

Worksheet 7a

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2022-12-9

R Markdown

Worksheet for R Programming

Instructions:

- Use RStudio or the RStudio Cloud accomplish this worksheet.
- Save the R script as RWorksheet_lastname#7a.R.
- On your own GitHub repository, push the R script, the Rmd file, as well as this pdf worksheet to the repo you have created before.
- Do not forget to comment your Git repo on our VLE
- Accomplish this worksheet by answering the questions being asked and writing the code manually.

Basic Statistics

```
library(Hmisc)
```

```
## Warning: package 'Hmisc' was built under R version 4.2.2
```

```
## Loading required package: lattice
```

```
## Loading required package: survival
```

```
## Loading required package: Formula
```

```
## Loading required package: ggplot2
```

```
## Warning: package 'ggplot2' was built under R version 4.2.2
```

```
##
```

```
## Attaching package: 'Hmisc'
```

```
## The following objects are masked from 'package:base':
```

```
##
```

```
## format.pval, units
```

```
library(pastecs)
```

```
## Warning: package 'pastecs' was built under R version 4.2.2
```

1. Create a data frame for the table below.

```
Students <- c(1:10)
PreTest <- c(55,54,47,57,51,61,57,54,63,58)
PostTest <- c(61,60,56,63,56,63,59,56,62,61)

one <- data.frame(Students,PreTest,PostTest)
one
```

```
##      Students PreTest PostTest
## 1          1      55       61
## 2          2      54       60
## 3          3      47       56
## 4          4      57       63
## 5          5      51       56
## 6          6      61       63
## 7          7      57       59
## 8          8      54       56
## 9          9      63       62
## 10         10      58       61
```

- a. Compute the descriptive statistics using different packages (Hmisc and pastecs). Write the codes and its result.

```
#Hmisc
describe(one)
```

```
## one
##
## 3 Variables      10 Observations
## -----
## Students
##      n missing distinct    Info    Mean    Gmd    .05    .10
##     10      0        10      1    5.5    3.667    1.45    1.90
##     .25    .50    .75    .90    .95
##     3.25    5.50    7.75    9.10    9.55
##
## lowest : 1 2 3 4 5, highest: 6 7 8 9 10
##
## Value      1  2  3  4  5  6  7  8  9 10
## Frequency  1  1  1  1  1  1  1  1  1  1
## Proportion 0.1 0.1 0.1 0.1 0.1 0.1 0.1 0.1 0.1 0.1
## -----
## PreTest
##      n missing distinct    Info    Mean    Gmd
##     10      0         8    0.988    55.7    5.444
##
```

```
## lowest : 47 51 54 55 57, highest: 55 57 58 61 63
##
## Value      47  51  54  55  57  58  61  63
## Frequency   1   1   2   1   2   1   1   1
## Proportion 0.1 0.1 0.2 0.1 0.2 0.1 0.1 0.1
## -----
## PostTest
##      n missing distinct      Info      Mean      Gmd
##     10         0         6     0.964     59.7     3.311
##
## lowest : 56 59 60 61 62, highest: 59 60 61 62 63
##
## Value      56  59  60  61  62  63
## Frequency   3   1   1   2   1   2
## Proportion 0.3 0.1 0.1 0.2 0.1 0.2
## -----
```

```
#pasteccs
stat.desc(one)
```

```
##           Students      PreTest      PostTest
## nbr.val    10.0000000  10.0000000  10.0000000
## nbr.null    0.0000000   0.0000000   0.0000000
## nbr.na      0.0000000   0.0000000   0.0000000
## min         1.0000000  47.0000000  56.0000000
## max        10.0000000  63.0000000  63.0000000
## range       9.0000000  16.0000000   7.0000000
## sum        55.0000000 557.0000000 597.0000000
## median      5.5000000  56.0000000  60.5000000
## mean        5.5000000  55.7000000  59.7000000
## SE.mean     0.9574271   1.46855938  0.89504811
## CI.mean.0.95 2.1658506   3.32211213  2.02473948
## var         9.1666667  21.56666667   8.01111111
## std.dev     3.0276504   4.64399254   2.83039063
## coef.var    0.5504819   0.08337509   0.04741023
```

2. The Department of Agriculture was studying the effects of several levels of a fertilizer on the growth of a plant. For some analyses, it might be useful to convert the fertilizer levels to an ordered factor.

