Answer Sketch for 432 Quiz 1 Honors Opportunity

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```
library(janitor)
library(leaps)
library(rms)
library(broom)
library(tidyverse)
```

Setup for Questions 1-3

The data in the honors.csv file contain information for 255 subjects on:

- a binary outcome (Good or Bad),
- a size (quantitative, between 60 and 200, in millimeters),
- an indicator of whether a treatment was used (1 = treatment was used or 0 = treatment was not used), and
- a specification as to which of five ordered groups (1 = lowest, 5 = highest) by socio-economic status (ses_group) the subject falls in, along with
- a subject ID.

Import the data into the honors frame, and then fit a logistic regression model to predict the log odds of a Good outcome using the subject's size, treatment status and ses_group, treating the ses_group as a categorical variable. Questions 1-3 use a complete case analysis. One such analysis yields these results.

Here is the actual m1 model that was fit:

term	estimate	std.error	statistic	p.value	conf.low	conf.high
(Intercept)	0.155	0.822	-2.267	0.023	0.030	0.751
size	1.010	0.006	1.677	0.093	0.998	1.022
treatment	0.557	0.291	-2.010	0.044	0.314	0.985
ses_group2	1.226	0.601	0.339	0.735	0.378	4.087
ses_group3	1.384	0.523	0.622	0.534	0.510	4.051
ses_group4	1.110	0.558	0.188	0.851	0.378	3.442
ses_group5	1.416	0.511	0.680	0.496	0.535	4.068

The output below comes from another approach to fitting the identical logistic regression model that we saw previously, still using only the complete cases. I'll call this model m1a, to emphasize that it contains the same outcome and predictors, put together in the same way. Here is what was fit in m1a:

```
d <- datadist(honors)
options(datadist = "d")</pre>
```

```
m1a <- lrm(goodoutcome ~ size + treatment + ses_group, data = honors)
m1a</pre>
```

```
Frequencies of Missing Values Due to Each Variable goodoutcome size treatment ses_group

0 5 4 14
```

Logistic Regression Model

lrm(formula = goodoutcome ~ size + treatment + ses_group, data = honors)

		Model Likelihood		Discrim	ination	Rank D	iscrim.
		Ratio Te	est	Inde	xes	Inde	exes
0bs	233	LR chi2	8.34	R2	0.049	C	0.614
0	160	d.f.	6	g	0.479	Dxy	0.228
1	73	Pr(> chi2)	0.2143	gr	1.615	gamma	0.229
max deriv	v 3e-11			gp	0.100	tau-a	0.099
				Brier	0.208		

