ST558 HomeWork 5

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Task 1 Conceptual questions

- 1. What is the purpose of using cross-validation when fitting a random forest model?
 - cross validation is to check the performance and generalization of the random forest model. Although they already reduce over fitting, cross validation make a more accurate guess of how the model will perform on unseen data
- 2. Describe the bagged tree algorithm.
 - Bagging is an ensemble technique that builds decision tress on different bootstrapped samples of an original dataset. each tree is created independently and the results are averages creating a more stable and accurate prediction.
- 3. What is meant by a general linear model?
 - A general linear model is a statistical model where the outcome is a linear combination of predictors. This includes, simple linear regression, multiple linear regression.

$$Y = X_B + e$$

- 4. When fitting a multiple linear regression model, what does adding an interaction term do? That is, what does it allow the model to do differently as compared to when it is not included in the model?
 - The interaction term allow for the model to check situation where one predictor depends on the value of another., It enables the model to check for more complex relationships between variables.
- 5. Why do we split our data into a training and test set?
 - Data is split into training and test sets to evaluate the model's ability to generalize. The model is trained on a portion of a set then used to see if that training holds up on unseen data. This prevents over fitting, and increases accuracy.

Task 2: Data Prep

Packages and Data

```
library(tidyverse)
library(tidymodels)
library(caret)
library(yardstick)
library(glmnet)
heart = as_tibble(read_csv("heart.csv"))
```

Question 1

summary(heart)

Age	Sex	${\tt ChestPainType}$	RestingBP	
Min. :28.00	Length:918	Length:918	Min. : 0.0	
1st Qu.:47.00	Class :character	Class :character	1st Qu.:120.0	
Median :54.00	Mode :character	Mode :character	Median:130.0	
Mean :53.51			Mean :132.4	
3rd Qu.:60.00			3rd Qu.:140.0	
Max. :77.00			Max. :200.0	
Cholesterol	FastingBS	RestingECG	MaxHR	
Min. : 0.0	Min. :0.0000	Length:918	Min. : 60.0	
1st Qu.:173.2	1st Qu.:0.0000	Class :character	1st Qu.:120.0	
Median :223.0	Median :0.0000	Mode :character	Median :138.0	
Mean :198.8	Mean :0.2331		Mean :136.8	
3rd Qu.:267.0	3rd Qu.:0.0000		3rd Qu.:156.0	
Max. :603.0	Max. :1.0000		Max. :202.0	
ExerciseAngina	Oldpeak	ST_Slope	HeartDisease	
Length:918	Min. :-2.600	00 Length:918	Min. :0.0000	
Class : characte:	r 1st Qu.: 0.000	00 Class :characte	er 1st Qu.:0.0000	
Mode :characte	r Median : 0.600	00 Mode :characte	er Median :1.0000	
	Mean : 0.88	74	Mean :0.5534	
	3rd Qu.: 1.500	00	3rd Qu.:1.0000	
	Max. : 6.200	00	Max. :1.0000	

- a. heart disease is being treated as a quantitative variable, it is a storing values as either 1 or 0.
- b. This does not make sense as we should eb treating the variables as categorical, as a yes or no response. While this does work in theory as a binary 1/0 response, it would be better to treat this as a facotrs with levels of yes or no, or T and F.

```
new_heart = heart %>%
  mutate(HeartDisease_FACTORED = factor(HeartDisease, levels = c(0,1), labels = c("N", "Y"))) %>%
  select(-ST_Slope, -HeartDisease)
head(new_heart)
```

A tibble: 6 x 11

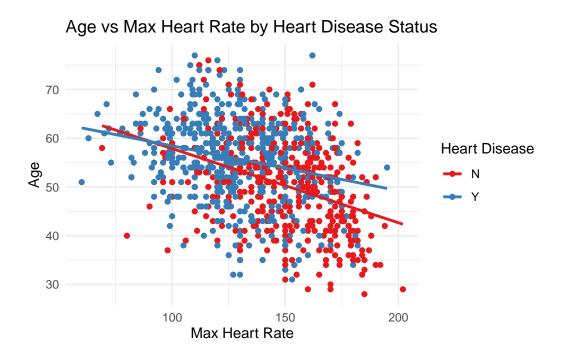
	Age	Sex	${\tt ChestPainType}$	RestingBP	${\tt Cholesterol}$	${\tt FastingBS}$	${\tt RestingECG}$	${\tt MaxHR}$
	<dbl></dbl>	<chr></chr>	<chr></chr>	<dbl></dbl>	<dbl></dbl>	<dbl></dbl>	<chr></chr>	<dbl></dbl>
1	40	M	ATA	140	289	0	Normal	172
2	49	F	NAP	160	180	0	Normal	156
3	37	M	ATA	130	283	0	ST	98
4	48	F	ASY	138	214	0	Normal	108
5	54	M	NAP	150	195	0	Normal	122
6	39	M	NAP	120	339	0	Normal	170

[#] i 3 more variables: ExerciseAngina <chr>, Oldpeak <dbl>,
HeartDisease_FACTORED <fct>

