

# Basic inferential data analysis on the tooth growth data in R

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A note in the beginning:

As stated in the introduction for the assignment, each report should not be longer than 3 pages + 3 pages of figures and code. I therefore went for the goal to not exceed 6 pages including all figures and code.

## Load the ToothGrowth data and perform some basic exploratory data analyses

Load the data and suppress messages and warnings.

```
data(ToothGrowth)
library(ggplot2)
suppressWarnings(suppressMessages(library(plyr)))
library(grid)
```

First step in the exploratory data analysis: get an overview of the dataframe

```
str(ToothGrowth)

## 'data.frame':   60 obs. of  3 variables:
## $ len : num  4.2 11.5 7.3 5.8 6.4 10 11.2 11.2 5.2 7 ...
## $ supp: Factor w/ 2 levels "OJ","VC": 2 2 2 2 2 2 2 2 2 2 ...
## $ dose: num  0.5 0.5 0.5 0.5 0.5 0.5 0.5 0.5 0.5 0.5 ...
```

then (step 2) check how many examples from each supplementation and dose exists

```
table(ToothGrowth$supp, ToothGrowth$dose, dnn=c("supp","dose"))

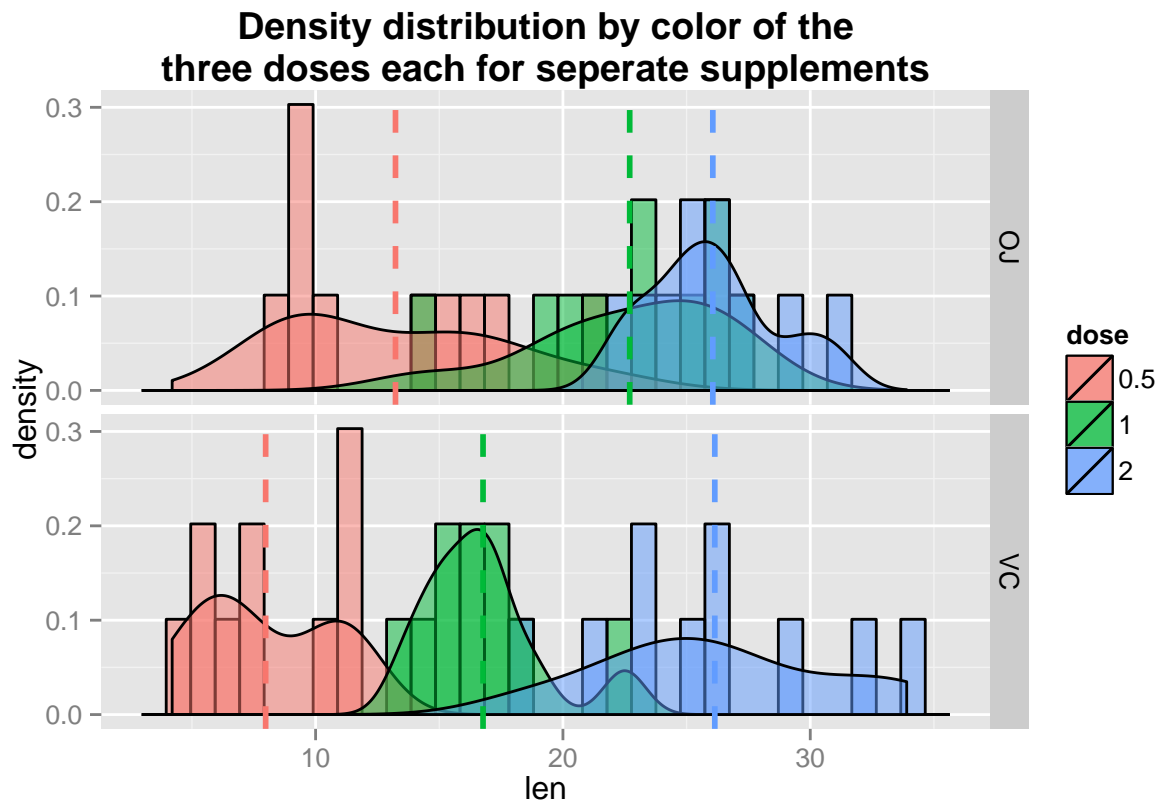
##      dose
## supp 0.5  1  2
##   OJ  10 10 10
##   VC  10 10 10
```

