BCB744 (BioStatistics): Summative Task 2, 12 April 2023

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Honesty Pledge

This assignment requires that you work as an individual and not share your code, results, or discussion with your peers. Penalties and disciplinary action will apply if you are found cheating.

Instructions

Please note the following instructions. Failing to comply with them in full will result in a loss of marks.

- **QUARTO** -> **HTML** Submit your assessment answers as an .html file compiled from your Quarto document. Produce *fully annotated reports*, including the meta-information at the top (name, date, purpose, etc.). Provide ample commentary explaining the purpose of the various tests/sections as necessary.
- TESTING OF ASSUMPTIONS For all questions, make sure that when formal inferential statistics are required, each is preceded by the appropriate tests for the assumptions, i.e., state the assumptions, state the statistical procedure for testing the assumptions and mention their corresponding H_0 . If a graphical approach is used to test assumptions, explain the principle behind the approach. Explain the findings emerging from the test

of assumptions, and justify your selection of the appropriate inferential test (e.g. t-test, ANOVA, etc.) that you will use.

- STATE HYPOTHESES When inferential statistics are required, please provide the full H₀ and H_A, and conclude the analysis with a statement of which is accepted or rejected.
- GRAPHICAL SUPPORT All descriptive and inferential statistics must be supported by the appropriate figures of the results.
- STATEMENT OF RESULTS Make sure that the textual statement of the final result is written exactly as required for it to be published in a journal article. Please consult a journal if you don't know how.
- **FORMATTING** Pay attention to formatting. Some marks will be allocated to the appearance of the script, including considerations of aspects of the tidiness of the file, the use of the appropriate headings, and adherence to code conventions (e.g. spacing etc.).
- MARK ALLOCATION Please see the Introduction Page for an explanation of the assessment approach that will be applied to these questions.

Submit the .html file wherein you provide answers to Questions 1–7 by no later than 19:00 today. Label the script as follows:

```
BCB744\_<Name>\_<Surname>\_Summative\_Task\_2.html, e.g.
```

BCB744 AJ Smit Summative Task 2.html.

Upload your .html files onto Google Forms.

Question 1

Chromosomal effects of mercury-contaminated fish consumption

These data reside in package **coin**, dataset **mercuryfish**. The dataframe contains the mercury level in blood, the proportion of cells with abnormalities, and the proportion of cells with chromosome aberrations in consumers of mercury-contaminated fish and a control group. Please see the dataset's help file for more information.

Analyse the dataset and answer the following questions:

- a. Does the presence of methyl-mercury in a diet containing fish result in a higher proportion of cellular abnormalities?
- b. Does the concentration of mercury in the blood influence the proportion of cells with abnormalities, and does this differ between the control and exposed groups?

c. Is there a relationship between the variables abnormal and ccells? This will have to be for the control and exposed groups, noting that an interaction effect *might* be present.

Question 2

Malignant glioma pilot study

Package **coin**, dataset **glioma**: A non-randomized pilot study on malignant glioma patients with pretargeted adjuvant radioimmunotherapy using yttrium-90-biotin.

- a. Do sex and group interact to affect survival time (time)?
- b. Do age and histology interact to affect survival time (time)?
- c. Show a full graphical exploration of the data. Are there any other remaining patterns visible in the data that should be explored statistically? Study your results, select the most promising and insightful question that remains, and do the analysis.

Question 3

Risk factors associated with low infant birth weight

Package MASS, dataset birthwt: A dataset about the risk factors associated with low infant birth mass collected at Baystate Medical Center, Springfield, Mass. during 1986.

State three hypotheses and test them. Make sure one of the tests makes use of the 95% confidence interval approach rather than a formal inferential methodology.

Question 4

The LungCapData.csv data

- a. Using the Lung Capacity data provided, please calculate the 95% CIs for the LungCap variable as a function of:
 - Gender
 - Smoke
 - Caesarean

```
lungs <- read.csv("../data/LungCapData.csv", sep = "\t")</pre>
library(rcompanion)
(gender_ci <- groupwiseMean(LungCap ~ Gender, data = lungs, conf = 0.95, digits = 3))
  Gender
           n Mean Conf.level Trad.lower Trad.upper
1 female 358 7.41
                         0.95
                                    7.14
                                                7.67
   male 367 8.31
                         0.95
                                    8.03
                                                8.58
(smoke_ci <- groupwiseMean(LungCap ~ Smoke, data = lungs, conf = 0.95, digits = 3))
  Smoke
          n Mean Conf.level Trad.lower Trad.upper
     no 648 7.77
                        0.95
                                   7.56
                                               7.98
1
    yes 77 8.65
                       0.95
                                   8.22
                                               9.07
(caesarean_ci <- groupwiseMean(LungCap ~ Caesarean, data = lungs, conf = 0.95, digits = 3))
              n Mean Conf.level Trad.lower Trad.upper
1
         no 561 7.83
                            0.95
                                       7.61
                                                   8.05
        yes 164 7.97
                                                   8.38
2
                            0.95
                                       7.56
```

