

In-class worksheet 2

Jan 24, 2019

1. t test

We will try the t test on the built-in data set `PlantGrowth`. However, first we need to reformat the data set, which we do with the function `unstack()`. We store the reformatted data set in a variable `plants`:

```
head(PlantGrowth)
```

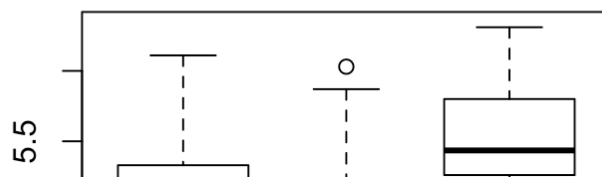
```
##   weight group
## 1   4.17  ctrl
## 2   5.58  ctrl
## 3   5.18  ctrl
## 4   6.11  ctrl
## 5   4.50  ctrl
## 6   4.61  ctrl
```

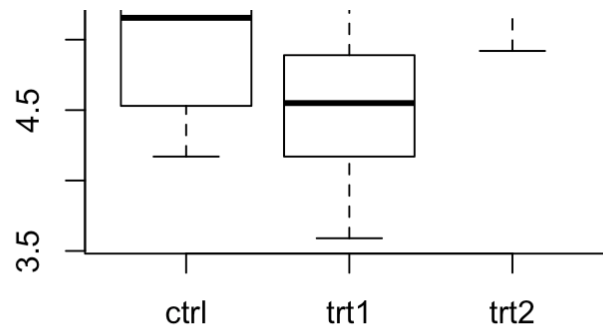
```
plants <- unstack(PlantGrowth)
head(plants)
```

```
##   ctrl trt1 trt2
## 1 4.17 4.81 6.31
## 2 5.58 4.17 5.12
## 3 5.18 4.41 5.54
## 4 6.11 3.59 5.50
## 5 4.50 5.87 5.37
## 6 4.61 3.83 5.29
```

The data set contains plant growth yield (dry weight) under one control and two treatment conditions:

```
boxplot(plants)
```





Question: Is the mean control weight significantly different from the mean weight under treatment 1? Is the mean weight under treatment 1 significantly different from the mean weight under treatment 2? Use the function `t.test()` to find out.

First, control vs. treatment 1:

```
t.test(plants$ctrl, plants$trt1)
```

```
##
## Welch Two Sample t-test
##
## data:  plants$ctrl and plants$trt1
## t = 1.1913, df = 16.524, p-value = 0.2504
## alternative hypothesis: true difference in means is not equal to 0
## 95 percent confidence interval:
## -0.2875162  1.0295162
## sample estimates:
## mean of x mean of y
##      5.032      4.661
```

The p-value is 0.25. We cannot reject H_0 . Control and treatment 1 do not appear to be different.

Second, treatment 1 vs. treatment 2:

```
t.test(plants$trt1, plants$trt2)
```

```
##  
## Welch Two Sample t-test  
##  
## data: plants$trt1 and plants$trt2  
## t = -3.0101, df = 14.104, p-value = 0.009298  
## alternative hypothesis: true difference in means is not equal to 0  
## 95 percent confidence interval:  
## -1.4809144 -0.2490856  
## sample estimates:  
## mean of x mean of y  
## 4.661 5.526
```

The p-value is 0.009. We reject H_0 . Plants seem to grow more under treatment 2 than under treatment 1.

2. Correlation

We will try the correlation test on the built-in data set `cars`. The data set contains the speed of cars and the distances taken to stop, measured in the 1920s:

```
head(cars)
```

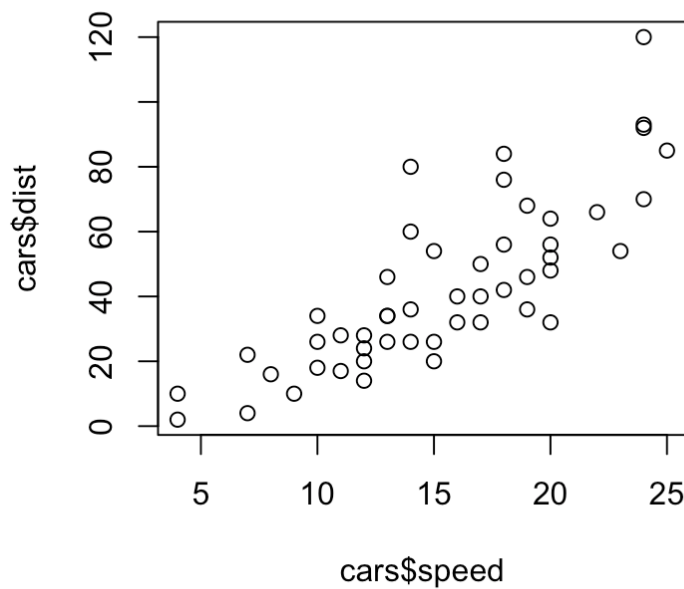
```
## speed dist  
## 1 4 2  
## 2 4 10  
## 3 7 4  
## 4 7 22  
## 5 8 16  
## 6 9 10
```