And as a third I step create a plot, that emphasises the differences in tooth growth per dose and supplementation

```
ToothGrowth$dose <- factor(ToothGrowth$dose)
try_cat <- ddply(ToothGrowth,.(dose,supp),summarize,means=mean(len))
q = ggplot(ToothGrowth, aes(x=len,fill=dose)) +
  geom_histogram(alpha=0.5, position='identity', aes(y=..density..),
    bin=(max(ToothGrowth$len)-min(ToothGrowth$len))/30, colour='black') +
  geom_density(alpha=0.5) + facet_grid(supp ~.) +
  geom_vline(data=try_cat, aes(xintercept=means, color = dose), linetype='dashed',
    size=1) +
  ggtitle('Density distribution by color of the
```

```
three doses each for separate supplements') +
  theme(plot.title=element_text(face='bold'))

print(q)
```

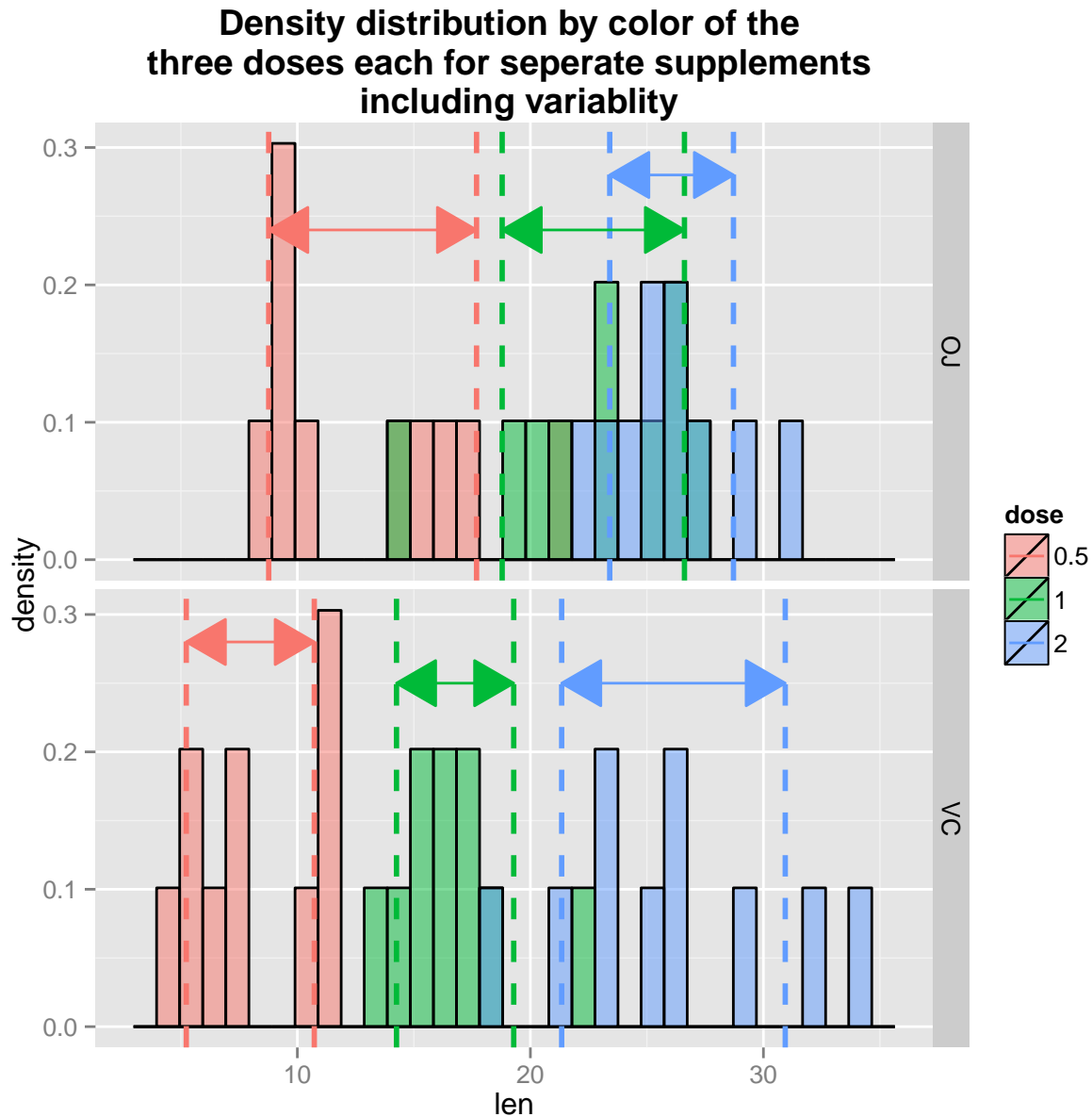


where the dashed lines represent the mean values for each separate distribution. This plot already gives a great overview on the data and how it is structured. The following plot shows the range of  $\mu \pm sd$  to give an overview over the variability of the data.

```
cat_sd <- ddply(ToothGrowth,.(dose,supp),summarize,sds=sd(len))
cat_sd <- merge(cat_sd, try_cat, by=c("dose","supp"))

q_sd = ggplot(data=ToothGrowth, aes(x=len,fill=dose)) +
  geom_histogram(alpha=0.5, position='identity', aes(y=..density..),color='black',
    bin=(max(ToothGrowth$len)-min(ToothGrowth$len))/30) +
  facet_grid(supp ~.) +
  geom_vline(data=cat_sd, aes(xintercept=means+sds, color = dose),
    linetype='dashed', size=1) +
  geom_vline(data=cat_sd, aes(xintercept=means-sds, color = dose),
    linetype='dashed', size=1) +
  geom_segment(data=cat_sd, aes(x=means-sds, xend= means+sds,
    y=c(0.24,0.24,0.28,0.28,0.25,0.25),
    yend=c(0.24,0.24,0.28,0.28,0.25,0.25), color = dose),
    arrow=arrow(ends='both',type='closed'),size=0.5)+
  ggtitle('Density distribution by color of the
three doses each for separate supplements \n including variability') +
```

```
theme(plot.title=element_text(face='bold'))
print(q_sd)
```



These two plots allow me to develop 3 Nullhypotheses

1. 2 doses of the supplement OJ are as good for toothgrowth as 2 doses of the supplement VC.
2. 1 dose of the supplement OJ has a similar effect on toothsgrowths as 2 doses of the supplement VC.
3. 2 doses of OJ have the same effect or less on toothgrowth than all other supplementations and doses, except for 2 doses of the supplement VC.

