# talk04 练习与作业

# 目录

0.1	练习和作业说明	1
0.2	Talk04 内容回顾	1
0.3	练习与作业: 用户验证	1
0.4	练习与作业 1: R session 管理	2
0.5	练习与作业 2: Factor 基础	3
0.6	练习与作业 3: 用 mouse genes 数据做图	6

## 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] "wchen"

Sys.getenv("HOME")

## [1] "/Users/wchen"
```

### 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;
# 定义一些变量(比如 x, y, z 并赋值; 内容随意
rm(list=ls())
x <- c("single", "married", "single");
y <- c(10,100,1000, 10000);
Z <- LETTERS[1:12];

# 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)
w=read.table(file="data/Table0.txt")
```

```
# 保存 workspace 到.RData
save.image(file = "data/TableO.RData");

# 列出当前工作空间内的所有变量
ls();

## [1] "w" "x" "y" "Z"

# 删除当前工作空间内所有变量
rm(list=ls());

# 从.RData 文件恢复保存的数据
load(file = "data/TableO.RData");

# 再次列出当前工作空间内的所有变量
ls();

## character(0)

0.5 练习与作业 2: Factor 基础
```

#### 0.5.1 factor 增加

• 创建一个变量:

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

- 为其增加两个 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 <- as.factor(x);
levels(x) <- c("single", "married");
#x[3] <- "widowed";
# 不行,因为 x 为 factor,只允许接受 single 和 married
levels(x) <- c(levels(x), "widowed");
x[ length(x) + 1 ] <- "widowed";
x[3] <- "widowed";
x;
```

## [1] married single widowed married widowed

## Levels: single married widowed

#### 0.5.2 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

#### 0.5.3 forcats 的问题

```
forcats 包中的 fct_inorder, fct_infreq 和 fct_inseq 函数的作用是什么?
```

This family of functions changes only the order of the levels

fct\_inorder: by the order in which they first appear.

 ${\it fct\_infreq}$  : by number of observations with each level ( large first)

fct\_inseq: by numeric value of level

请使用 forcats 包中的 gss\_cat 数据举例说明

```
## 代码写这里,并运行;
if (!require("forcats")){
   chooseCRANmirror();
   install.packages("forcats",destdir = "D:/resourse/software/Rproject4.1.1/download pace)
}
## Loading required package: forcats
library("forcats");
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
                                                1
                   None
```

```
## 6
            Protestant Southern baptist
                                            NA
attach(gss_cat);
head(fct_inorder(marital), n=20)
   [1] Never married Divorced
                                   Widowed
                                                Never married Divorced
   [6] Married
                     Never married Divorced
                                                              Married
##
                                                Married
## [11] Married
                     Married
                                  Married
                                                              Divorced
                                                Married
## [16] Married
                     Widowed
                                   Never married Married
                                                              Married
## Levels: Never married Divorced Widowed Married Separated No answer
head(fct_infreq(rincome), n=30)
   [1] $8000 to 9999 $8000 to 9999 Not applicable Not applicable Not applicable
   [6] $20000 - 24999 $25000 or more $7000 to 7999 $25000 or more $25000 or more
##
## [11] $25000 or more $25000 or more $25000 or more $25000 or more
## [16] $25000 or more Not applicable $25000 or more $10000 - 14999 Not applicable
## [21] $25000 or more Refused
                                     Not applicable $25000 or more Not applicable
## [26] Not applicable Not applicable Not applicable Not applicable
## 16 Levels: $25000 or more Not applicable $20000 - 24999 ... No answer
f<-factor(1:6,levels=c("1 ","2","3","4","5","6"))
fct_inseq(f)
## [1] <NA> 2
                3
                          5
                               6
## Levels: 1 2 3 4 5 6
     练习与作业 3: 用 mouse genes 数据做图
```

#### 0.6.1 画图

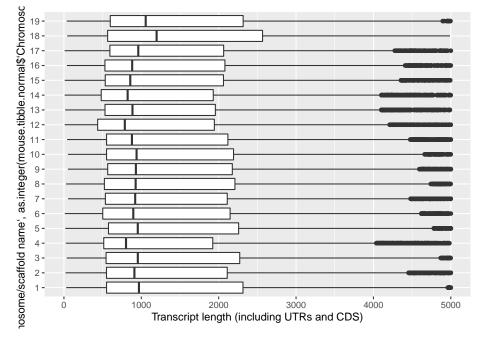
1. 用 readr 包中的函数读取 mouse genes 文件 (从本课程的 Github 页面 下载 data/talk04/)

##

```
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;
if (!require("dplyr")){
 chooseCRANmirror();
 install.packages("dplyr",destdir = "D:/resourse/software/Rproject4.1.1/download packages")
}
## Loading required package: dplyr
##
## Attaching package: 'dplyr'
## The following objects are masked from 'package:stats':
##
##
      filter, lag
## The following objects are masked from 'package:base':
##
##
      intersect, setdiff, setequal, union
library("dplyr");
#devtools::install_github("tidyverse/dplyr")
library(readr)
mouse.tibble<-read_delim(file="../data/talk04/mouse_genes_biomart_sep2018.txt",delim="\
## Rows: 138532 Columns: 6
## -- Column specification -----
## Delimiter: "\t"
## chr (5): Gene stable ID, Transcript stable ID, Protein stable ID, Transcript...
## dbl (1): Transcript length (including UTRs and CDS)
```

```
## i Use `spec()` to retrieve the full column specification for this data.
## i Specify the column types or set `show_col_types = FALSE` to quiet this message.
head(mouse.tibble);
## # A tibble: 6 x 6
     `Gene stable ID`
                       `Transcript stabl~ `Protein stable ~ `Transcript length (in~
##
##
     <chr>
                       <chr>
                                           <chr>
                                                                               <dbl>
## 1 ENSMUSG000000643~ ENSMUST00000082423 <NA>
                                                                                  67
## 2 ENSMUSG000000643~ ENSMUST00000082422 <NA>
                                                                                  67
## 3 ENSMUSG000000643~ ENSMUST00000082421 ENSMUSP000000810~
                                                                                 1144
## 4 ENSMUSG000000643~ ENSMUST00000082420 <NA>
                                                                                  69
## 5 ENSMUSG000000643~ ENSMUST00000082419 ENSMUSP000000810~
                                                                                 519
## 6 ENSMUSG000000643~ ENSMUST00000082418 ENSMUSP000000810~
                                                                                 1824
## # ... with 2 more variables: Transcript type <chr>,
       Chromosome/scaffold name <chr>
colnames(mouse.tibble);
## [1] "Gene stable ID"
## [2] "Transcript stable ID"
## [3] "Protein stable ID"
## [4] "Transcript length (including UTRs and CDS)"
## [5] "Transcript type"
## [6] "Chromosome/scaffold name"
mouse.tibble.normal <-mouse.tibble %>%
  filter( `Chromosome/scaffold name` %in% c( 1:19))
mouse.tibble.xy<-mouse.tibble%>%
  filter( `Chromosome/scaffold name` %in% c('X','Y'))
mouse.tibble.20<-
  bind_rows(mouse.tibble.normal,mouse.tibble.xy)
library(ggplot2)
plot1 <-
  ggplot( data = mouse.tibble.normal,
```

## Warning: Removed 6377 rows containing non-finite values (stat\_boxplot).



```
ylim(0, 5000);
plot2;
```

## Warning: Removed 6639 rows containing non-finite values (stat\_boxplot).

