STAT 420: Project 1

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4/13/2020

Part 1

```
library(faraway)
data(prostate)
RMSE = function(model) {
  sqrt(mean(resid(model) ^ 2))
model_1 = lm(lpsa ~ ., data = prostate) # all possible predictors
model_2 = lm(lpsa ~ lcavol, data = prostate) # only lcavol
model_3 = lm(lpsa ~ lcavol + lweight + svi, data = prostate) # best
model_4 = lm(lpsa ~ lcavol + lweight + age + lbph, data = prostate)
model_5 = lm(lpsa ~ lcavol + lweight + svi + lbph + lcp + pgg45, data = prostate)
rs_1 = summary(model_1)$r.squared
rs_2 = summary(model_2)$r.squared
rs_3 = summary(model_3)$r.squared
rs_4 = summary(model_4)$r.squared
rs_5 = summary(model_5)$r.squared
rs = c(rs_1, rs_2, rs_3, rs_4, rs_5)
rmse_1 = RMSE(model_1)
rmse_2 = RMSE(model_2)
rmse_3 = RMSE(model_3)
rmse_4 = RMSE(model_4)
rmse_5 = RMSE(model_5)
rmse = c(rmse_1, rmse_2, rmse_3, rmse_4, rmse_5)
summary(model_1)
```

```
##
## Call:
## lm(formula = lpsa ~ ., data = prostate)
##
## Residuals:
## Min 1Q Median 3Q Max
## -1.7331 -0.3713 -0.0170 0.4141 1.6381
##
```

```
## Coefficients:
##
              Estimate Std. Error t value Pr(>|t|)
## (Intercept) 0.669337 1.296387 0.516 0.60693
## lcavol
             ## lweight
              0.454467
                       0.170012
                                 2.673 0.00896 **
## age
             -0.019637 0.011173 -1.758 0.08229 .
## lbph
             0.107054 0.058449
                                1.832 0.07040 .
                                 3.136 0.00233 **
## svi
              0.766157
                        0.244309
## lcp
             -0.105474
                        0.091013 -1.159 0.24964
## gleason
             0.045142
                        0.157465
                                 0.287 0.77503
## pgg45
              0.004525
                        0.004421
                                  1.024 0.30886
## ---
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
##
## Residual standard error: 0.7084 on 88 degrees of freedom
## Multiple R-squared: 0.6548, Adjusted R-squared: 0.6234
## F-statistic: 20.86 on 8 and 88 DF, p-value: < 2.2e-16
cbind(rs, rmse)
##
             rs
                     rmse
## [1,] 0.6547541 0.6747510
## [2,] 0.5394319 0.7793386
## [3,] 0.6264403 0.7018742
```

I think model 3 (predictors: lcavol, lweight, svi) is the best model because the r squared is not that low and the RMSE is not that high compared to the other models. According to the table provided by summary(model_1), the p-value of these 3 predictors are the lowest (<0.005), which means there is a significant relationship between these predictors and lpsa.

Part 2

[4,] 0.5975750 0.7284869 ## [5,] 0.6425760 0.6865484

```
library(MASS)
data(Boston)

set.seed(42)
train_index = sample(1:nrow(Boston), 400)
train = Boston[train_index,]
test = Boston[-train_index,]

RMSE_2 = function(model, data) {
    yi = data[,c(length(data))]
    yi_hat = predict(model, data)
    sqrt(sum((yi_hat - yi)^2)/nrow(data))
}

model_1 = lm(medv ~ ., data = train) # all possible predictors
model_2 = lm(medv ~ crim, data = train) # only crim
```

```
model_3 = lm(medv ~ crim + nox + rm + dis + rad + ptratio + lstat, data = train)
model_4 = lm(medv \sim crim + nox + rm + dis + rad + ptratio + lstat + tax + zn + black, data = train) # b
model_5 = lm(medv ~ tax + zn + black, data = train)
train1_rmse = RMSE_2(model_1, train)
train2_rmse = RMSE_2(model_2, train)
train3_rmse = RMSE_2(model_3, train)
train4 rmse = RMSE 2(model 4, train)
train5_rmse = RMSE_2(model_5, train)
train_rmse = c(train1_rmse, train2_rmse, train3_rmse, train4_rmse, train5_rmse)
test1_rmse = RMSE_2(model_1, test)
test2_rmse = RMSE_2(model_2, test)
test3_rmse = RMSE_2(model_3, test)
test4_rmse = RMSE_2(model_4, test)
test5_rmse = RMSE_2(model_5, test)
test_rmse = c(test1_rmse, test2_rmse, test3_rmse, test4_rmse, test5_rmse)
summary(model_1)
##
## Call:
## lm(formula = medv ~ ., data = train)
## Residuals:
               1Q Median
                               3Q
## -14.3126 -2.7134 -0.5522 1.5431 25.5431
## Coefficients:
##
              Estimate Std. Error t value Pr(>|t|)
## (Intercept) 40.211363 5.823305 6.905 2.07e-11 ***
             ## crim
## zn
              0.037754 0.016166
                                 2.335 0.020038 *
## indus
              ## chas
              1.918167 0.999327 1.919 0.055663 .
             -17.987178  4.304668  -4.179  3.63e-05 ***
## nox
## rm
              3.478935
                        0.457299
                                  7.608 2.16e-13 ***
             -0.003087
                        0.014798 -0.209 0.834880
## age
## dis
             0.074539
                                 4.167 3.81e-05 ***
## rad
              0.310637
## tax
              -0.011081
                        0.004234 -2.617 0.009212 **
             ## ptratio
## black
              -0.533910  0.055318  -9.652  < 2e-16 ***
## 1stat
## ---
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
## Residual standard error: 4.759 on 386 degrees of freedom
## Multiple R-squared: 0.7262, Adjusted R-squared: 0.7169
## F-statistic: 78.73 on 13 and 386 DF, p-value: < 2.2e-16
```

```
cbind(train_rmse, test_rmse)
```

```
## train_rmse test_rmse
## [1,] 4.675465 4.767746
## [2,] 8.238496 9.318085
## [3,] 4.812485 5.232587
## [4,] 4.698149 4.884765
## [5,] 7.705864 8.000684
```