Task 3: EDA

```
new_heart %>%
    ggplot(
    aes(
        x = MaxHR,
        y = Age,
        colour = HeartDisease_FACTORED
    )
) +
geom_point() +
geom_smooth(method = "lm", se = F) +
scale_color_brewer(palette = "Set1") +
labs(
    title = "Age vs Max Heart Rate by Heart Disease Status",
    x = "Max Heart Rate",
    y = "Age",
    color = "Heart Disease"
) +
theme_minimal()
```

`geom_smooth()` using formula = 'y ~ x'



Question 2

An interaction model would be more accurate. The relationship between max heart rate and age differed depending on whether or not the person has heart disease. The have different slops indicating that max hr depends on heart disease status. This is algined with an interactive model not an additive model.

Task 4: Testing and Training

```
set.seed(101)
heart_split = initial_split(new_heart, prop = 0.8)
train = training(heart_split)
test = testing(heart_split)
```

Task 5: OLS and LASSO

Question 1

```
ols_mlr = lm(Age ~ MaxHR + HeartDisease_FACTORED + MaxHR*HeartDisease_FACTORED, data = train)
summary(ols_mlr)
Call:
lm(formula = Age ~ MaxHR + HeartDisease_FACTORED + MaxHR * HeartDisease_FACTORED,
   data = train)
Residuals:
    Min 1Q Median 3Q
-22.7703 -5.7966 0.4516 5.7772 20.6378
Coefficients:
                          Estimate Std. Error t value Pr(>|t|)
(Intercept)
                          75.58896 3.07510 24.581 < 2e-16 ***
MaxHR
                          HeartDisease_FACTOREDY
                         -8.58502 3.83433 -2.239 0.02546 *
MaxHR:HeartDisease_FACTOREDY 0.08343 0.02716 3.072 0.00221 **
Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
Residual standard error: 8.478 on 730 degrees of freedom
Multiple R-squared: 0.1839,
                            Adjusted R-squared: 0.1806
F-statistic: 54.84 on 3 and 730 DF, p-value: < 2.2e-16
```

Question 2

```
yardstick::rmse_vec(test$Age, predict(ols_mlr, test))
```

[1] 9.100206

```
LASSO_recipe = recipe(Age ~ MaxHR + HeartDisease_FACTORED, data = train) %>%
 step_dummy(all_nominal_predictors()) %>%
 step_normalize(all_predictors()) %>%
 step_interact(~ MaxHR:starts_with("HeartDisease_FACTORED"))
LASSO_recipe
-- Inputs
Number of variables by role
outcome:
predictor: 2
-- Operations
* Dummy variables from: all_nominal_predictors()
* Centering and scaling for: all_predictors()
* Interactions with: MaxHR:starts_with("HeartDisease_FACTORED")
Question 4
set.seed(101)
cv_folds = vfold_cv(train, v = 10)
lasso_model = linear_reg(penalty = tune(), mixture = 1) %>%
 set_engine("glmnet")
lasso_workflow = workflow() %>%
 add_model(lasso_model) %>%
 add_recipe(LASSO_recipe)
lasso_workflow
Preprocessor: Recipe
Model: linear_reg()
-- Preprocessor -----
3 Recipe Steps
* step_dummy()
```

```
* step_normalize()
* step_interact()
Linear Regression Model Specification (regression)
Main Arguments:
 penalty = tune()
 mixture = 1
Computational engine: glmnet
LASSO grid = lasso workflow %>%
  tune_grid(resamples = cv_folds, grid = grid_regular(penalty(), levels = 200))
LASSO_grid
# Tuning results
# 10-fold cross-validation
# A tibble: 10 x 4
   splits
                           .metrics
                                              .notes
   <list>
                    <chr> <chr>> <chr>>
                                               t>
 1 <split [660/74]> Fold01 <tibble [400 \times 5]> <tibble [0 \times 3]>
 2 <split [660/74]> Fold02 <tibble [400 x 5]> <tibble [0 x 3]>
 3 <split [660/74]> Fold03 <tibble [400 \times 5]> <tibble [0 \times 3]>
 4 <split [660/74] > Fold04 <tibble [400 x 5] > <tibble [0 x 3] >
 5 <split [661/73] > Fold05 <tibble [400 x 5] > <tibble [0 x 3] >
 6 <split [661/73]> Fold06 <tibble [400 \times 5]> <tibble [0 \times 3]>
7 <split [661/73]> Fold07 <tibble [400 x 5]> <tibble [0 x 3]>
 8 <split [661/73] > Fold08 <tibble [400 \times 5] > <tibble [0 \times 3] >
 9 <split [661/73] > Fold09 <tibble [400 \times 5] > <tibble [0 \times 3] >
10 <split [661/73] > Fold10 <tibble [400 x 5] > <tibble [0 x 3] >
LASSO_grid %>%
  collect_metrics() %>%
 filter(.metric == "rmse")
# A tibble: 200 x 7
   penalty .metric .estimator mean
                                         n std_err .config
      <dbl> <chr> <dbl> <int>
                                             <dbl> <chr>
       e-10 rmse
                    standard 8.47
                                        10
                                             0.124 Preprocessor1_Model001
 1 1
 2 1.12e-10 rmse
                    standard 8.47
                                        10 0.124 Preprocessor1_Model002
 3 1.26e-10 rmse
                    standard
                              8.47
                                        10
                                             0.124 Preprocessor1_Model003
 4 1.41e-10 rmse
                    standard
                              8.47
                                        10
                                             0.124 Preprocessor1_Model004
 5 1.59e-10 rmse
                                        10
                    standard
                              8.47
                                             0.124 Preprocessor1_Model005
 6 1.78e-10 rmse
                    standard
                              8.47
                                             0.124 Preprocessor1_Model006
7 2.00e-10 rmse
                               8.47
                    standard
                                        10
                                             0.124 Preprocessor1_Model007
                                             0.124 Preprocessor1_Model008
 8 2.25e-10 rmse
                    standard
                                8.47
                                        10
 9 2.52e-10 rmse
                                8.47
                                        10
                                             0.124 Preprocessor1_Model009
                    standard
10 2.83e-10 rmse
                    standard
                              8.47
                                        10
                                             0.124 Preprocessor1_Model010
# i 190 more rows
LASSO_grid %>%
  collect_metrics() %>%
 filter(.metric == "rmse") %>%
```

```
ggplot(aes(penalty, mean, color = .metric)) +
geom_line()
```

```
lowest_rmse <- LASSO_grid %>%
 select_best(metric = "rmse")
lowest_rmse
# A tibble: 1 x 2
     penalty .config
       <dbl> <chr>
1 0.0000000001 Preprocessor1_Model001
lasso_workflow %>%
 finalize_workflow(lowest_rmse)
Preprocessor: Recipe
Model: linear_reg()
-- Preprocessor ------
3 Recipe Steps
* step_dummy()
* step_normalize()
* step_interact()
-- Model -----
Linear Regression Model Specification (regression)
Main Arguments:
 penalty = 1e-10
 mixture = 1
```