## Provide a basic summary of the data

In order to properly construct the plots for the exploratory data analysis, I already generated a summary of the data and stored it in the variable `cat_sd`:

```
cat_sd
```

```
##   dose supp      sds means
## 1  0.5   OJ 4.459709 13.23
## 2  0.5   VC 2.746634  7.98
## 3   1    OJ 3.910953 22.70
## 4   1    VC 2.515309 16.77
## 5   2    OJ 2.655058 26.06
## 6   2    VC 4.797731 26.14
```

## Use hypothesis testing to compare different tooth growths by supp and dose.

In the following I create a dataframe containing p values and the t confidence intervals for all combinations of supplementation and doses. The assumptions are

1. Each pair of dose and supplementation has a different variance than the other ones.
2. We use a two sided test, as two of the hypotheses test for equality for the mean value of two distributions.
3. We use a one sided test for the last hypothesis as our Null hypothesis is that all other dose-supplement pairs are as effective as the OC.2 pair or better.

```
suppressWarnings(library(data.table))
final_set <- data.frame(combination=character(),p.value=numeric(),
                        interval.lower=numeric(), interval.upper=numeric())
in_groups <- split(ToothGrowth, list(ToothGrowth$supp, ToothGrowth$dose))
names = names(in_groups)
for (i in 1:length(names)) {
  for (m in 1:length(names)){
    if (i < m) {
      temp <- t.test(in_groups[[i]][[1]], in_groups[[m]][[1]], var.equal = FALSE)
      temp2 <- c("",0,0,0)
      temp2[1] <- paste(names(in_groups)[i], names(in_groups)[m], sep="~")
      temp2[2] <- round(temp$p.value,7)
      temp2[3:4] <- round(temp$conf.int,7)
      final_set <- rbindlist(list(final_set,as.list(temp2)))
    }
  }
}
final_set$contains0 <- final_set$interval.lower <= 0 & final_set$interval.upper >= 0
print(final_set)
```

```
##      combination  p.value interval.lower interval.upper contains0
## 1: OJ.0.5~VC.0.5 0.0063586      1.7190573      8.7809427      FALSE
## 2: OJ.0.5~OJ.1  8.78e-05     -13.4156344     -5.5243656      FALSE
## 3: OJ.0.5~VC.1 0.0460103      -7.008109      -0.071891      FALSE
## 4: OJ.0.5~OJ.2  1.3e-06     -16.3352406     -9.3247594      FALSE
## 5: OJ.0.5~VC.2  7.2e-06     -17.2635219     -8.5564781      FALSE
```