```
Coef S.E. Wald Z Pr(>|Z|)
Intercept -1.8628 0.8216 -2.27 0.0234
size 0.0099 0.0059 1.68 0.0935
treatment -0.5852 0.2911 -2.01 0.0444
ses_group=2 0.2035 0.6007 0.34 0.7348
ses_group=3 0.3251 0.5226 0.62 0.5339
ses_group=4 0.1047 0.5576 0.19 0.8511
ses_group=5 0.3479 0.5115 0.68 0.4964
```

summary(m1a) %>% knitr::kable(digits = 3)

	Low	High	Diff.	Effect	S.E.	Lower 0.95	Upper 0.95	Type
size	100.525	135.5	34.975	0.346	0.207	-0.058	0.751	1
Odds Ratio	100.525	135.5	34.975	1.414	NA	0.943	2.120	2
treatment	0.000	1.0	1.000	-0.585	0.291	-1.156	-0.015	1
Odds Ratio	0.000	1.0	1.000	0.557	NA	0.315	0.986	2
$ses_group - 1:5$	5.000	1.0	NA	-0.348	0.511	-1.350	0.655	1
Odds Ratio	5.000	1.0	NA	0.706	NA	0.259	1.924	2
$ses_group - 2:5$	5.000	2.0	NA	-0.144	0.481	-1.086	0.798	1
Odds Ratio	5.000	2.0	NA	0.866	NA	0.337	2.220	2
$ses_group - 3:5$	5.000	3.0	NA	-0.023	0.379	-0.765	0.720	1
Odds Ratio	5.000	3.0	NA	0.978	NA	0.465	2.054	2
$ses_group - 4:5$	5.000	4.0	NA	-0.243	0.425	-1.075	0.589	1
Odds Ratio	5.000	4.0	NA	0.784	NA	0.341	1.802	2

0.1 Note on the importance of using goodoutcome rather than outcome

- Note that if you incorrectly used outcome in the glm fit, rather than goodoutcome, you get an error message, since the values aren't 0 or 1.
- But if you used outcome in the 1rm fit, rather than goodoutcome, the machine assumes that what you want to look at is the factor version of outcome, and in this case, because Good comes alphabetically

after Bad, it fits the same model (m1a) as above.

1 Question 1 (1 point)

What do you conclude from the m1a summary about the odds ratio and confidence interval associated with the treatment variable? To answer this question, provide a complete description (in complete English sentences) of the odds ratio effect associated with treatment in the summary(m1a) output. This should require two or three sentences.

1.1 Answer for Question 1

The odds of a good outcome are estimated to be 0.557 times as large for a subject receiving the treatment than they are for a subject of the same size and the same ses_group who is not receiving the treatment. It wasn't necessary here to mention the 95% confidence interval for this odds ratio, which is (0.315, 0.986), which indicates that this effect is large enough to reach our usual standard to declare the effect of treatment to be statistically significant. Apparently, adjusting for size and ses_group, treatment is associated with a higher chance of a bad outcome in this model.

• English can be tricky. It is reasonable to write 0.557 times as high, but it isn't reasonable to write 0.557 times higher. That (higher) would only work if the value of the odds ratio was larger than 1. So I'd stick with "XXX times as large" or "XXX times as high".

2 Question 2 (1 point)

Why is the odds ratio shown in the m1a output for size different from that shown in the earlier presentation using tidy for the m1 model? Keep your answer to two or three sentences.

2.1 Answer for Question 2

In m1a, the default choices for summary (in this 1rm model) describe the impact of moving from a size at the 25th percentile of the data (100.525 mm) to a size at the 75th percentile of the data (135.5 mm). In m1, the default choice of summary (in this glm fit) describe the impact of moving 1 mm (for example, from 100.525 mm to 101.525 mm). Hence, the estimated effect of such a change on the odds of a good outcome appears much larger in the m1a output.

Note that while it is also true that the baseline category for ses_group changes from category 1 (in m1) to category 5 (in m1a) this has no impact on the odds ratio for size in the summary(m1a) output. Suppose, for instance that we reran m1 but now forcing ses_group = 5 to be the baseline category. The odds ratios estimated for size would not change, but everything else would match up perfectly with the m1a output. We would get:

```
honors_rev <- honors %>%
    mutate(ses_group = fct_relevel(ses_group, "5", "1", "2", "3", "4"))

m1_rev <- glm(goodoutcome ~ size + treatment + ses_group, data = honors_rev, family = binomial)