I think model 4 (predictors: crim, nox, rm, dis, rad, ptratio, lstat, tax, zn, black) is the best model because the RMSE is the second lowest. According to the table provided by summary(model_1), the p-value of these predictors are < 0.001, which means there is a significant relationship between these predictors and medv.

Part 3

(a)

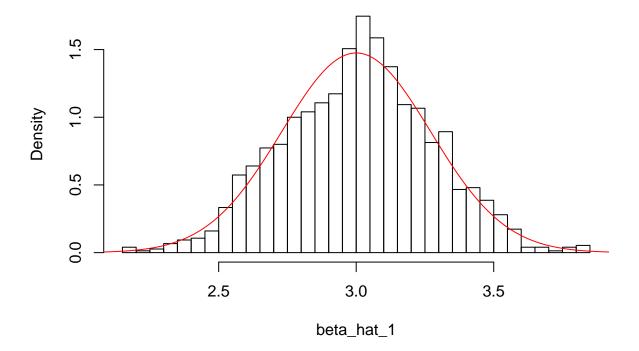
```
set.seed(42)
n = 25
x0 = rep(1, n)
x1 = runif(n, 0, 10)
x2 = runif(n, 0, 10)
x3 = runif(n, 0, 10)
x4 = runif(n, 0, 10)
X = cbind(x0, x1, x2, x3, x4)
C = solve(t(X) %*% X)
y = rep(0, n)
ex_4_data = data.frame(y, x1, x2, x3, x4)
diag(C)
                                     x2
                        x1
## 0.744784994 0.004573055 0.005091328 0.005898213 0.005058979
ex_4_data[10,]
##
              x1
                          x2
                                   xЗ
                                            x4
## 10 0 7.050648 0.03948339 5.144129 7.758234
(b)
```

(c)

beta_hat_1 = numeric(1500)
beta_2_pval = numeric(1500)
beta_3_pval = numeric(1500)

```
for (i in 1:1500) {
  ex_4_data[, 1] = 2 + 3*x1 + 4*x2 + 0*x3 + 1*x4 + rnorm(25, 0, 4)
  y = ex_4_data[,1]
  model = lm(y \sim x1 + x2 + x3 + x4)
  beta_hat_1[i] = summary(model)$coef[2]
  beta_2_pval[i] = summary(model)$coef[3,4]
  beta_3_pval[i] = summary(model)$coef[4,4]
(d)
true_var = (16*C)[2,2]
true_var
## [1] 0.07316889
True distribution of \hat{\beta}_1 is mean: 3 and variance: 0.0731689.
beta_hat_1_mean = mean(beta_hat_1)
beta_hat_1_var = var(beta_hat_1)
hist(beta_hat_1, prob = TRUE, breaks = 50)
x = seq(0, 6, length=1000)
y = dnorm(x, 3, sqrt(true_var))
lines(x,y, col = "red")
```

Histogram of beta_hat_1



```
beta_hat_1_mean
```

[1] 3.006391

beta_hat_1_var

[1] 0.07303341

The mean and variance of beta_hat_1 are close to 3 and 0.0731689. The curve also matches the histogram.

(f)

```
prop = mean(beta_3_pval < 0.05)
prop</pre>
```

[1] 0.0466667

The proportion is 0.0466667, which means most of the tests fail to reject the null hypothesis $(H_0: \beta_3 = 0)$ with $\alpha = 0.05$.

It is expected because the true value of β_3 is 0. (g)

```
mean(beta_2_pval < 0.05)</pre>
```

[1] 1

The proportion is 1, which means all of the tests reject the null hypothesis $(H_0: \beta_2 = 0)$ with $\alpha = 0.05$. It is expected because the true value of β_2 is 4, which is far from 0.