

Problem set, group 3

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```
# keep this chunk in all your RMarkdown scripts
```

```
knitr::opts_chunk$set(echo = TRUE)
```

```
knitr::opts_chunk$set(tidy.opts = list(width.cutoff = 60), tidy = TRUE)
```

```
# List required packages
```

```
library(tidyverse)
```

```
## -- Attaching core tidyverse packages ----- tidyverse 2.0.0 --
```

```
## v dplyr      1.1.4      v readr      2.1.5
```

```
## v forcats    1.0.0      v stringr    1.5.1
```

```
## v ggplot2    3.5.1      v tibble     3.2.1
```

```
## v lubridate  1.9.3      v tidyr      1.3.1
```

```
## v purrr      1.0.2
```

```
## -- Conflicts ----- tidyverse_conflicts() --
```

```
## x dplyr::filter() masks stats::filter()
```

```
## x dplyr::lag()     masks stats::lag()
```

```
## i Use the conflicted package (<http://conflicted.r-lib.org/>) to force all conflicts to become errors
```

```
library(openxlsx)
```

```
library(lmtest)
```

```
## Loading required package: zoo
```

```
##
```

```
## Attaching package: 'zoo'
```

```
##
```

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

```
##
```

```
##      as.Date, as.Date.numeric
```

```
## Initial data loading, transforming
```

```
mammals <- read.xlsx("mammals_sleep.xlsx")
```

```
colnames(mammals) <- sub(pattern = "\\.(\\((.*)\\))", replacement = "",  
  colnames(mammals))
```

```
colnames(mammals) <- sub(pattern = "\\((.*)\\)", replacement = "",  
  colnames(mammals))
```

```
mammals[, 3:12][mammals[, 3:12] == -999] <- NA
```

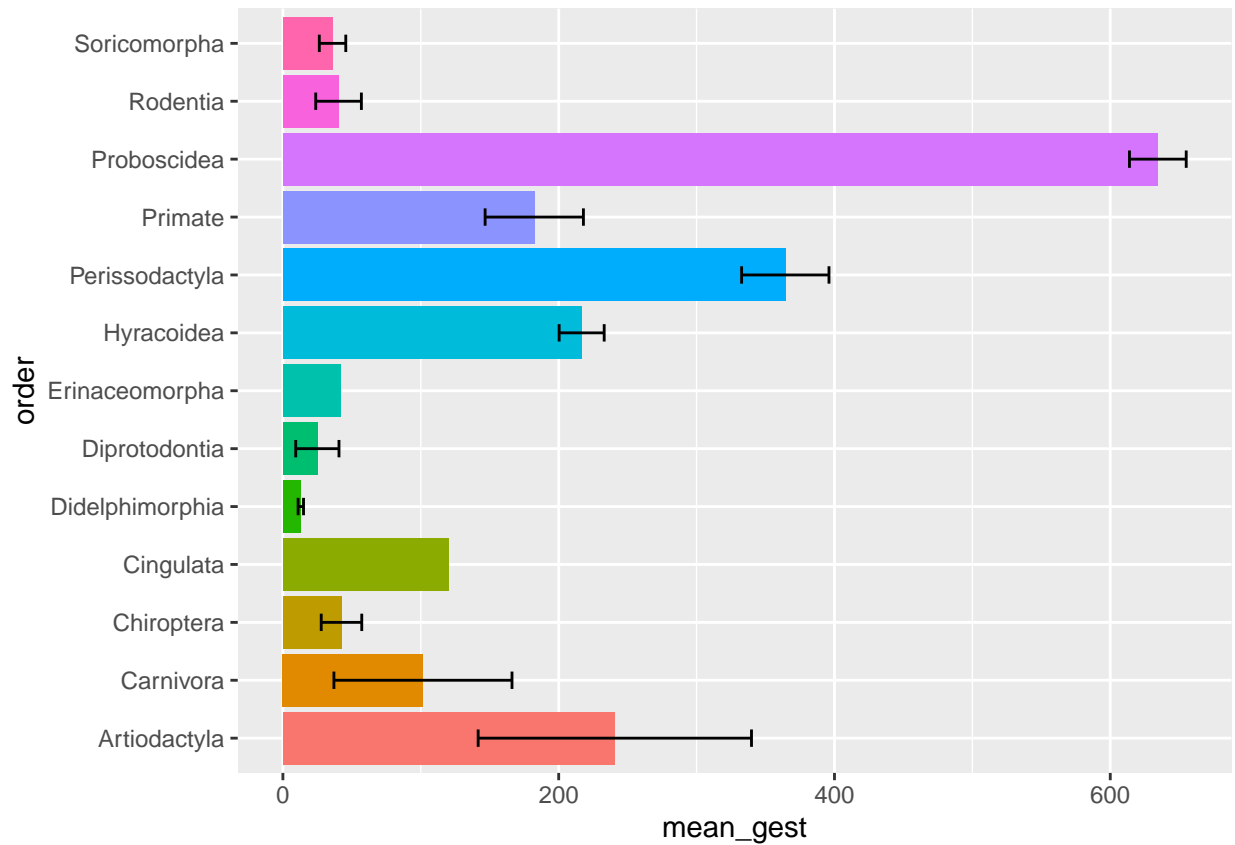
```
## analysis
```

