hw2 sml

Group2

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Exercise 1

We start by focusing on the ridge expression. In the first part we want to center the response and the regressors:

$$\sum_{i=1}^{N} \left(y_i - \beta_0 - \sum_{j=1}^{p} (x_{ij} - \overline{x}_j) \beta_j \right) =$$

$$\sum_{i=1}^{N} \left(y_i - \overline{y} - \beta_0 - \sum_{j=1}^{p} \overline{x_j} \beta_j - \sum_{j=1}^{p} (x_{ij} - \overline{x}_j) \beta_j \right) =$$

$$\sum_{i=1}^{N} \left(y_i - \beta_0^c - \sum_{j=1}^{p} (x_{ij} - \overline{x}_j) \beta_j^c \right).$$

Hence, we have that:

$$\beta_0^c = \beta_0 + \sum_{j=1}^p \overline{x_j} \beta_j - \overline{y}$$
$$\beta_j^c = \beta_j \ \ j = 1, 2, \dots, p$$

As a final remark, this centering procedure consist in shifting all the variables x_j and the response y to have mean zero. As a result, only β_0^c which is the intercept, is going to change to be equal to 0, whereas the slope of the regression line β_i^c remain the same.

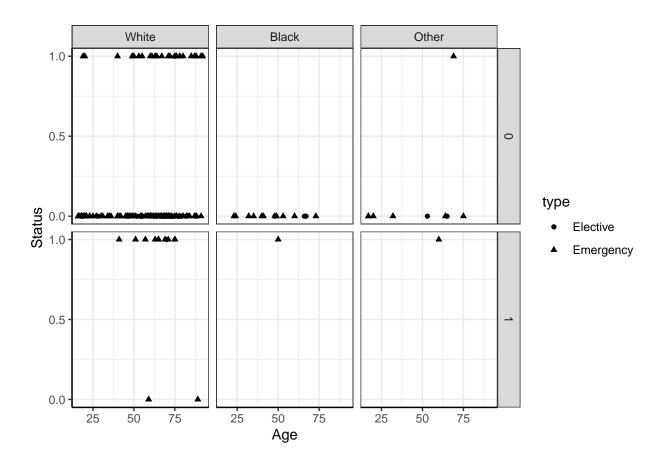
Exercise 9

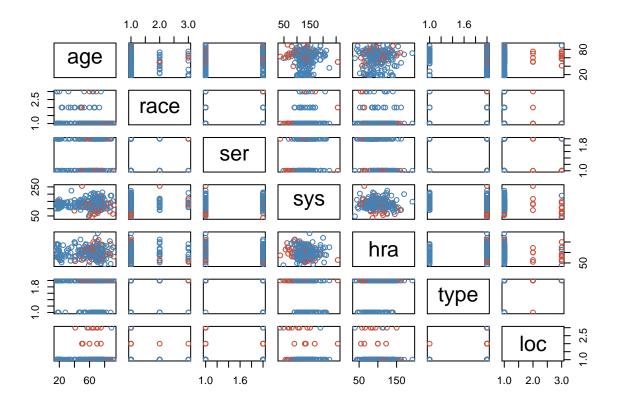
In this exercise, we delve deeper into common problems we might encounter when fitting a logistic regression model for the prediction of categorical data. The problem mentioned here consists of Complete or Quasi-Complete Separation, where the issue is not really connected to the model itself, but it is mainly due to "thin" data. This means, we might have a really skewed distribution in the response variable; in this case we observe only 20% of cases in which the patient has died, in a sample size of 200.

Point A

```
## A - Fit a logistic regression model with all regressors
## A.1 Short exploratory Data Analysis
# Dataframe
df <- icu
# Variables classes
#unlist(lapply(df, class))
# NAs
#unlist(lapply(df, function(x) sum(is.na(x))))
# Change variables of interest into numeric, binarize variables
df <- within(df,</pre>
             # Response variable
             sta <- ifelse(sta == "Died", 1, 0)
             loc_bin <- ifelse(loc == "Nothing", 0, 1)</pre>
             race_bin <- ifelse(race == "White", 1, 0)</pre>
## A.2 Model Fitting and summary. Note locComa and Stupor are both negative,
## but might have same pred power.
m1 <- glm(sta ~., family = binomial(link = "logit"), data = df[, 2:(length(df)-2)])
coef <- m1$coefficients</pre>
```

After running the lositic regression on all the variables we encounter the warning about fitted probabilities being numerically 0 or 1, which is a first sign of **separation**. To graphically illustrate the issue, we can look into the data with a grid plot:





