Lab 0 Answer Sketch

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li li li li li	<pre>brary(here) brary(janitor) brary(magrittr) brary(Hmisc) brary(tableone) brary(tidyverse) eme_set(theme_bw()) # I like theme_bw in my ggplots.</pre>	

```
lab0 <- read_csv(here("data", "lab0.csv")) %>%
    type.convert() # converts all characters to factors
```

1 Looking Over the Data Set

Treated

I will start with a quick summary to be sure things are imported properly in the lab0 data set...

1st Qu.:41.74

1st Qu.:46.48

summary(lab0)

subject treatment cov1 cov2

Min. :101.0 Not Treated:95 Min. :27.03 Min. :29.49

Median :47.70 Median :53.48 Median :228.0 :210.2 :48.36 :52.68 Mean Mean Mean 3rd Qu.:261.5 3rd Qu.:56.62 3rd Qu.:59.99 Max. :295.0 Max. :73.28 Max. :73.25

:40

 cov3
 cov4
 cov5

 Min. : 8.00
 Min. : 9.00
 Min. : 0.0000

 1st Qu.:17.00
 1st Qu.:0.0000

 Median : 20.00
 Median : 20.00
 Median : 0.0000

Mean :20.28 Mean :20.06 Mean :0.4296 3rd Qu.:24.00 3rd Qu.:23.00 3rd Qu.:1.0000 Max. :33.00 Max. :33.00 Max. :1.0000

1.1 Using the describe function

Alternatively, we could use the describe function, which is part of the Hmisc package...

Hmisc::describe(lab0)

1st Qu.:134.5

lab0

210.2 107.7 135 0 135 1 70.54 114.4 .25 .50 .75 .90 .95 228.0 261.5 134.5 281.6 288.3

lowest: 101 102 103 104 105, highest: 291 292 293 294 295

treatment n missing distinct 135 0 Value Not Treated Treated Frequency Proportion 0.704 0.296 cov1 Info . 05 n missing distinct Mean Gmd . 10 48.36 11.83 30.70 34.36 0 133 1 48.36 .50 .75 .90 .95 135 . 25 41.75 47.70 56.62 61.84 64.94 lowest: 27.03 28.30 28.39 28.73 29.60, highest: 66.37 67.30 67.89 68.39 73.28 n missing distinct Info Mean Gmd . 05 . 10 1 52.68 11.08 36.81 39.15 0 133 .75 .90 . 25 .50 . 95 53.48 59.99 65.39 67.05 46.48 lowest: 29.49 31.93 32.24 34.24 34.25, highest: 68.90 69.55 70.10 72.03 73.25 ______ cov3 Mean 20.28 .10 n missing distinct Info Gmd .05 135 0 24 0.995 5.87 12 14 .90 . 95 .50 .75 . 25 20 24 27 17 29 lowest: 8 9 11 12 13, highest: 28 29 30 32 33 ______ cov4 Mean Gmd .05 . 10 n missing distinct Info 135 23 0.994 20.06 4.929 13 15 0 .75 .90 .95 . 25 .50 17 20 23 26 28 lowest: 9 11 12 13 14, highest: 28 29 31 32 33 ______

 $\operatorname{\mathtt{Sum}}$

58

Mean

0.4296

Gmd

0.4938

Info

0.735

2

cov5

135

n missing distinct

0

1.2 Glimpsing the data's structure

Or, perhaps we just want to see the structure of the data and some of the first few values in each variable, in which case, the str command would help, or we could use the dplyr package's glimpse function...

glimpse(lab0)

The two treatment options are named "Treated" and "Not Treated", as opposed to "Treated" and "Untreated". Anything so that the thing I wanted to evaluate probabilities for (i.e. Treated as compared to Not Treated) came second alphabetically is appealing, because R, by default, treats the first level in a binary categorical variable as unsuccessful and the second level as successful and generally orders levels of binary variables alphabetically.

