

Work Sheet 7

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1. Create a data frame for the table below

```
Student <- seq(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)
```

```
num1 <- data.frame(Student,PreTest,PostTest)
num1
```

##	Student	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.

```
library(Hmisc)

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

## Loading required package: lattice

## Loading required package: survival

## Warning: package 'survival' was built under R version 4.2.2

## 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

describe(num1)

## num1
##
## 3 Variables      10 Observations
##
-----
---
## Student
##      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
```

```
##
-----
---

stat.desc(num1)

##           Student      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 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.

```
num2 <- c(10,10,10,20,20,50,10,
          20,10,50,20,50,20,10)
num2

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

a. Write the codes and describe the result.

```
num2factor <- factor(num2, ordered = TRUE)
num2factor

## [1] 10 10 10 20 20 50 10 20 10 50 20 50 20 10
## Levels: 10 < 20 < 50
```

3. Abdul Hassan, president of Floor Coverings Unlimited, has asked you to study the exercise levels undertaken by 10 num3 were "l", "n", "n", "i", "l", "l", "n", "n", "i", "l"; n=none, l=light, i=intense

```
num3 <- c("l","n","n","i","l","l","n","n","i","l")
```

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

Factor

```
factor(num3, levels = c("n", "l", "i"))
```

```
## [1] l n n i l l n n i l  
## Levels: n l i
```

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")
```

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

factor function and factor level

```
num4a <- factor(state)  
num4a  
  
## [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
```

The result shows the levels of the states of the given data.

Getting factor level of states

5. From #4 - continuation:

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

```
incomes <- 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)
```

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

```
num5a <- tapply(incomes, state, mean)  
num5a  
  
##      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.

```
num5a  
  
##      act      nsw      nt      qld      sa      tas      vic      wa  
## 44.50000 57.33333 55.50000 53.60000 55.00000 60.50000 56.00000 52.25000
```

The result show the level and means of the income of each states.

6. Calculate the standard errors of the state income means (refer again to number 3)

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

```
num6.n <- length(num5a)
num6.sd <- sd(num5a)
num6.se <- num6.sd/sqrt(num6.n)
num6.se

## [1] 1.653911
```

b. Interpret the result.

This is how I get the state income means by dividing the sd() to sqrt() or length() and that is how I get the standard errors of the state income means and this was the result.

7. Use the titanic dataset.

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

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

Survived

```
survive <- subset(Titanic, Survived == "Yes")
survive
```

```
##   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
```

Not Survived

```
died <- subset(Titanic, Survived == "No")
died
```

```
##      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 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*

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

```
library("readxl")

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

library(tinytex)

num8b <- read_excel("C:\\Users\\neil navaroo\\Documents\\Breast_Cancer.xlsx")
num8b

## # A tibble: 49 × 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>
## 1 1000025           5       1       1       1       2 1         3
## 2 1002945           5       4       4       5       7 10        3
## 3 1015425           3       1       1       1       2 2         3
## 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.

```

num8c1.n <- length(num8b$`CL. thickness`)
num8c1.sd <- sd(num8b$`CL. thickness`)
num8c1.se <- num8c1.sd/sqrt(num8b$`CL. thickness`)
num8c1.se

## [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.

```

sd(num8b$`Marg. Adhesion`) / mean(num8b$`Marg. Adhesion`) * 100

## [1] 97.67235

```

c.3 Number of null values of Bare Nuclei.

```

num8c3 <- subset(num8b, `Bare. Nuclei` == "NA")
num8c3

## # A tibble: 2 × 11
##       Id CL. t...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>
##       <dbl>

```

```
## 1 1.06e6      8      4      5      1      2 NA      7      3
1
## 2 1.10e6      6      6      6      9      6 NA      7      8
1
## # ... with 1 more variable: 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.4 Mean and standard deviation for Bland Chromatin

```
mean(num8b$`Bl. Cromatin`)
```

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

```
sd(num8b$`Bl. Cromatin`)
```

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

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

Calculate the mean

```
num8c5 <- mean(num8b$`Cell Shape`)
num8c5
```

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

Calculate the standard error of the mean

```
numA <- length(num8b$`Cell Shape`)
numB <- sd(num8b$`Cell Shape`)
numC <- numB/sqrt(numA)
numC
```

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

Find the t-score that corresponds to the confidence level

```
numD = 0.05
numE = numA - 1
numF = qt(p=numD/2, df=numE, lower.tail=F)
numF
```

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

Constructing the confidence interval

```
numG <- numF * numC
```

Lower

```
numH <- num8c5 - numG
```

Upper


```
numI <- num8c5 + numG
```

Upper and Lower

```
c(numH,numI)
```

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

d. How many attributes?

```
attributes(num8b)
```

```
## $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"
```

There are 3 and these are the class(3), row.name(49) and col.name/length(11).

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

```
num8e <- subset(num8b, Class == "malignant")
num8e
```

```
## # A tibble: 17 × 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 1102573      5      6      5      6     10 1      3
1
## 16 1103608     10     10     10      4      8 1      8
10
## 17 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`
```

There 17 respondents who are malignant. And there are total of 49 respondent.

Getting the percentage

```
17 / 49 * 100
```

```
## [1] 34.69388
```

There are 34.69388 or 35% of respondents who are malignant.

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)
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
```

```
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
```

Exporting the data abalone to the Microsoft excel file

```
library(xlsx)
```

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

```
write.xlsx("abalone", "C:\\Users\\neil navaroo\\Documents\\abalone.xlsx")
```