Also in the summary we that the coefficients for locStupor, Black race, Emergency and other regressors are huge, indicating the maximum likelihood estimates do not exist. We can also notice the standard errors of these coefficients to be particularly big in magnitude.

```
##
## Call:
  glm(formula = sta ~ ., family = binomial(link = "logit"), data = df[,
       2:(length(df) - 2)])
##
##
##
  Deviance Residuals:
##
        Min
                          Median
                                        3Q
                   1Q
                                                  Max
   -1.50525
            -0.53717
                       -0.17867
                                  -0.00019
                                             3.01708
##
##
  Coefficients:
##
                   Estimate Std. Error z value Pr(>|z|)
                 -5.548e+00
                                         -2.444
##
  (Intercept)
                              2.271e+00
                                                 0.01454 *
                                          3.055
## age
                  5.645e-02
                              1.848e-02
                                                 0.00225 **
## genderFemale
                 -7.215e-01
                              5.460e-01
                                         -1.321
                                                 0.18639
                                         -0.012
## raceBlack
                 -1.617e+01
                              1.314e+03
                                                 0.99018
## raceOther
                  5.829e-01
                              1.313e+00
                                          0.444
                                                 0.65696
## serSurgical
                 -6.739e-01
                              6.289e-01
                                         -1.071
                                                 0.28398
## canYes
                  3.483e+00
                              1.121e+00
                                          3.106
                                                 0.00189 **
## crnYes
                  1.191e-01
                              8.449e-01
                                          0.141
                                                 0.88786
## infYes
                 -1.081e-01
                              5.557e-01
                                         -0.195
                                                 0.84573
## cprYes
                  1.032e+00
                              9.901e-01
                                          1.043
                                                 0.29714
## sys
                 -2.084e-02 9.443e-03
                                        -2.207 0.02732 *
```

```
## hra
                 -2.915e-03
                              1.032e-02
                                         -0.282
                                                 0.77761
                                          1.822
## preYes
                  1.279e+00
                              7.022e-01
                                                 0.06842
                                          2.792
## typeEmergency
                  3.748e+00
                              1.342e+00
                                                 0.00523 **
## fraYes
                              1.093e+00
                                          1.509
                  1.649e+00
                                                  0.13139
## po2<= 60
                 -6.765e-01
                              9.402e-01
                                         -0.720
                                                 0.47179
## ph< 7.25
                  1.771e+00
                             1.212e+00
                                          1.461
                                                 0.14410
## pco> 45
                 -2.084e+00
                              1.165e+00
                                         -1.789
                                                  0.07361
## bic< 18
                 -2.623e-01
                              8.967e-01
                                         -0.293
                                                  0.76985
## cre> 2.0
                  1.004e-01
                              1.131e+00
                                          0.089
                                                  0.92925
## locStupor
                  3.771e+01
                              2.487e+03
                                          0.015
                                                  0.98790
## locComa
                  3.458e+00
                              1.342e+00
                                          2.578
                                                 0.00994 **
##
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
##
  (Dispersion parameter for binomial family taken to be 1)
##
##
                              on 199
##
       Null deviance: 200.16
                                       degrees of freedom
## Residual deviance: 112.17
                               on 178
                                       degrees of freedom
  AIC: 156.17
##
##
## Number of Fisher Scoring iterations: 17
##
       locStupor
                      raceBlack
                                  (Intercept) typeEmergency
                                                                     canYes
    37.705271929
                                                 3.747879245
                                                               3.482602382
##
                  16.174546113
                                  5.548262182
##
                                     ph< 7.25
                                                                    preYes
         locComa
                        pco> 45
                                                      fraYes
##
     3.458374688
                   2.083580267
                                  1.770978482
                                                 1.649453062
                                                               1.279496861
##
          cprYes
                  genderFemale
                                     po2<= 60
                                                 serSurgical
                                                                 raceOther
##
     1.032231804
                   0.721456637
                                  0.676510682
                                                 0.673862547
                                                               0.582918678
##
         bic< 18
                         crnYes
                                       infYes
                                                    cre> 2.0
                                                                        age
##
                   0.119137902
                                  0.108119860
                                                 0.100402678
     0.262346172
                                                               0.056452457
##
             sys
                            hra
##
     0.020839745
                   0.002915162
```

Point B

So, as we observed in the previous point, we can absolutely make a case for a separation problem. Therefore, we start by pool stratified variable levels into fewer ones.

##	gender1	gender2	race1	race2	race3	ser1
##	"Male"	"Female"	"White"	"Black"	"Other"	"Medical"
##	ser2	can1	can2	crn1	crn2	inf1
##	"Surgical"	"No"	"Yes"	"No"	"Yes"	"No"
##	inf2	cpr1	cpr2	pre1	pre2	type1
##	"Yes"	"No"	"Yes"	"No"	"Yes"	"Elective"
##	type2	fra1	fra2	po21	po22	ph1
##	"Emergency"	"No"	"Yes"	"> 60"	"<= 60"	">= 7.25"
##	ph2	pco1	pco2	bic1	bic2	cre1
##	"< 7.25"	"<= 45"	"> 45"	">= 18"	"< 18"	"<= 2.0"
##	cre2	loc1	loc2	loc3		
##	"> 2.0"	"Nothing"	"Stupor"	"Coma"		