```
summary_table <- mammals %>%
  group_by(order) %>%
  summarize(total = n(), mean_gest = mean(gestation.time, na.rm = TRUE),
    sd_gest = sd(gestation.time, na.rm = TRUE), median_gest = median(gestation.time,
    na.rm = TRUE), iqr = IQR(gestation.time, na.rm = TRUE),
    var = var(gestation.time, na.rm = TRUE), se = sd_gest/sqrt(n()),
    CI_low = mean_gest - 1.96 * se, CI_high = mean_gest +
    1.96 * se)
summary_table
```

```
## # A tibble: 18 x 10
```

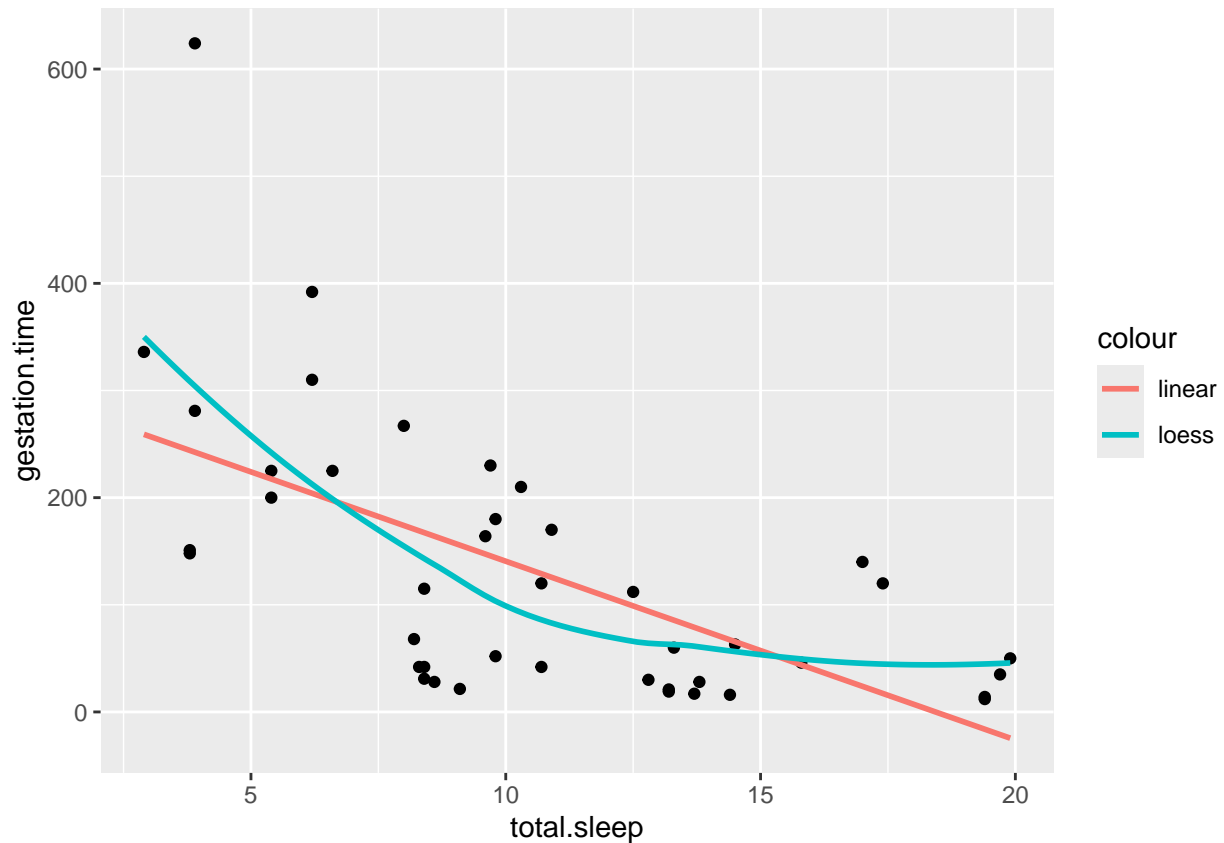
##	order	total	mean_gest	sd_gest	median_gest	iqr	var	se	CI_low	CI_high
##	<chr>	<int>	<dbl>	<dbl>	<dbl>	<dbl>	<dbl>	<dbl>	<dbl>	<dbl>
##	1 Afroso~	1	60	NA	60	0	NA	NA	NA	NA
##	2 Artiod~	7	241.	134.	151	192.	17898.	50.6	142.	340.
##	3 Carniv~	8	102.	93.2	63	20	8682.	32.9	37.0	166.
##	4 Chiro~	2	42.5	10.6	42.5	7.5	112.	7.5	27.8	57.2
##	5 Cingul~	2	120	NA	120	0	NA	NA	NA	NA
##	6 Didelp~	2	13	1.41	13	1	2	1	11.0	15.0
##	7 Diprot~	2	25	11.3	25	8	128	8	9.32	40.7
##	8 Erinac~	2	42	NA	42	0	NA	NA	NA	NA
##	9 Eulipo~	1	21.5	NA	21.5	0	NA	NA	NA	NA
##	10 Hyraco~	3	217.	14.4	225	12.5	208.	8.33	200.	233
##	11 Lagomo~	1	31	NA	31	0	NA	NA	NA	NA
##	12 Monotr~	1	28	NA	28	0	NA	NA	NA	NA
##	13 Periss~	3	364.	28.0	365	28	784.	16.2	333.	396.
##	14 Primate	10	182.	57.5	175	79	3306.	18.2	147.	218.
##	15 Probos~	2	634.	14.8	634.	10.5	220.	10.5	614.	655.
##	16 Rodent~	11	40.4	28.0	30	20.5	785.	8.45	23.8	56.9
##	17 Scande~	1	46	NA	46	0	NA	NA	NA	NA
##	18 Sorico~	3	36	8.49	36	6	72	4.90	26.4	45.6