- The data were 10,10,10, 20,20,50,10,20,10,50,20,50,20,10.

a. Write the codes and describe the result.

```
agriculture <- c(10,10,10,20,20,50,10,20,10,50,20,50,20,10)
order_one <- sort(agriculture, decreasing = FALSE)
order_one
```

```
## [1] 10 10 10 10 10 20 20 20 20 20 50 50 50
```

3. Abdul Hassan, president of Floor Coverings Unlimited, has asked you to study the exercise levels undertaken by 10 subjects were “l”, “n”, “n”, “i”, “l”, “l”, “n”, “n”, “i”, “l” ; n=none, l=light, i=intense

a. What is the best way to represent this in R?

```
Subjects <- c("l","n","n","i","l","l","n","n","i","l")
one <- data.frame(Subjects)
one
```

```
##   Subjects
## 1         l
## 2         n
## 3         n
## 4         i
## 5         l
## 6         l
## 7         n
## 8         n
## 9         i
## 10        l
```

4. Sample of 30 tax accountants from all the states and territories of Australia and their individual state of origin is specified by a character vector of state mnemonics as:

```
state <- c("tas", "sa", "qld", "nsw", "nsw", "nt", "wa", "wa", "qld",
           "vic", "nsw", "vic", "qld", "qld", "sa", "tas", "sa", "nt",
           "wa", "vic", "qld", "nsw", "nsw", "wa", "sa", "act", "nsw",
           "vic", "vic", "act")

state
```

```
## [1] "tas" "sa"  "qld" "nsw" "nsw" "nt"  "wa"  "wa"  "qld" "vic" "nsw" "vic"
## [13] "qld" "qld" "sa"  "tas" "sa"  "nt"  "wa"  "vic" "qld" "nsw" "nsw" "wa"
## [25] "sa"  "act" "nsw" "vic" "vic" "act"
```

a. Apply the factor function and factor level. Describe the results.

```
fs <- factor(state)
fs
```

```
## [1] tas sa  qld nsw nsw nt  wa  wa  qld vic nsw vic qld qld sa  tas sa  nt  wa
## [20] vic qld nsw nsw wa  sa  act nsw vic vic act
## Levels: act nsw nt qld sa tas vic wa
```

```
levels(state)
```

```
## NULL
```

5. From #4 - continuation:

- Suppose we have the incomes of the same tax accountants in another vector (in suitably large units of money)

```
income <- c(60, 49, 40, 61, 64, 60, 59, 54,
            62, 69, 70, 42, 56, 61, 61, 61, 58, 51, 48,
            65, 49, 49, 41, 48, 52, 46, 59, 46, 58, 43)
income
```

```
## [1] 60 49 40 61 64 60 59 54 62 69 70 42 56 61 61 61 58 51 48 65 49 49 41 48 52
## [26] 46 59 46 58 43
```

- a. Calculate the sample mean income for each state we can now use the special function `tapply()`:

```
incmeans <- tapply(income, state, mean)
incmeans
```

```
##      act      nsw      nt      qld      sa      tas      vic      wa
## 44.50000 57.33333 55.50000 53.60000 55.00000 60.50000 56.00000 52.25000
```

- b. Copy the results and interpret.

Answer: The Result shows the income of tax accountants in each state.

6. Calculate the standard errors of the state income means (refer again to number 3) Note: After this assignment, the standard errors are calculated by: `incster <- tapply(incomes, statef, stdError)`

- a. What is the standard error? Write the codes.

```
stdError <- function(x) sqrt(var(x)/length(x))

calc_StandardDev <- length(incmeans)
calc1 <- sd(incmeans)
calc2 <- calc1/sqrt(calc_StandardDev)
calc2
```

```
## [1] 1.653911
```

- b. Interpret the result. Answer: By getting the state income means, I divided the `sd()`– the standard deviation to `sqrt()` which has the result of length of `incmeans` –the `tapply`. Through this process the result of standard errors was taken.