tidy(m1_rev, conf.int = TRUE, conf.level = 0.95,
    exponentiate = TRUE) %>% knitr::kable(digits = 3)
```

term	estimate	std.error	statistic	p.value	conf.low	conf.high
(Intercept)	0.220	0.752	-2.013	0.044	0.049	0.945
size	1.010	0.006	1.677	0.093	0.998	1.022
treatment	0.557	0.291	-2.010	0.044	0.314	0.985
ses_group1	0.706	0.511	-0.680	0.496	0.246	1.869

term	estimate	std.error	statistic	p.value	conf.low	conf.high
ses_group2	0.866	0.481	-0.300	0.764	0.327	2.184
ses_group3	0.978	0.379	-0.060	0.952	0.463	2.054
ses_group4	0.784	0.425	-0.573	0.567	0.335	1.786

3 Question 3 (1 point)

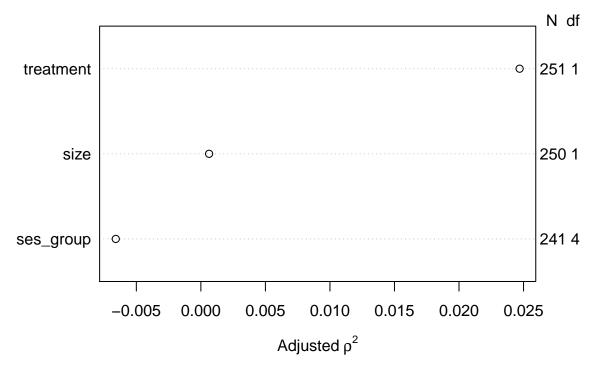
Using the honors data (again without imputing any missing values), obtain a Spearman ρ^2 plot and use it to identify a good way to add a single additional non-linear term to this model (you may spend only a single additional degree of freedom). What addition would you make? This should be explained in one or more complete English sentences.

3.1 Answer for Question 3

The Spearman ρ^2 plot without doing any imputation is:

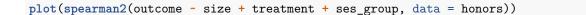
```
plot(spearman2(goodoutcome ~ size + treatment + ses_group, data = honors))
```



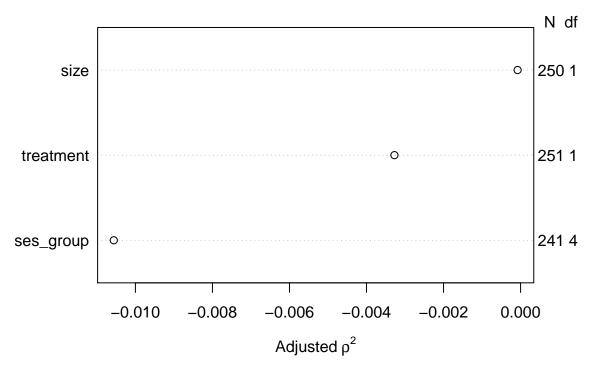


treatment is the most promising variable, and is binary. size is next, which is quantitative, so it looks like adding a treatment-size interaction is the most promising way to spend a single additional degree of freedom.

- Note the importance of using goodoutcome as the outcome of interest here. It would be equally reasonable to use outcome = "good", but not to just use outcome, because outcome sets up different results for the squared Spearman correlation.
- The plot below demonstrates the **wrong** approach...



Spearman ρ^2 Response : outcome



Setup for Questions 4 and 5

Using the honors data, fit the model you specified in Question 3 (including the non-linear term), while also accounting for missing data using **multiple imputation**. Set your seed to be 432432, and impute the predictors that need imputation using all available observations on all available variables, with 20 imputations. Be sure to show the code you used to fit your imputation model and your outcome model in your HTML file. Call the imputation model model_imp and the outcome model m2.

3.2 Models Fit by Dr. Love in Developing this Sketch

Here's a count by variable of the missingness in the data:

So we're missing nothing in our goodoutcome variable, but are missing some size, treatment and ses_group information.

Here is the imputation model I fit:

```
set.seed(432432)
d <- datadist(honors)</pre>
```

• Note that I included goodoutcome in this model, and that I didn't include any interaction term.

Here is the outcome model, including the interaction term:

Variance Inflation Factors Due to Imputation:

ses_group=2	treatment	size	Intercept
1.06	1.03	1.02	1.03
<pre>size * treatment</pre>	ses_group=5	ses_group=4	ses_group=3
1.03	1.07	1.08	1.03

Rate of Missing Information:

ses_group=2	treatment	size	Intercept
0.06	0.03	0.02	0.03
<pre>size * treatment</pre>	ses_group=5 si	ses_group=4	ses_group=3
0.02	0.07	0.07	0.03

d.f. for t-distribution for Tests of Single Coefficients:

```
Intercept size treatment ses_group=2 27411.89 65215.54 23496.55 5272.14 ses_group=3 ses_group=4 ses_group=5 size * treatment 23016.94 3627.65 4371.00 30883.55
```

The following fit components were averaged over the 20 model fits:

```
stats linear.predictors
```

Logistic Regression Model

```
fit.mult.impute(formula = goodoutcome ~ size + treatment + ses_group +
    size * treatment, fitter = lrm, xtrans = model_imp, data = honors,
    x = TRUE, y = TRUE)
```

Model Likelihood Discrimination Rank Discrim.

]	Ratio Te	est	Inde	exes	Inde	exes
Obs 25	5 LR	chi2	12.85	R2	0.069	C	0.633
0 17	3 d.f		7	g	0.579	Dxy	0.266
1 8	2 Pr(> chi2)	0.0809	gr	1.786	gamma	0.267
max deriv 2e-0	8			gp	0.118	tau-a	0.117
				Brier	0.208		
	Coef	S.E.	${\tt Wald}\ {\tt Z}$	Pr(> Z)			
Intercept	-0.6330	1.0801	-0.59	0.5578			
size	-0.0007	0.0083	-0.08	0.9356			
treatment	-2.3813	1.3930	-1.71	0.0874			
ses_group=2	0.2779	0.5988	0.46	0.6425			
ses_group=3	0.5416	0.5043	1.07	0.2828			
ses_group=4	0.2280	0.5557	0.41	0.6817			
ses_group=5	0.4747	0.5033	0.94	0.3456			
size * treatment	0.0140	0.0115	1.22	0.2213			

4 Question 4 (1 point)

If Harry was size 100 mm and fell into group 4 in socio-economic status and Sally was size 120 mm and fell into group 3 in socio-economic status, and both Harry and Sally received the treatment, which of the two would have a larger probability of a Good outcome according to your model? How do you know? Your answer should be given in complete English sentences.

4.1 Answer for Question 4

The direct predictions in terms of probabilities can be made from the lrm model as follows:

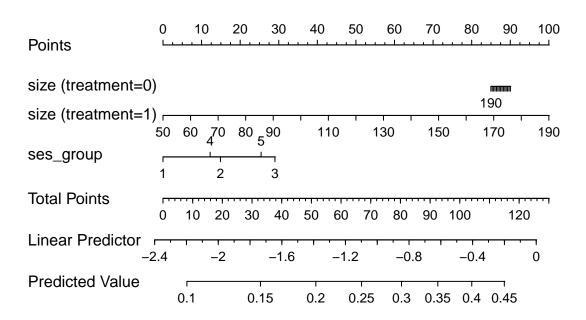
1 2 0.1895625 0.2947395

So Sally's probability is definitely larger than Harry's.

• Note: In the Friday piece, I used a tibble here, rather than data.frame but the tibble forces you to say something about treating the ses_group as a factor.

Another way to assess this would be with a nomogram. Sally's larger size (since they each were treated) and group 3 status (as opposed to Harry's group 4) will yield a clearly larger probability of a good outcome for Sally.

plot(nomogram(m2, fun = plogis))



• Note the need for plogis in the nomogram call in order to make predictions in terms of probabilities.

4.1.1 What if we used a different imputation method?

Other reasonable imputation models would have been:

Variance Inflation Factors Due to Imputation:

Intercept size treatment ses_group=2

```
1.02 1.02 1.03 1.11

ses_group=3 ses_group=4 ses_group=5 size * treatment

1.09 1.08 1.06 1.03
```

Rate of Missing Information:

```
Intercept size treatment ses_group=2 0.02 0.02 0.03 0.10 ses_group=3 ses_group=4 ses_group=5 size * treatment 0.08 0.08 0.06 0.02
```

d.f. for t-distribution for Tests of Single Coefficients:

Intercept	size	treatment	ses_group=2
42713.59	70102.29	28019.38	1832.94
ses_group=3	ses_group=4	ses_group=5	<pre>size * treatment</pre>
3094.34	3110.13	5342.33	30918.64

The following fit components were averaged over the 20 model fits:

stats linear.predictors

```
predict(m2Z, newdat, type = "fitted")
```

0.1904870 0.2959125

which yields predictions of essentially 0.19 for Harry and 0.30 for Sally,

and

Variance Inflation Factors Due to Imputation:

```
Intercept size treatment ses_group=2
1.03 1.02 1.03 1.08
ses_group=3 ses_group=4 ses_group=5 size * treatment
1.04 1.09 1.07 1.02
```

Rate of Missing Information:

ses_group=2	treatment	size	Intercept
0.07	0.03	0.02	0.03
<pre>size * treatment</pre>	ses_group=5	ses_group=4	ses_group=3
0.02	0.07	0.08	0.04

d.f. for t-distribution for Tests of Single Coefficients:

ses_group=2	treatment	size	Intercept
3786.92	27581.92	72518.76	28265.31
<pre>size * treatment</pre>	ses_group=5	ses_group=4	ses_group=3
36559.09	4169.24	2649.40	13779.68

The following fit components were averaged over the 20 model fits:

stats linear.predictors

```
predict(m2Y, newdat, type = "fitted")
```

1 2 0.1901077 0.2957259

which yields predictions of essentially 0.19 for Harry and 0.30 for Sally, as well.

5 Question 5 (1 point)

Write a few English sentences describing how the addition of imputation and a non-linear term changes (or doesn't change) the conclusions that you draw in m2 from what you saw in the m1 (or, equivalently, the m1a) model examined earlier.

5.1 Answer for Question 5

The main things I was looking for:

- The appropriate conclusion is that the addition of imputation and an interaction effect have had at most a modest impact on the conclusions of the model. It remains a weak model.
- In the model post-imputation, (m2) it looks like the impact of size for those without the treatment is much more modest than the impact of size when the treatment is received, according to the nomogram for model m2 shown earlier.

There were lots of ways to get to those conclusions.

m1a

Logistic Regression Model

lrm(formula = goodoutcome ~ size + treatment + ses_group, data = honors)

		Model Likelihood		Discr	imination	Rank I	Discrim.
		Ratio T	est	Ind	dexes	Inc	dexes
0bs	233	LR chi2	8.34	R2	0.049	С	0.614
0	160	d.f.	6	g	0.479	Dxv	0.228