## 6:	VC.0.5~OJ.1	0	-17.9214925	-11.5185075	FALSE
## 7:	VC.0.5~VC.1	7e-07	-11.265712	-6.314288	FALSE
## 8:	VC.0.5~OJ.2	0	-20.6181832	-15.5418168	FALSE
## 9:	VC.0.5~VC.2	0	-21.901512	-14.418488	FALSE
## 10:	OJ.1~VC.1	0.0010384	2.8021482	9.0578518	FALSE
## 11:	OJ.1~OJ.2	0.0391951	-6.5314425	-0.1885575	FALSE
## 12:	OJ.1~VC.2	0.0965261	-7.5643336	0.6843336	TRUE
## 13:	VC.1~OJ.2	2e-07	-11.7203326	-6.8596674	FALSE
## 14:	VC.1~VC.2	9.16e-05	-13.0542667	-5.6857333	FALSE
## 15:	OJ.2~VC.2	0.9638516	-3.7980705	3.6380705	TRUE

## Discussion and conclusion

Going back to our Null-hypotheses:

1. 2 doses of the supplement OJ are as good for toothgrowth as 2 doses of the supplement VC.
2. 1 dose of the supplement OJ has a similar effect on toothgrowths as 2 doses of the supplement VC.
3. 2 doses of OJ have the same effect or less on toothgrowth than all other supplementations and doses, except for 2 doses of the supplement VC.

and the corresponding  $H_A$ 's are:

1. 2 doses of the supplement OJ(OC.2) have a different effect on toothgrowth as 2 doses of the supplement VC (VC.2).
2. 1 dose of the supplement OJ (OC.1) has a different effect on toothgrowth as 2 doses of VC (VC.2).
3. 2 doses of OJ (OC.2) have a greater effect on toothgrowth than all other supplementation, except for 2 doses of VC (VC.2).

Checking the first Null Hypothesis:

##	combination	p.value	interval.lower	interval.upper	contains0
## 1:	OJ.2~VC.2	0.9638516	-3.7980705	3.6380705	TRUE

The upper subset makes it clear that we cannot reject the Null-Hypothesis and that the probability of the means being equal is 96%. The t-confidence interval backs this conclusion as it contains 0 and is actually centered around 0. In 95% of the time this interval contains the true difference of the means.

Checking the second Null Hypothesis:

##	combination	p.value	interval.lower	interval.upper	contains0
## 1:	OJ.1~VC.2	0.0965261	-7.5643336	0.6843336	TRUE

This result is way less clear, but we cannot reject the Null-Hypothesis either. 9% of data can be explained by the null hypothesis, which is not enough to reject it, but not a lot either. The confidence interval contains 0, but is not centered around it, with a tendency towards negative values. This means, that the mean more often than not, is smaller than 0. Which can be interpreted as toothgrowth is more often bigger under VC.2 than under OJ.1, but this difference is not big enough to be significant.

Checking the third Null Hypothesis: I have to redo the t-test as a one sided test:

```

final_set2 <- data.frame(combination=character(),p.value=numeric(),
                        interval.lower=numeric(), interval.upper=numeric())
for (m in 1:length(names(in_groups))){
  if (names[m] != "OJ.2") {
    temp <- t.test(in_groups$OJ.2[[1]], in_groups[[m]][[1]], var.equal = FALSE,
                  alternative="greater")
    temp2[1] <- paste("OJ.2", names(in_groups)[m], sep="~")
    temp2[2] <- round(temp$p.value,7)
    temp2[3:4] <- round(temp$conf.int,7)
    final_set2 <- rbindlist(list(final_set2,as.list(temp2)))
  }
}
print(final_set2)

```

```

##      combination    p.value interval.lower interval.upper
## 1: OJ.2~OJ.0.5      7e-07      9.9484504          Inf
## 2: OJ.2~VC.0.5        0      15.985071          Inf
## 3:  OJ.2~OJ.1 0.0195976      0.7486236          Inf
## 4:  OJ.2~VC.1      1e-07      7.2841492          Inf
## 5:  OJ.2~VC.2 0.5180742     -3.1334996          Inf

```

I set the alternative to 'greater' as our hypothesis states OJ.2 has a greater impact, than the other supplementations and doses, except for VC.2.

Going through the p-values we can safely reject all the Nullhypothesis (all doses and supplementations have the same or a greater effect than OJ.2) and conclude that OJ.2 has a greater effect on toothgrowth than the others (except VC.2). The significance level is always lower than 5%.