7. Use the titanic dataset.

- a. subset the titatic dataset of those who survived and not survived. Show the codes and its result.

```
data("Titanic")
titanic <- data.frame(Titanic)

survived <- subset(titanic, Survived == "Yes")
survived
```

##	Class	Sex	Age	Survived	Freq
## 17	1st	Male	Child	Yes	5
## 18	2nd	Male	Child	Yes	11
## 19	3rd	Male	Child	Yes	13
## 20	Crew	Male	Child	Yes	0
## 21	1st	Female	Child	Yes	1
## 22	2nd	Female	Child	Yes	13
## 23	3rd	Female	Child	Yes	14
## 24	Crew	Female	Child	Yes	0
## 25	1st	Male	Adult	Yes	57
## 26	2nd	Male	Adult	Yes	14
## 27	3rd	Male	Adult	Yes	75
## 28	Crew	Male	Adult	Yes	192
## 29	1st	Female	Adult	Yes	140
## 30	2nd	Female	Adult	Yes	80
## 31	3rd	Female	Adult	Yes	76
## 32	Crew	Female	Adult	Yes	20

```
survived2 <- subset(titanic, Survived == "No")
survived2
```

##	Class	Sex	Age	Survived	Freq
## 1	1st	Male	Child	No	0
## 2	2nd	Male	Child	No	0
## 3	3rd	Male	Child	No	35
## 4	Crew	Male	Child	No	0
## 5	1st	Female	Child	No	0
## 6	2nd	Female	Child	No	0
## 7	3rd	Female	Child	No	17
## 8	Crew	Female	Child	No	0
## 9	1st	Male	Adult	No	118
## 10	2nd	Male	Adult	No	154
## 11	3rd	Male	Adult	No	387
## 12	Crew	Male	Adult	No	670
## 13	1st	Female	Adult	No	4
## 14	2nd	Female	Adult	No	13
## 15	3rd	Female	Adult	No	89
## 16	Crew	Female	Adult	No	3

8. The data sets are about the breast cancer Wisconsin. The samples arrive periodically as Dr. Wolberg reports his clinical cases. The database therefore reflects this chronological grouping of the data. You can create this dataset in Microsoft Excel.

a. describe what is the dataset all about.

Answer: The dataset are all about breast cancer Wisconsin that has been reported by Dr. Wolberg as his clinical case. It reflects the chronological group of data, that shows the ID, other information regarding the breast cancer case. It is also to identify the number of malignant and benign case from the biopsy.

b. Import the data from MS Excel. Copy the codes

```
library(readxl)
```

```
## Warning: package 'readxl' was built under R version 4.2.2
```

```
data <- read_excel("D:\\bbmamon\\Worksheet 7a\\Breast_Cancer.xlsx")
data
```

```
## # A tibble: 49 x 11
##       Id CL. thickne~1 Cell ~2 Cell ~3 Marg.~4 Epith~5 Bare.~6 Bl. C~7 Norma~8
##       <dbl>         <dbl>   <dbl>   <dbl>   <dbl>   <dbl> <chr>     <dbl>   <dbl>
##  1 1000025           5         1         1         1         2 1           3         1
##  2 1002945           5         4         4         5         7 10          3         2
##  3 1015425           3         1         1         1         2 2           3         1
##  4 1016277           6         8         8         1         3 4           3         7
##  5 1017023           4         1         1         3         2 1           3         1
##  6 1017122           8        10        10         8         7 10          9         7
##  7 1018099           1         1         1         1         2 10          3         1
##  8 1018561           2         1         2         1         2 1           3         1
##  9 1033078           2         1         1         1         2 1           1         1
## 10 1033078           4         2         1         1         2 1           2         1
## # ... with 39 more rows, 2 more variables: Mitoses <dbl>, Class <chr>, and
## # abbreviated variable names 1: 'CL. thickness', 2: 'Cell size',
## # 3: 'Cell Shape', 4: 'Marg. Adhesion', 5: 'Epith. C.size',
## # 6: 'Bare. Nuclei', 7: 'Bl. Cromatin', 8: 'Normal nucleoli'
```