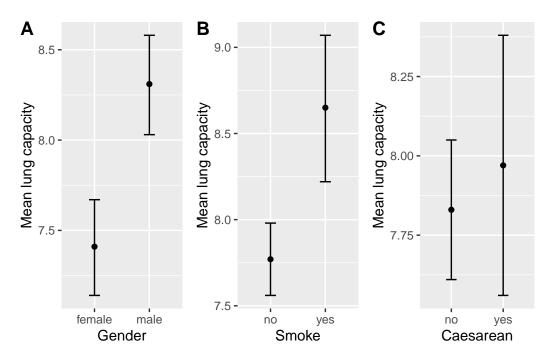
b. Create a graph of the mean \pm 95% CIs and determine if there are statistical differences in LungCap between the levels of Gender, Smoke, and Caesarean. Do the same using inferential statistics. Are your findings the same using these two approaches?

```
plt1 <- ggplot(gender_ci, aes(x = Gender, y = Mean)) +
    geom_point() +
    geom_errorbar(aes(ymin = Trad.lower, ymax = Trad.upper), width = 0.2) +
    ylab("Mean lung capacity")

plt2 <- ggplot(smoke_ci, aes(x = Smoke, y = Mean)) +
    geom_point() +
    geom_errorbar(aes(ymin = Trad.lower, ymax = Trad.upper), width = 0.2) +
    ylab("Mean lung capacity")

plt3 <- ggplot(caesarean_ci, aes(x = Caesarean, y = Mean)) +
    geom_point() +
    geom_errorbar(aes(ymin = Trad.lower, ymax = Trad.upper), width = 0.2) +
    ylab("Mean lung capacity")</pre>
```

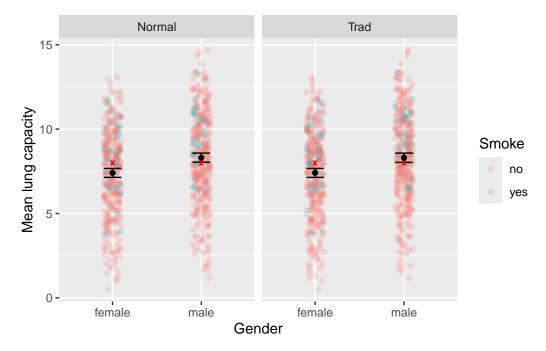
ggarrange(plt1, plt2, plt3, ncol = 3, labels = "AUTO")



c. Produce all the associated tests for assumptions—i.e. the assumptions to be met when deciding whether to use your choice of inferential test or its non-parametric counterpart.

```
lungs %>%
  group_by(Smoke) %>%
  summarise(LungCap var = round(two assum(LungCap)[1], 3),
            LungCap_norm = round(two_assum(LungCap)[2], 3))
# A tibble: 2 x 3
  Smoke LungCap_var LungCap_norm
              <dbl>
  <chr>
                            <dbl>
               7.43
                            0.008
1 no
2 yes
               3.54
                            0.622
lungs %>%
  group_by(Caesarean) %>%
  summarise(LungCap_var = round(two_assum(LungCap)[1], 3),
            LungCap norm = round(two assum(LungCap)[2], 3))
# A tibble: 2 x 3
  Caesarean LungCap_var LungCap_norm
  <chr>
                  <dbl>
                                <dbl>
                                0.004
1 no
                   7.13
                   6.97
                                0.554
2 yes
# It would be best to continue with a Wilcoxon test
```

