","May","Sep","Dec",</pre>

[1] Mar Mar May Jun Jul Aug Sep Sep Oct Nov Nov Nov Nov Dec
Levels: Jan Feb Mar Apr May Jun Jul Aug Sep Oct Nov Dec

0.5.5 forcats 的问题

forcats 包中的 fct_inorder, fct_infreq 和 fct_inseq 函数的作用是什么? 请使用 forcats 包中的 gss_cat 数据举例说明

```
## 代码写这里,并运行;
library(forcats)
```

Warning: 程辑包'forcats'是用R版本4.1.3 来建造的

```
head(gss_cat)
```

```
## year marital age race rincome partyid
## 1 2000 Never married 26 White $8000 to 9999 Ind,near rep
## 2 2000 Divorced 48 White $8000 to 9999 Not str republican
```

```
## 3 2000
               Widowed 67 White Not applicable
                                                       Independent
\#\# 4 2000 Never married 39 White Not applicable
                                                      Ind, near rep
## 5 2000
              Divorced 25 White Not applicable
                                                  Not str democrat
## 6 2000
               Married 25 White $20000 - 24999
                                                   Strong democrat
##
                 relig
                                  denom tvhours
## 1
            Protestant Southern baptist
                                             12
## 2
            Protestant Baptist-dk which
                                             NA
## 3
            Protestant No denomination
                                              2
## 4 Orthodox-christian
                         Not applicable
                                              4
## 5
                         Not applicable
                  None
                                              1
## 6
            Protestant Southern baptist
                                             NA
# fct_inord er: 按出现顺序为 levels 排序
fct_inorder(gss_cat$marital) %>% levels()
## [1] "Never married" "Divorced"
                                      "Widowed"
                                                      "Married"
## [5] "Separated"
                      "No answer"
# fct_infreq: 按出现频率为 levels 排序, 出现频率高的拍在前面
fct_infreq(gss_cat$marital) %>% levels()
## [1] "Married"
                       "Never married" "Divorced"
                                                      "Widowed"
## [5] "Separated"
                       "No answer"
# fct_inseq: 根据 level 的数字大小为 levels 排序,要求 factor levels 为数字
factor(gss_cat$age, levels = 80:60) %>% levels()
   [1] "80" "79" "78" "77" "76" "75" "74" "73" "72" "71" "70" "69" "68" "67" "66"
## [16] "65" "64" "63" "62" "61" "60"
factor(gss_cat$age, levels = 80:60) %>% fct_inseq() %>% levels()
   [1] "60" "61" "62" "63" "64" "65" "66" "67" "68" "69" "70" "71" "72" "73" "74"
## [16] "75" "76" "77" "78" "79" "80"
```

0.6 练习与作业 3: 用 mouse genes 数据做图

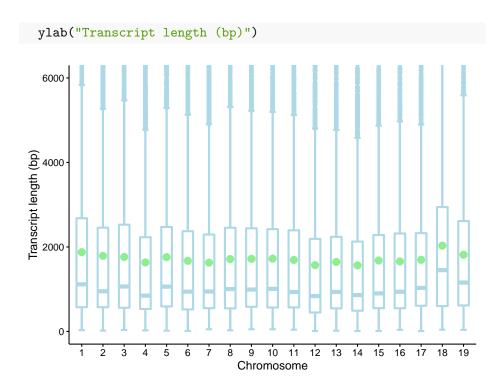
0.6.1 画图

1. 用 readr 包中的函数读取 mouse genes 文件(从本课程的 Github 页 面下载 data/talk04/)

- 2. 选取常染色体的基因
- 3. 画以下两个基因长度 boxplot:
- 按染色体序号排列, 比如 1, 2, 3 X, Y
- 按基因长度中值排列, 从短 -> 长 ...

```
## 代码写这里,并运行;
# 不显示 warning 信息 和 message
options(warn = -1, message = -1)
library(readr)
library(ggplot2)
library(dplyr)
##
## 载入程辑包: 'dplyr'
## The following objects are masked from 'package:stats':
##
##
      filter, lag
## The following objects are masked from 'package:base':
##
      intersect, setdiff, setequal, union
##
```

```
options(warn = 0, message = 0)
mouse_genes <- read_tsv("../data/talk04/mouse_genes_biomart_sep2018.txt",</pre>
  col_names = TRUE, show_col_types = FALSE)
colnames(mouse_genes) <- gsub(" ", "_", colnames(mouse_genes))</pre>
colnames(mouse_genes) <- gsub("/", "_", colnames(mouse_genes))</pre>
colnames(mouse_genes) <- gsub("\\(", "", colnames(mouse_genes))</pre>
colnames(mouse_genes) <- gsub("\\)", "", colnames(mouse_genes))</pre>
autosome_genes <- filter(mouse_genes, Chromosome_scaffold_name %in% 1:19)
# 按染色体序号排列
ggplot(autosome_genes, aes(x = factor(as.numeric(Chromosome_scaffold_name)),
  y = Transcript_length_including_UTRs_and_CDS)) +
  stat_boxplot(geom = "errorbar", width = 0.3, lwd = 1, color = "lightblue") +
  geom_boxplot(width = 0.5, lwd = 1, color = "lightblue") +
  theme minimal() +
  theme(panel.grid.major.x = element blank(),
    panel.grid.minor.x = element_blank(),
    panel.grid.major.y = element_blank(),
    panel.grid.minor.y = element_blank(),
    panel.background = element_blank(),
    panel.border = element blank(),
    axis.line = element line(color = "black", size = 0.5),
    axis.ticks = element_line(color = "black", size = 0.5),
    axis.text.x = element_text(size = 10, color = "black"),
    axis.text.y = element_text(size = 10, color = "black"),
    axis.title.x = element_text(size = 12, color = "black"),
    axis.title.y = element_text(size = 12, color = "black"),
    legend.position = "none") +
  coord_cartesian(ylim = c(0, 6000)) +
  stat_summary(fun = mean, geom = "point", shape = 20,
    size = 5, color = "lightgreen", fill = "lightgreen") +
  xlab("Chromosome") +
```



```
#按基因长度 中值 排列,从短 => 长
ggplot(autosome_genes, aes(x = reorder(Chromosome_scaffold_name,
    Transcript_length_including_UTRs_and_CDS, median),
    y = Transcript_length_including_UTRs_and_CDS)) +
  stat_boxplot(geom = "errorbar", width = 0.3, lwd = 1, color = "lightblue") +
  geom_boxplot(width = 0.5, lwd = 1, color = "lightblue") +
  theme_minimal() +
  theme(panel.grid.major.x = element_blank(),
    panel.grid.minor.x = element_blank(),
    panel.grid.major.y = element_blank(),
    panel.grid.minor.y = element_blank(),
    panel.background = element_blank(),
    panel.border = element_blank(),
    axis.line = element_line(color = "black", size = 0.5),
    axis.ticks = element_line(color = "black", size = 0.5),
    axis.text.x = element_text(size = 10, color = "black"),
```

```
axis.text.y = element_text(size = 10, color = "black"),
axis.title.x = element_text(size = 12, color = "black"),
axis.title.y = element_text(size = 12, color = "black"),
legend.position = "none") +
coord_cartesian(ylim = c(0, 6000)) +
stat_summary(fun = mean, geom = "point", shape = 20,
size = 5, color = "lightgreen", fill = "lightgreen") +
xlab("Chromosome") +
ylab("Transcript length (bp)")
```