- c. Compute the descriptive statistics using different packages. Find the values of: c.1 Standard error of the mean for clump thickness.

```
Clump <- length(data$`CL. thickness`)
Clump_A <- sd(data$`CL. thickness`)
Clump_A2 <- Clump_A/sqrt(data$`CL. thickness`)
Clump_A2
```

```
## [1] 1.2812754 1.2812754 1.6541194 1.1696391 1.4325095 1.0129371 2.8650189
## [8] 2.0258743 2.0258743 1.4325095 2.8650189 2.0258743 1.2812754 2.8650189
## [15] 1.0129371 1.0828754 1.4325095 1.4325095 0.9059985 1.1696391 1.0828754
## [22] 0.9059985 1.6541194 1.0129371 2.8650189 1.2812754 1.6541194 1.2812754
## [29] 2.0258743 2.8650189 1.6541194 2.0258743 0.9059985 2.0258743 1.6541194
## [36] 2.0258743 0.9059985 1.1696391 1.2812754 2.0258743 1.1696391 0.9059985
## [43] 1.1696391 1.2812754 0.9059985 2.8650189 1.6541194 2.8650189 1.4325095
```

- c.2 Coefficient of variability for Marginal Adhesion.

```
co_ef <- sd(data$`Marg. Adhesion`)/mean(data$`Marg. Adhesion`)*100
co_ef
```

```
## [1] 97.67235
```

- c.3 Number of null values of Bare Nuclei.

```
null_value <- sum(data$`Bare. Nuclei` == "NA")  
null_value
```

```
## [1] 2
```

c.4 Mean and standard deviation for Bland Chromatin

```
b1 <- mean(data$`Bl. Cromatin`)  
b1
```

```
## [1] 3.836735
```

```
b12 <- sd(data$`Bl. Cromatin`)  
b12
```

```
## [1] 2.085135
```

c.5 Confidence interval of the mean for Uniformity of Cell Shape Calculate the mean

```
calculate_mean <- mean(data$`Cell Shape`)  
calculate_mean
```

```
## [1] 3.163265
```

```
std_A <- length(data$`Cell Shape`)  
std_A
```

```
## [1] 49
```

```
std_B <- sd(data$`Cell Shape`)  
std_B
```

```
## [1] 2.910806
```

```
std_C <- std_B/sqrt(std_A)  
std_C
```

```
## [1] 0.4158294
```

```
A = 0.05  
cs = std_A - 1  
cs2 = qt(p = A/2, df = cs, lower.tail = F)  
cs2
```

```
## [1] 2.010635
```



```
margin_error <- cs2 * std_C
margin_error
```

```
## [1] 0.836081
```

```
lower_bound <- calculate_mean - margin_error
lower_bound
```

```
## [1] 2.327184
```

```
upper_bound <- calculate_mean + margin_error
upper_bound
```

```
## [1] 3.999346
```

```
print(c(lower_bound, upper_bound))
```

```
## [1] 2.327184 3.999346
```

d. How many attributes?

```
breast_cancer <- attributes(data)
breast_cancer
```

```
## $class
## [1] "tbl_df"      "tbl"        "data.frame"
##
## $row.names
## [1] 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22 23 24 25
## [26] 26 27 28 29 30 31 32 33 34 35 36 37 38 39 40 41 42 43 44 45 46 47 48 49
##
## $names
## [1] "Id"           "CL. thickness" "Cell size"      "Cell Shape"
## [5] "Marg. Adhesion" "Epith. C.size" "Bare. Nuclei"   "Bl. Cromatin"
## [9] "Normal nucleoli" "Mitoses"       "Class"
```

Answer: It has 10 attributes

e. Find the percentage of respondents who are malignant. Interpret the results.

```
percent <- subset(data, Class == "malignant")
percent
```

```
## # A tibble: 18 x 11
##       Id CL. thickne~1 Cell ~2 Cell ~3 Marg.~4 Epith~5 Bare.~6 Bl. C~7 Norma~8
##       <dbl>         <dbl>   <dbl>   <dbl>   <dbl>   <dbl> <chr>     <dbl>   <dbl>
## 1 1017122           8       10      10       8       7 10         9       7
## 2 1041801           5        3       3       3       2 3         4       4
## 3 1044572           8        7       5      10       7 9         5       5
```

```
## 4 1047630      7      4      6      4      6 1      4      3
## 5 1050670     10      7      7      6      4 10      4      1
## 6 1054590      7      3      2     10      5 10      5      4
## 7 1054593     10      5      5      3      6 7      7     10
## 8 1057013      8      4      5      1      2 NA      7      3
## 9 1065726      5      2      3      4      2 7      3      6
## 10 1072179     10      7      7      3      8 5      7      4
## 11 1080185     10     10     10      8      6 1      8      9
## 12 1084584      5      4      4      9      2 10      5      6
## 13 1091262      2      5      3      3      6 7      7      5
## 14 1099510     10      4      3      1      3 3      6      5
## 15 1100524      6     10     10      2      8 10      7      3
## 16 1102573      5      6      5      6     10 1      3      1
## 17 1103608     10     10     10      4      8 1      8     10
## 18 1105257      3      7      7      4      4 9      4      8
## # ... with 2 more variables: Mitoses <dbl>, Class <chr>, and abbreviated
## #   variable names 1: 'CL. thickness', 2: 'Cell size', 3: 'Cell Shape',
## #   4: 'Marg. Adhesion', 5: 'Epith. C.size', 6: 'Bare. Nuclei',
## #   7: 'Bl. Cromatin', 8: 'Normal nucleoli'
```

```
#Getting the percentage
18 / 49 * 100
```

```
## [1] 36.73469
```