```
summary_table %>%
  filter(total > 1) %>%
  ggplot(aes(y = order, x = mean_gest, fill = order)) + geom_col(show.legend = FALSE) +
  geom_errorbar(aes(xmin = CI_low, xmax = CI_high), width = 0.3,
    color = "black")
```



```
na.omit(mammals) %>%
  ggplot(aes(x = total.sleep, y = gestation.time)) + geom_point() +
  geom_smooth(aes(color = "linear"), se = FALSE, method = "lm") +
  geom_smooth(aes(color = "loess"), se = FALSE, span = 0.9)
```

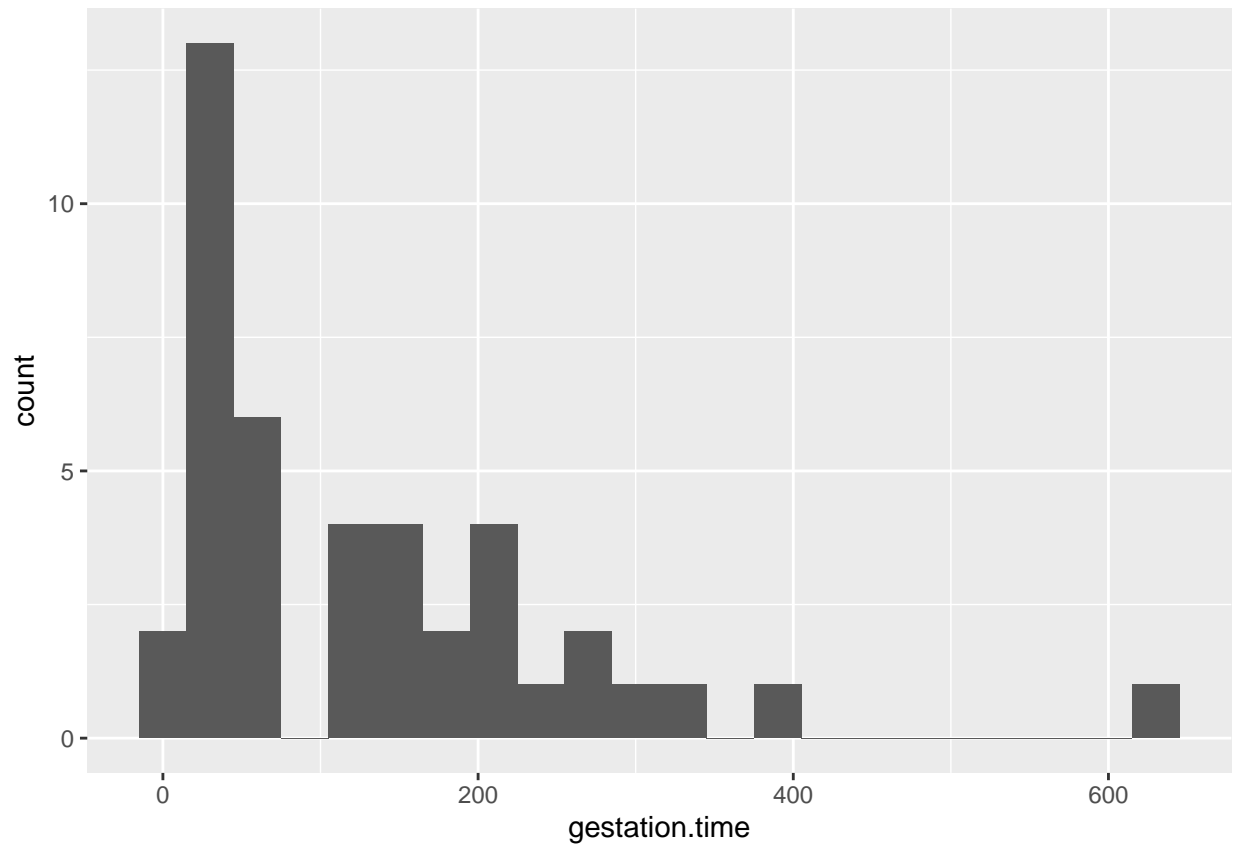
```
## 'geom_smooth()' using formula = 'y ~ x'
## 'geom_smooth()' using method = 'loess' and formula = 'y ~ x'
```



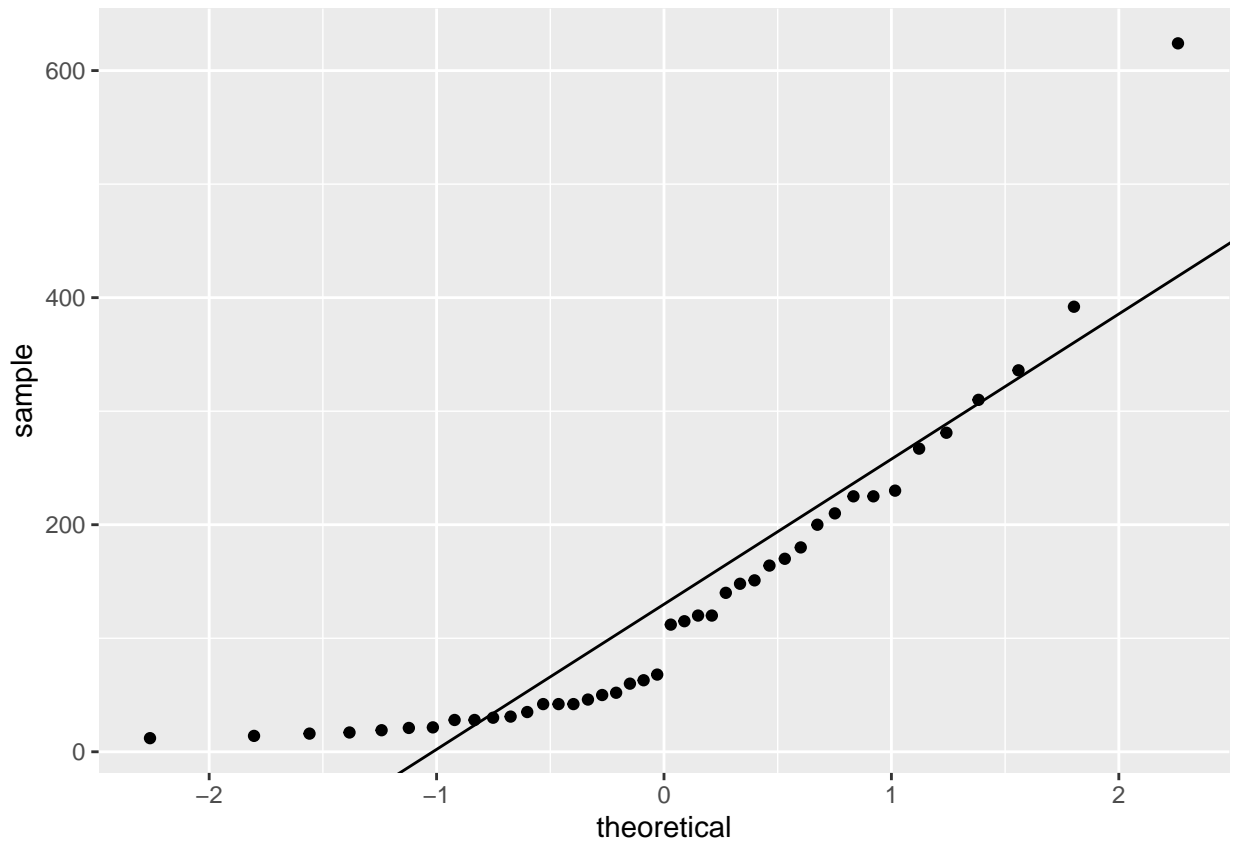
```
cor.test(x = na.omit(mammals)$total.sleep, y = na.omit(mammals)$gestation.time)
```