2 Fitting a Logistic Regression Model using the glm function

We are fitting a model to predict the probability of "Treated" here. If we want to see what's in m1, we can type it in, and see what we get, or ask for a summary, and get some additional details.

```
m1
```

Coefficients:

```
(Intercept) cov1 cov2 cov3 cov4 cov5 0.23905 0.04159 -0.02512 -0.18594 0.06993 0.79492
```

```
Degrees of Freedom: 134 Total (i.e. Null); 129 Residual
Null Deviance:
                    164.1
Residual Deviance: 135 AIC: 147
summary (m1)
Call:
glm(formula = treatment == "Treated" ~ cov1 + cov2 + cov3 + cov4 +
    cov5, family = binomial(), data = lab0)
Deviance Residuals:
    Min
              1Q
                   Median
                                        Max
                                3Q
                -0.4914
-1.9216
        -0.7418
                                     2.4184
                            0.8746
Coefficients:
            Estimate Std. Error z value Pr(>|z|)
                                  0.119
(Intercept) 0.23905
                        2.00561
                                          0.9051
cov1
             0.04159
                        0.02198
                                  1.892
                                          0.0585 .
cov2
            -0.02512
                        0.02303 -1.091
                                          0.2754
                        0.04735 -3.927 8.6e-05 ***
cov3
            -0.18594
             0.06993
                        0.05053
                                  1.384
                                          0.1664
cov4
cov5
             0.79492
                        0.44205
                                  1.798
                                          0.0721 .
Signif. codes:
               0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
(Dispersion parameter for binomial family taken to be 1)
    Null deviance: 164.08 on 134 degrees of freedom
Residual deviance: 135.02 on 129 degrees of freedom
AIC: 147.02
```

2.1 Storing Probabilities and Linear Predictions

Number of Fisher Scoring iterations: 4

To store the linear predictions (i.e. log odds of the estimated probabilities,) and the estimated probabilities themselves as part of the lab0 data file, I'll use the following commands:

```
lab0$linpred <- m1$linear.predictors
lab0$prob <- m1$fitted.values</pre>
```

The remaining tasks in the assignment essentially require you to obtain some numerical (perhaps) and graphical (mandatory) summaries of the estimated probabilities broken down into the two treatment groups.

3 Some Numerical Summaries of the Fitted Probabilities by Treatment Group

3.1 Using dplyr and summarise

The dplyr library can be used to compare the probs across the two treatment groups, along with some piping commands, to create a little data frame of the summaries you're interested in, as follows:

3.2 Using the by command

by(lab0\$prob, lab0\$treatment, describe)

Some of you may be more familiar with the by command - that works, as well...

```
labO$treatment: Not Treated
dd[x,]
                                                             .05
          missing distinct
                                Info
                                         Mean
                                                   Gmd
                                                                       .10
      95
                0
                        95
                                   1
                                       0.2343
                                                0.1891 0.04319 0.05634
     .25
              .50
                        .75
                                 .90
                                          .95
 0.10597
          0.18805 0.31990 0.47906 0.55551
```

lowest: 0.01772653 0.02853116 0.03021174 0.03724054 0.04076704 highest: 0.60114577 0.60689182 0.71653717 0.82781846 0.84216789

```
lab0$treatment: Treated
dd[x,]
          missing distinct
                                 Info
                                           Mean
                                                      Gmd
                                                                .05
                                                                          .10
      40
                 0
                          40
                                     1
                                         0.4435
                                                   0.2411
                                                             0.1284
                                                                      0.1543
                                   .90
     .25
               .50
                         .75
                                             .95
  0.2972
           0.4299
                     0.6032
                               0.6858
                                         0.7658
```

lowest: 0.05370511 0.10947830 0.12944077 0.15246158 0.15452007

3.3 Using the tableone library

Or, you could use the tableone library to produce a summarized Table 1 describing our results...