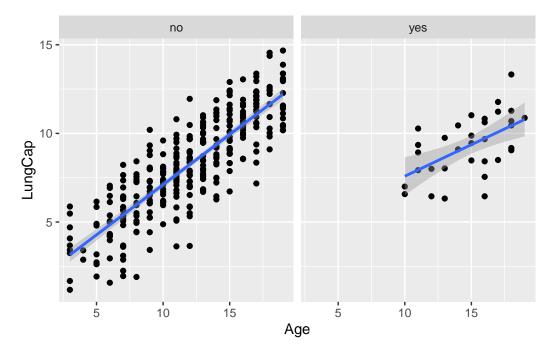
d. Create a combined tidy dataframe (observe tidy principles) with the estimates for the 95% CI for the LungCap data (LungCap as a function of Gender), estimated using both the traditional and bootstrapping approaches. Create a plot comprising two panels (one for the traditional estimates, one for the bootstrapped estimates) of the mean, median, scatter of raw data points, and the upper and lower 95% CI.



e. Undertake a statistical analysis that incorporates both the effect of Age and one of the categorical variables on LungCap. What new insight does this provide?

```
# above we see that within males, the subgroups based on whether or not
# they smoke are normally distributed in both instances
mod1 <- lm(LungCap ~ Smoke * Age, data = lungs[lungs$Gender == "male", ])</pre>
summary(mod1)
Call:
lm(formula = LungCap ~ Smoke * Age, data = lungs[lungs$Gender ==
    "male", ])
Residuals:
    Min
            1Q Median
                            3Q
                                   Max
-4.5903 -0.9875 0.0920 1.0286 3.7097
Coefficients:
            Estimate Std. Error t value Pr(>|t|)
(Intercept) 1.44681 0.25191 5.743 1.96e-08 ***
Smokeyes
              2.57844
                        1.48497
                                 1.736
                                          0.0833 .
              0.56613
                        0.01996 28.367 < 2e-16 ***
Age
                        0.09924 -2.118 0.0348 *
Smokeyes: Age -0.21021
Signif. codes: 0 '*** 0.001 '** 0.01 '* 0.05 '.' 0.1 ' ' 1
Residual standard error: 1.484 on 363 degrees of freedom
Multiple R-squared: 0.6968,
                               Adjusted R-squared: 0.6943
F-statistic: 278.1 on 3 and 363 DF, p-value: < 2.2e-16
# lung capacity of males is affected by age (disregarding effect of smoke),
# lung capacity is not affected by smoke (disregarding effect of age), but
# there is a significant interaction between them, i.e. the effect of age
# is more pronounced in non-smoker than it is in smokers...
lungs[lungs$Gender == "male", ] |>
  ggplot(aes(x = Age, y = LungCap)) +
   geom_point() +
    geom smooth(method = "lm") +
```

facet_wrap(~Smoke)



the figure shows the interaction effect: in non-smokers their lung capacity
increases more rapidly with age, whereas in smokers, the development of lung
capacity with age seems to be stunted.

Question 5

The air quality data

Package datasets, dataset airquality. These are daily air quality measurements in New York, May to September 1973. See the help file for details.

a. Which two of the four response variables are best correlated with each other?

Question 6

The shells.csv data

This dataset contains measurements of shell widths and lengths of the left and right valves of two species of mussels, *Aulacomya* sp. and *Choromytilus* sp. Length and width measurements are presented in mm.

Fully analyse this dataset.

Question 7

The fertiliser_crop_data.csv data

The data represent an experiment designed to test whether or not fertiliser type and the density of planting have an effect on the yield of wheat. The dataset contains the following variables:

- Final yield (kg per acre)—make sure to convert this to the most suitable SI unit before continuing with your analysis
- Type of fertiliser (fertiliser type A, B, or C)
- Planting density (1 = low density, 2 = high density)
- Block in the field (north, east, south, west)

SD = sd(mass),

Fully analyse this dataset.

```
fert <- read.csv("../data/fertiliser_crop_data.csv")</pre>
# convert to SI units
fert <- fert |>
  mutate(mass = mass / 0.40468564224)
\# are assumptions met? note that I also calculate the mean +/- SD here
fert %>%
  group_by(density) %>%
  summarise(mean = mean(mass),
            SD = sd(mass),
            mass_var = round(two_assum(mass)[1], 3),
            mass_norm = round(two_assum(mass)[2], 3))
# A tibble: 2 x 5
  density
            mean
                    SD mass_var mass_norm
          <dbl> <dbl>
    <int>
                           <dbl>
                                     <dbl>
        1 11889. 40.8
                           1668.
1
                                     0.469
2
        2 11920. 43.3
                           1877.
                                     0.529
fert %>%
  group_by(block) %>%
  summarise(mean = mean(mass),
```