Interpret the results.

For the total of 49 respondents, there are 18 patient who has malignant breast cancer and 31 patient who has benign case of breast cancer. Patient with benign case is greater than those patient who has malignant case. In conclusion, there are 37% of respondents who has a malignant case of breast cancer.

9. Export the data abalone to the Microsoft excel file. Copy the codes.

```
library("AppliedPredictiveModeling")
```

```
## Warning: package 'AppliedPredictiveModeling' was built under R version 4.2.2
```

```
data(abalone)
View(abalone)
head(abalone)
```

```
##   Type LongestShell Diameter Height WholeWeight ShuckedWeight VisceraWeight
## 1    M      0.455    0.365  0.095    0.5140      0.2245      0.1010
## 2    M      0.350    0.265  0.090    0.2255      0.0995      0.0485
## 3    F      0.530    0.420  0.135    0.6770      0.2565      0.1415
## 4    M      0.440    0.365  0.125    0.5160      0.2155      0.1140
## 5    I      0.330    0.255  0.080    0.2050      0.0895      0.0395
## 6    I      0.425    0.300  0.095    0.3515      0.1410      0.0775
##   ShellWeight Rings
## 1      0.150    15
## 2      0.070     7
```

```
## 3      0.210      9
## 4      0.155     10
## 5      0.055      7
## 6      0.120      8
```

```
summary(abalone)
```

```
## Type      LongestShell      Diameter      Height      WholeWeight
## F:1307   Min.    :0.075   Min.    :0.0550   Min.    :0.0000   Min.    :0.0020
## I:1342   1st Qu.:0.450   1st Qu.:0.3500   1st Qu.:0.1150   1st Qu.:0.4415
## M:1528   Median :0.545   Median :0.4250   Median :0.1400   Median :0.7995
##          Mean    :0.524   Mean    :0.4079   Mean    :0.1395   Mean    :0.8287
##          3rd Qu.:0.615   3rd Qu.:0.4800   3rd Qu.:0.1650   3rd Qu.:1.1530
##          Max.    :0.815   Max.    :0.6500   Max.    :1.1300   Max.    :2.8255
## ShuckedWeight  VisceraWeight  ShellWeight  Rings
## Min.    :0.0010   Min.    :0.0005   Min.    :0.0015   Min.    : 1.000
## 1st Qu.:0.1860   1st Qu.:0.0935   1st Qu.:0.1300   1st Qu.: 8.000
## Median :0.3360   Median :0.1710   Median :0.2340   Median : 9.000
## Mean    :0.3594   Mean    :0.1806   Mean    :0.2388   Mean    : 9.934
## 3rd Qu.:0.5020   3rd Qu.:0.2530   3rd Qu.:0.3290   3rd Qu.:11.000
## Max.    :1.4880   Max.    :0.7600   Max.    :1.0050   Max.    :29.000
```

```
save(file = "abalone.xlsx")
```

```
## Warning in save(file = "abalone.xlsx"): nothing specified to be save()d
```

```
library(xlsx)
```

```
## Warning: package 'xlsx' was built under R version 4.2.2
```

```
install.packages("xlsx")
```

```
## Warning: package 'xlsx' is in use and will not be installed
```

```
write.xlsx("abalone", "D:\\bbmamon\\Worksheet 7a\\abalone.xlsx")
```