Stratified by treatment Not Treated Treated test р 40 95 prob (mean (SD)) 0.23 (0.18) 0.44 (0.21) < 0.001 linpred (mean (SD)) -1.44 (1.09) -0.29 (1.01) <0.001 cov1 (mean (SD)) 47.07 (10.20) 51.42 (10.01) 0.025 53.40 (9.79) cov2 (mean (SD)) 50.98 (9.30) 0.187 cov3 (mean (SD)) 21.45 (4.98) 17.50 (4.74) <0.001 cov4 (mean (SD)) 19.66 (4.17) 21.00 (4.83) 0.107 cov5 = 1 (%)38 (40.0) 20 (50.0) 0.378

You could even use non-parametric tests, and report quartiles for the continuous covariates...

```
print(tab1, nonnorm=c("prob", "linpred", "cov1", "cov2", "cov3", "cov4"))
```

```
Stratified by treatment
                       Not Treated
                                             Treated
                                                                   p
                           95
                                                 40
n
prob (median [IQR])
                        0.19 [0.11, 0.32]
                                              0.43 [0.30, 0.60]
                                                                   <0.001
linpred (median [IQR]) -1.46 [-2.13, -0.75] -0.28 [-0.86, 0.42]
                                                                   <0.001
cov1 (median [IQR])
                       46.42 [39.97, 53.94] 51.07 [43.32, 60.79]
                                                                    0.046
                       54.08 [47.08, 60.36] 51.66 [44.02, 55.49]
cov2 (median [IQR])
                                                                    0.122
cov3 (median [IQR])
                       21.00 [18.00, 24.00] 18.00 [14.75, 21.00] < 0.001
                       20.00 [17.00, 23.00] 19.50 [18.00, 24.00]
cov4 (median [IQR])
                                                                    0.267
                           38 (40.0)
                                                20 (50.0)
cov5 = 1 (\%)
                                                                    0.378
                      Stratified by treatment
                       test
n
prob (median [IQR])
                       nonnorm
linpred (median [IQR]) nonnorm
cov1 (median [IQR])
                       nonnorm
cov2 (median [IQR])
                       nonnorm
```

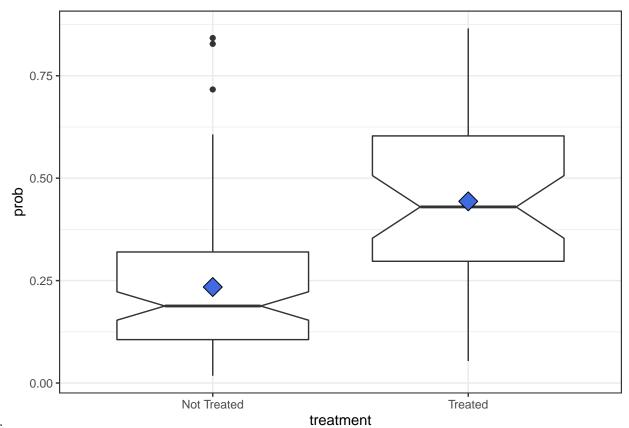
4 Plotting the Fitted Probabilities by Treatment Group

OK. So we've seen a numerical summary - let's focus on the important issue - a plot.

4.1 A Boxplot using ggplot2, with Notches and Means Indicated

```
ggplot(lab0, aes(x = treatment, y = prob)) +
  geom_boxplot(notch=TRUE) +
  stat_summary(fun.y="mean", geom="point", shape=23, size = 5, fill = "royalblue")
```

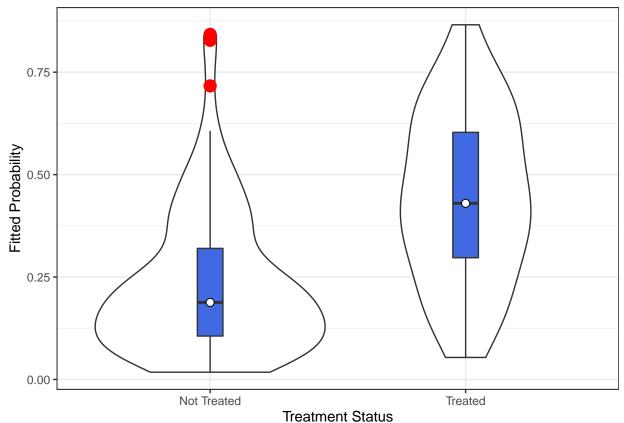
Warning: `fun.y` is deprecated. Use `fun` instead.