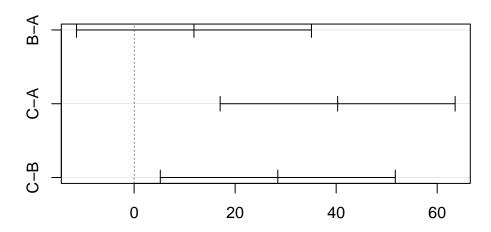
mass_var = round(two_assum(mass)[1], 3),
mass_norm = round(two_assum(mass)[2], 3))

```
# A tibble: 4 x 5
  block mean
                 SD mass_var mass_norm
  <chr> <dbl> <dbl>
                      <dbl>
                                <dbl>
1 east 11925. 43.4
                       1882.
                                0.422
2 north 11894. 42.2 1781.
                                0.77
3 south 11884. 39.7 1578.
                                0.212
4 west 11915. 43.7 1906.
                                0.21
fert %>%
  group_by(fertilizer) %>%
  summarise(mean = mean(mass),
           SD = sd(mass),
           mass_var = round(two_assum(mass)[1], 3),
           mass_norm = round(two_assum(mass)[2], 3))
# A tibble: 3 x 5
  fertilizer mean
                      SD mass_var mass_norm
  <chr>
            <dbl> <dbl> <dbl>
                                     <dbl>
1 A
           11887. 46.1
                         2122.
                                     0.774
2 B
            11899. 38.6 1490.
                                     0.887
            11927. 40.3 1623.
3 C
                                     0.254
# yes, all assumptions check out, proceed with normal paramatric stats
# do an ANOVA and look at main effects first
aov1 <- aov(mass ~ density + block + fertilizer, data = fert)</pre>
summary(aov1)
           Df Sum Sq Mean Sq F value
                                      Pr(>F)
            1 23164 23164 15.224 0.000184 ***
density
                2199
                       1099
                             0.723 0.488329
block
fertilizer 2 27444 13722
                             9.018 0.000269 ***
Residuals 90 136940
                      1522
Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
# the block effect is not significant but density and fertilizer are
# let's check if the fertilizer type interacts with density
aov2 <- aov(mass ~ density * fertilizer, data = fert)</pre>
summary(aov2)
```

```
Df Sum Sq Mean Sq F value
                                              Pr(>F)
                               23164 15.195 0.000186 ***
density
                    1 23164
fertilizer
                    2 27444
                               13722
                                      9.001 0.000273 ***
density:fertilizer 2 1935
                                 967
                                       0.635 0.532500
                   90 137203
Residuals
                                1524
Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
# no interaction effect is present, so the fertilizer has the same
# effect regardless of at which planting density it is applied
# lets see which planting fertilizer results in the greatest mass
TukeyHSD(aov2, which = "fertilizer", ordered = TRUE)
  Tukey multiple comparisons of means
    95% family-wise confidence level
    factor levels have been ordered
Fit: aov(formula = mass ~ density * fertilizer, data = fert)
$fertilizer
        diff
                    lwr
                            upr
                                     p adj
B-A 11.84752 -11.414312 35.10935 0.4482026
C-A 40.29177 17.029945 63.55360 0.0002393
C-B 28.44426
             5.182428 51.70609 0.0123951
TukeyHSD(aov2, which = "fertilizer", ordered = TRUE)
  Tukey multiple comparisons of means
    95% family-wise confidence level
    factor levels have been ordered
Fit: aov(formula = mass ~ density * fertilizer, data = fert)
$fertilizer
                    lwr
                             upr
                                     p adj
B-A 11.84752 -11.414312 35.10935 0.4482026
C-A 40.29177 17.029945 63.55360 0.0002393
C-B 28.44426 5.182428 51.70609 0.0123951
```

```
plot(TukeyHSD(aov2, which = "fertilizer", ordered = TRUE))
```

95% family-wise confidence level



Differences in mean levels of fertilizer

```
# here we can see that the mass of crop produced by fertilizer C is the
# greatest, significantly more so compared to both A and B; the effect
# of fertilizer B is no different than that of A
#
# the second planting density also yields a greater mass per ha
#
# make sure the results are written up as appropriate for a journal,
# so indicate the d.f., S.S., and p-value
```

The end

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BCB744_<Name>_<Surname>_Summative_Task_2.html, e.g.

BCB744_AJ_Smit_Summative_Task_2.html.

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