```
1
               73
                     Pr(> chi2) 0.2143
                                                     1.615
                                                                       0.229
                                            gr
                                                              gamma
max |deriv| 3e-11
                                                     0.100
                                                                       0.099
                                                              tau-a
                                           gp
                                           Brier
                                                     0.208
            Coef
                     S.E.
                            Wald Z Pr(>|Z|)
            -1.8628 0.8216 -2.27
                                   0.0234
Intercept
             0.0099 0.0059 1.68
                                   0.0935
size
            -0.5852 0.2911 -2.01
treatment
                                   0.0444
ses_group=2
             0.2035 0.6007
                             0.34
                                   0.7348
ses_group=3
             0.3251 0.5226
                             0.62
                                   0.5339
ses_group=4
             0.1047 0.5576
                             0.19
                                   0.8511
ses_group=5
             0.3479 0.5115
                             0.68
                                   0.4964
```

m2

Logistic Regression Model

```
fit.mult.impute(formula = goodoutcome ~ size + treatment + ses_group +
    size * treatment, fitter = lrm, xtrans = model_imp, data = honors,
    x = TRUE, y = TRUE)
```

		Model Likelihood Ratio Test		Discrimination Indexes		Rank Discrim. Indexes	
0bs	255	LR chi2	12.85	R2	0.069	C	0.633
0	173	d.f.	7	g	0.579	Dxy	0.266
1	82	Pr(> chi2)	0.0809	gr	1.786	gamma	0.267
max deri	v 2e-08			gp	0.118	tau-a	0.117
				Brier	0.208		

```
Coef
                         S.E.
                                Wald Z Pr(>|Z|)
Intercept
                 -0.6330 1.0801 -0.59 0.5578
size
                 -0.0007 0.0083 -0.08
                                       0.9356
treatment
                 -2.3813 1.3930 -1.71
                                       0.0874
                  0.2779 0.5988
                                0.46
                                       0.6425
ses_group=2
ses group=3
                  0.5416 0.5043
                                 1.07
                                       0.2828
                  0.2280 0.5557
ses_group=4
                                0.41
                                       0.6817
ses group=5
                  0.4747 0.5033 0.94
                                       0.3456
size * treatment 0.0140 0.0115 1.22 0.2213
```

In the m1 models, the effect of treatment is statistically significant at the 5% level after accounting for size and ses_group, but in the m2 model, after the inclusion of the interaction term, this no longer appears to be the case. The interaction term, like the rest of m2, carries no statistical significance by Wald tests in m2, whereas in m1a, the treatment effect appeared to be just under our usual standard for statistical significance. In ANOVA testing for m2, shown below, we can see the combined impact of treatment (main effect + interaction) does still exhibit a p value below 0.05.

anova(m2)

Wald Statistics	Response:	goodoutcome
-----------------	-----------	-------------

Factor	Chi-Square	d.f.	P
size (Factor+Higher Order Factors)	2.91	2	0.2339
All Interactions	1.50	1	0.2213
treatment (Factor+Higher Order Factors)	7.97	2	0.0186

All Interactions				1.50	1	0.2213
ses_group				1.55	4	0.8173
<pre>size * treatment</pre>	(Factor+Higher	Order	Factors)	1.50	1	0.2213
TOTAL				11.21	7	0.1296

- In the model post-imputation, (m2) it looks like the impact of size for those without the treatment is much more modest than the impact of size when the treatment is received, according to the nomogram we saw earlier.
- In neither model does the effect of size or ses_group appear to meet the standard for statistical significance.
- The model discrimination is a bit better in the model with imputation and the interaction term. Specifically, the C statistic for m2, with imputation and interaction, is 0.633, as compared to 0.614 for the original model, but both are still weak.
- The Nagelkerke R-squared for m2 is 0.069, a bit larger than the 0.049 for the original model, though still weak.
- The appropriate conclusion, then, is that the addition of imputation and an interaction effect have had at most a modest impact on the conclusions of the model. It remains a weak model.