3-1.pdf

4.2 A Violin Plot

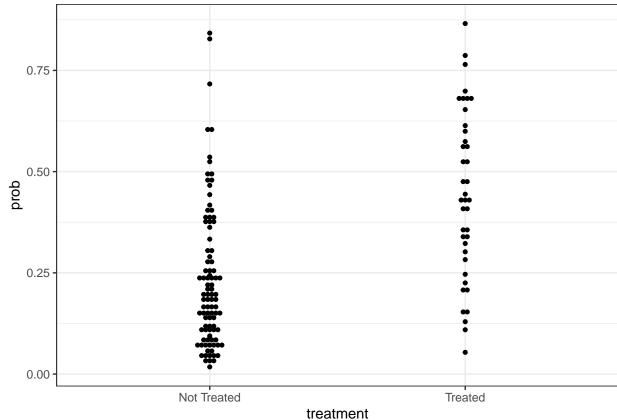
Warning: `fun.y` is deprecated. Use `fun` instead.



5-1.pdf

4.3 A DotPlot to compare the probabilities, via ggplot2

```
ggplot(lab0, aes(x = treatment, y = prob)) +
  geom_dotplot(binaxis="y", binwidth=0.01, stackdir="center")
```

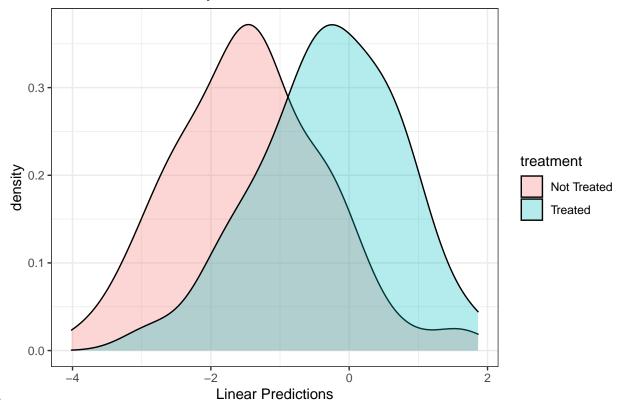


7-1.pdf

4.4 A Density Plot, using ggplot2

A possibly more impressive picture would be a density plot. The best way to get this (here, I'll look at the linear probability (i.e. log odds of treatment) results rather than the raw probabilities on a 0-1 scale just to see if we observe something different) uses the ggplot2 library again...

Linear Predictions By Treatment



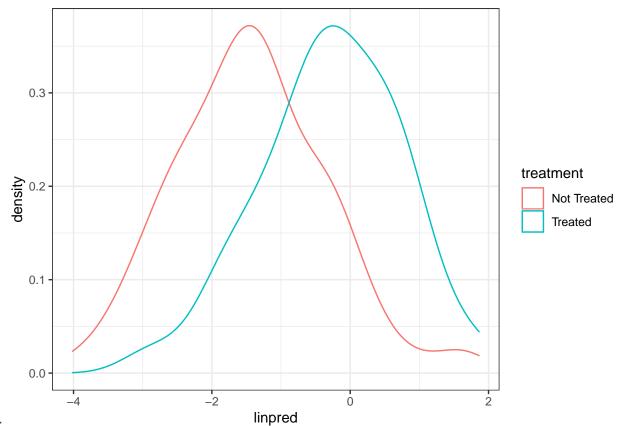
8-1.pdf

One advantage of the linear probabilities over the raw probability estimates is that the log odds results (linear probabilities) are a bit more likely to follow a normalish distribution. Again, it looks like there is fairly substantial overlap in the fitted probabilities across the treatment groups.

4.5 Another Density Plot, using ggplot2

We can use color instead of fill to indicate the densities.