Is there a relationship between speed and stopping distance? Use the function `cor.test()` to find out. Then make a scatterplot of speed vs. stopping distance, using the function `plot()`.

```
cor.test(cars$speed, cars$dist)
```

```
##  
## Pearson's product-moment correlation  
##  
## data: cars$speed and cars$dist  
## t = 9.464, df = 48, p-value = 1.49e-12  
## alternative hypothesis: true correlation is not equal to 0  
## 95 percent confidence interval:  
## 0.6816422 0.8862036  
## sample estimates:  
## cor  
## 0.8068949
```

```
plot(cars$speed, cars$dist)
```



There is a significant positive relationship between a car's speed and its stopping distance. The correlation coefficient is 0.81, i.e., 66% of the variation in a car's stopping distance is explained by the car's speed. (Remember, the square of the correlation coefficient, i.e. here $0.81^2=0.66$, tells us the amount of variation explained by the correlation.)

3. Regression

We will do a regression analysis on the data set `cabbages` from the R package `MASS`. The data set contains the weight (`HeadWt`), vitamin C content (`VitC`), the cultivar (`Cult`), and the planting date (`Date`) for 60 cabbage heads:

```
library(MASS) # load the MASS library to make the data set available
head(cabbages)
```

```
##   Cult Date HeadWt VitC
## 1  c39  d16    2.5   51
## 2  c39  d16    2.2   55
## 3  c39  d16    3.1   45
## 4  c39  d16    4.3   42
## 5  c39  d16    2.5   53
## 6  c39  d16    4.3   50
```

Use a multivariate regression to find out whether weight and cultivar have an effect on the vitamin C content. You will need to use the functions `lm()` and `summary()`.

To run a linear regression, you need to first fit the model to the data. This is done with the function `lm()` (`lm` stands for **L**inear **M**odel). The `lm()` function takes two arguments, the formula (here `VitC ~ Cult + HeadWt`) and the data (here `cabbages`). The formula describes what kind of model we want to fit. On the left-hand side of the symbol `~`, we write the response variable, here `VitC`. On the right-hand side, we write the predictor variables we want to use, separated by a `+` sign. Here, we use `Cult` and `HeadWt` as predictor variables. (You can learn more about formulas in R by typing `?formula` into the R console.)

```
fit <- lm(VitC ~ Cult + HeadWt, data=cabbages)
```

Once you have run the linear model, you can then display the results using the `summary()` function:

```
summary(fit)
```

```
##
## Call:
## lm(formula = VitC ~ Cult + HeadWt, data = cabbages)
##
## Residuals:
##      Min       1Q   Median       3Q      Max
## -12.233  -3.796  -1.065   4.542  14.061
##
## Coefficients:
##              Estimate Std. Error t value Pr(>|t|)
## (Intercept)  67.9297     3.1159  21.801 < 2e-16 ***
## Cultc52       9.3578     1.7433   5.368 1.52e-06 ***
## HeadWt      -5.6524     0.9962  -5.674 4.88e-07 ***
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
##
## Residual standard error: 6.304 on 57 degrees of freedom
## Multiple R-squared:  0.625, Adjusted R-squared:  0.6119
## F-statistic: 47.5 on 2 and 57 DF, p-value: 7.234e-13
```

We see that both the cultivar and the weight have a significant effect on vitamin C content. The negative estimate for `HeadWt` indicates that as the weight increases, vitamin C content decreases.

Often, the function `anova()` provides a simpler and cleaner summary of the model fit:

```
anova(fit)
```

```
## Analysis of Variance Table
##
## Response: VitC
##           Df Sum Sq Mean Sq F value    Pr(>F)
## Cult       1 2496.2  2496.15   62.811 9.145e-11 ***
## HeadWt     1 1279.5  1279.48   32.196 4.884e-07 ***
## Residuals 57 2265.2    39.74
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

The conclusions remain unchanged. Both the cultivar and the weight have a significant effect on vitamin C content.

4. If this was easy

Look into the function `predict()`. Can you use it to estimate the vitamin C content of a c52 cultivar with a weight of 4? Can you use it to calculate the residuals of the regression model?

The predict function allows us to predict values from a linear model that we have previously fit. It takes two arguments, the fitted model (`fit` from the previous subsection) and a data frame that has the same columns as were used as predictor variables in the linear model. Thus, to estimate the vitamin C content of a c52 cultivar with a weight of 4, we first create a data frame with one row. Then we run `predict()` :

```
d <- data.frame(Cult="c52", HeadWt=4) # make a data frame with 1 row
predict(fit, d) # run predict on previously fitted model with new data frame
```

```
##          1
## 54.67786
```

We predict that the vitamin C content of a c52 cultivar with a weight of 4 is 54.7.

If we run `predict` with the original data frame then we get the model estimate for each data point (these values are also called the fitted values). By subtracting these values from the original y values we obtain the residuals:

```
residuals <- cabbages$VitC - predict(fit, cabbages)
```

Note that the residuals are also available as `fit$residuals` . The following plot shows that the two sets of numbers are identical:

```
plot(residuals, fit$residuals)
abline(0, 1) # add one-one line
```