```
##
## Pearson's product-moment correlation
##
## data: na.omit(mammals)$total.sleep and na.omit(mammals)$gestation.time
## t = -4.926, df = 40, p-value = 1.499e-05
## alternative hypothesis: true correlation is not equal to 0
## 95 percent confidence interval:
## -0.7738778 -0.3818580
## sample estimates:
## cor
## -0.6144743
```

```
na.omit(mammals) %>%
  ggplot(aes(x = gestation.time)) + geom_histogram(binwidth = 30,
  na.rm = TRUE)
```



```
ggplot(na.omit(mammals), aes(sample = gestation.time)) + stat_qq(show.legend = FALSE) +  
  geom_abline(aes(intercept = mean(x = gestation.time), slope = sd(x = gestation.time)))
```



```
test_hom <- lmtest::bptest(na.omit(mammals)$total.sleep ~ na.omit(mammals)$gestation.time)
test_hom
```

```
##
## studentized Breusch-Pagan test
##
## data: na.omit(mammals)$total.sleep ~ na.omit(mammals)$gestation.time
## BP = 0.83182, df = 1, p-value = 0.3617
```

```
## 1. A rationale for your hypothesis, and a clear
## formulation of what you predict if your hypothesis is
## supported.
```

```
## Hypothesis: Gestation time correlates with sleep time
## within the an animal's order. Prediction: Longer
## gestation time will relate to longer sleep time on
## average by order. Null hypothesis: There is no
## relationship between sleep time and gestation time.
```

```
## 2. A description of any potential changes to the dataset
## that were necessary to prepare it for analysis, such as
## recoding of variables, transformations, etc.
```

```
## We had to add a new column containing the order of each
## animal in order to group them.
```

3. A summary of your exploratory analyses, with key
results reported in a standard format. As part of this
section, please include some basic descriptive stats of
your variables, incl groups

4. Results of your main analyses, reported in a standard
format (can include a well formatted table, if more
efficient than writing out all details). As part of this
section, you need to clearly state what variables went
into your analysis in what fashion (dependent,
independent, original, transformed, recoded, whatever),
and you need to report any special parameters in R
functions (beyond the default ones) and justify why you
selected them. This section also includes any post-hoc
analyses that you conducted.

5. A graphical depiction of your results appropriately
labeled and referenced in your writing. Each graph
should have a short, descriptive caption that explains
what is shown in the graph.

6. A brief discussion of the biological significance of
your findings, i.e. an interpretation of your results
based on the outcome of your hypothesis testing.