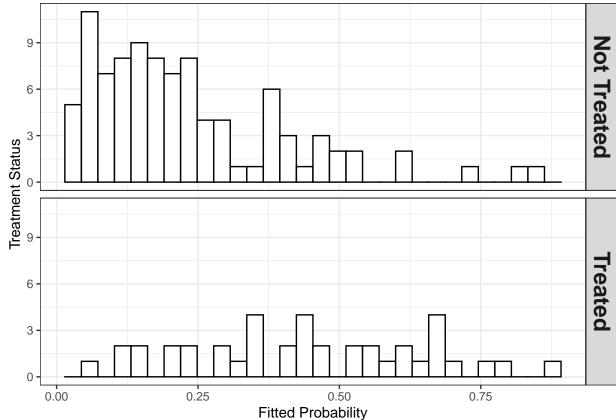
```
ggplot(lab0, aes(x=linpred, color=treatment)) +
  geom_density() +
  theme_bw()
```



4.6 Our Old Standby - Comparing Distributions via Histograms

The slickest approach I have here is this:

```
ggplot(lab0, aes(x = prob)) +
  geom_histogram(fill="white", color="black") +
  facet_grid(treatment ~ .) +
  theme(strip.text = element_text(face="bold", size=rel(1.5))) +
  xlab("Fitted Probability") + ylab("Treatment Status")
```

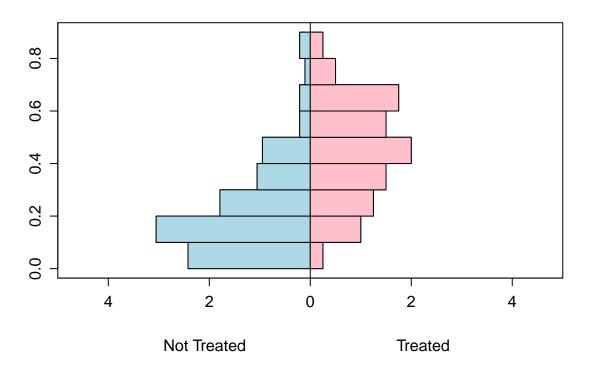


10-1.pdf

4.7 A Back-to-Back Histogram

A former student suggested this approach, from the Hmisc library. There are likely better ways to get such a plot out of R.

Back to Back Histogram of Fitted Probabilities



11-1.pdf

5 What About the ROC Curve and C Statistic?

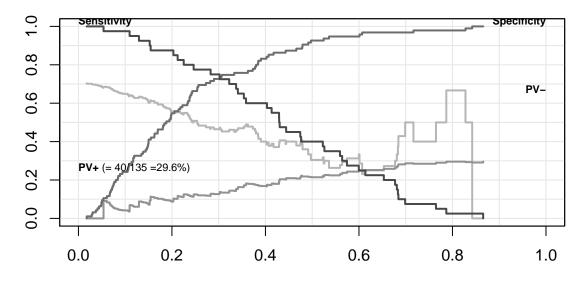
Recall that our model m1 was

```
Call: glm(formula = treatment == "Treated" ~ cov1 + cov2 + cov3 + cov4 +
cov5, family = binomial(), data = lab0)
```

Since we're looking at a logistic regression, someone in a previous version of this class asked if I could show you how I get the C statistic (area under the ROC curve) for such a model. I usually use the Epi library . . .

Note that we need to specify the formula (abbreviated form in the ROC function) again, but that's it to get these results.

- The C statistic (area under the curve) for this logistic regression model is 0.786
- Very briefly, the ability of the model's predicted values to discriminate between patients with one outcome vs. the other is quantified by the area under the curve, also called the C statistic or concordance index, which ranges from 0.5 (discrimination is not better than chance) to 1.0 (perfect discriminating power.)
- The ROC procedure comes from signal detection theory and has been adopted into the language of diagnostic testing, essentially treating the response in the logistic regression model as the true status variable, and the set of predictors as the test to be evaluated by things like sensitivity, specificity, and positive and negative predictive values based on dichotomizing along the levels of the predictor set.
- For more on the ROC, visit Wikipedia for Receiver Operating Characteristic. Or try Google.
- A value of 0.786 would indicate a less-than-terrific model in terms of this issue. Values of 0.8 or even 0.9 are usually needed to declare the model to be reasonably accurate in this sense.



Cutpoint for predicted probability