From the listing above we see that only two factors, namely **race** and **loc** have such more than 2 levels, which were already binarized in the beginning. By refitting the full model, we see that pooling the levels together

somewhat improves the modelling, but we still are far from a satisfying results, given the large coefficients still, and standard errors. At this point, one could turn some of the categorical variables into continuous variables, or eliminate some variables. The second option will introduce some bias, but this might be the only thing to do given the sample that we have. We also note that with interaction terms, this problem would be even more accentuated, and we would not be able to determine whether such interactions would be relevant given the separation problem.

```
##
## Call:
##
   glm(formula = sta ~ ., family = binomial(link = "logit"), data = df2[,
##
       2:length(df2)])
##
## Deviance Residuals:
        Min
##
                    10
                          Median
                                         3Q
                                                  Max
  -1.80396 -0.56064
                       -0.20440
                                  -0.08635
                                              2.97729
##
##
  Coefficients:
##
                  Estimate Std. Error z value Pr(>|z|)
## (Intercept)
                  -6.086979
                              2.260515
                                         -2.693
                                                 0.00709 **
## age
                  0.056393
                              0.018624
                                          3.028
                                                 0.00246 **
  genderFemale
                  -0.639725
                              0.531393
                                         -1.204
                                                 0.22864
## serSurgical
                  -0.673522
                              0.601902
                                         -1.119
                                                 0.26315
## canYes
                  3.107051
                              1.045846
                                          2.971
                                                 0.00297 **
## crnYes
                  -0.035708
                              0.801647
                                         -0.045
                                                 0.96447
## infYes
                  -0.204933
                              0.553191
                                         -0.370
                                                 0.71104
## cprYes
                  1.053483
                                          1.047
                                                 0.29530
                              1.006614
## sys
                  -0.015472
                              0.008497
                                         -1.821
                                                 0.06864
                                         -0.288
## hra
                  -0.002769
                              0.009607
                                                 0.77317
                   1.131942
                              0.671450
                                          1.686
                                                 0.09183
## preYes
## typeEmergency
                  3.079583
                              1.081584
                                          2.847
                                                 0.00441 **
## fraYes
                   1.411402
                              1.029705
                                          1.371
                                                 0.17047
## po2<= 60
                  0.073822
                              0.857044
                                          0.086
                                                 0.93136
## ph< 7.25
                              1.208804
                                          1.947
                  2.354078
                                                 0.05148
## pco> 45
                  -3.018442
                              1.253448
                                         -2.408
                                                 0.01604 *
## bic< 18
                                         -0.780
                  -0.709284
                              0.909777
                                                 0.43561
## cre> 2.0
                  0.295143
                              1.116925
                                          0.264
                                                 0.79159
## race_bin
                                          0.610 0.54160
                  0.565729
                              0.926828
                  5.232292
## loc_bin
                              1.226303
                                          4.267 1.98e-05 ***
##
                   0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
## Signif. codes:
##
## (Dispersion parameter for binomial family taken to be 1)
##
##
       Null deviance: 200.16
                               on 199
                                        degrees of freedom
## Residual deviance: 120.78
                               on 180
                                        degrees of freedom
  AIC: 160.78
##
## Number of Fisher Scoring iterations: 6
```

Point D

Now we want to improve the base model and the separation problem by using stepwise procedures based on AIC and BIC. We fit backward stepwise model selection.

The first based on AIC gives out:

$$sta \sim age + can + sys + type + ph + pco + loc_bin$$

 $AIC = 144.4$

For the backward stepwise model selection absed on BIC, we have:

```
sta \sim age + can + type + loc_bin

AIC = 165.63
```

```
s1_AIC<- step(m2, direction = "backward", k = 2)
n <- dim(df2)[1]
s1_BIC <- step(m2, direction = "backward", k = log(n))</pre>
```

Point E

Now that we have obtained our models, we want to compare the full model with the stepwise selected models by benchmarking their log-likelihoods and the in-sample misclassification.

```
pred_AIC <- ifelse(s1_AIC$fitted.values > 0.5, 1, 0)
pred_BIC <- ifelse(s1_BIC$fitted.values > 0.5, 1, 0)
pred_full <- ifelse(m2$fitted.values > 0.5, 1, 0)
## Misclassification errors
mis_AIC <- mean(pred_AIC != df2$sta)
mis_BIC <- mean(pred_BIC != df2$sta)
mis_FULL <- mean(pred_full != df2$sta)</pre>
results_mis <- c(mis_AIC, mis_BIC, mis_FULL)</pre>
## Loglikelihoods
lik_AIC <- logLik(s1_AIC)</pre>
lik_BIC <- logLik(s1_BIC)</pre>
lik_FULL <- logLik(m2)</pre>
results_lik <- c(lik_AIC, lik_BIC, lik_FULL)
## Matrix
result_mat <- matrix(c(results_mis, results_lik), nrow = 2)</pre>
rownames(result_mat) <- c("Misclassification", "Log-Lik")</pre>
colnames(result_mat) <- c("AIC", "BIC", "FULL")</pre>
result_mat
```

```
## AIC BIC FULL
## Misclassification 0.13 0.11 -69.56732
## Log-Lik 0.14 -64.22 -60.38917
```

From these results, we can see that the full model performs better in both metrics. However we need to keep in mind that this is only true in-sample, and a thorough analysis should be carried out also out-of-sample in such cases. This is also expected, since by eliminating covariates, we increase bias to reduce possible variance out-of-sample.