Computational engine: glmnet

```
#fit it to the entire training set to see the model fit
LASSO_final <- lasso_workflow %>%
   finalize_workflow(lowest_rmse) %>%
   fit(train)

tidy(LASSO_final)
```

```
# A tibble: 4 x 3
term estimate penalty
<chr> <chr> < dbl> < dbl> < dbl>
1 (Intercept) 54.0 0.0000000001
2 MaxHR -3.08 0.0000000001
3 HeartDisease_FACTORED_Y 1.36 0.000000001
4 MaxHR_x_HeartDisease_FACTORED_Y 1.03 0.000000001
```

Question 5

I would expect the outputs to be generally the same between the OLS and LASSO models.

based on the tidy output of LASSO_final we can see that the LASSO model has the same predictors that the OLS model has. This would indiacte that RMSE would also be very similar.

Question 6

```
ols_mlr %>%
  predict(test) %>%
  rmse_vec(truth = test$Age)
```

[1] 9.100206

```
LASSO_final %>%
predict(test) %>%
pull() %>%
rmse_vec(truth = test$Age)
```

[1] 9.095981

Question 7

They are both using the same predictors, and their interactions are also the same and simple.

Task 6: Logistic regression

```
logrec1 <- recipe(HeartDisease_FACTORED ~ MaxHR, data = train) %>%
    step_normalize(MaxHR)

logrec2 <- recipe(HeartDisease_FACTORED ~ Age + Sex + ChestPainType + RestingBP, data = train) %>%
    step_normalize(all_numeric(), -HeartDisease_FACTORED) %>%
    step_dummy(all_nominal_predictors())
```

```
set.seed(101)
cv_repeats <- vfold_cv(train, 10, 5)</pre>
log_spec <- logistic_reg() %>%
 set_engine("glm")
LR1_wkf <- workflow() |>
add recipe(logrec1) |>
add_model(log_spec)
LR2_wkf <- workflow() |>
add_recipe(logrec2) |>
add_model(log_spec)
LR1 fit <- LR1 wkf |>
fit_resamples(cv_repeats, metrics = metric_set(accuracy, mn_log_loss))
LR2_fit <- LR2_wkf |>
fit_resamples(cv_repeats, metrics = metric_set(accuracy, mn_log_loss))
rbind(LR1_fit |> collect_metrics(),
LR2 fit |> collect metrics())|>
mutate(Model = c("Model1", "Model1", "Model2", "Model2")) |>
select(Model, everything())
# A tibble: 4 x 7
  Model .metric
                     .estimator mean
                                          n std_err .config
                      <chr> <dbl> <int>
  <chr> <chr>
                                               <dbl> <chr>
1 Model1 accuracy binary 0.677 50 0.00670 Preprocessor1_Model1 2 Model1 mn_log_loss binary 0.607 50 0.00550 Preprocessor1_Model1
                      binary 0.776
3 Model2 accuracy
                                           50 0.00676 Preprocessor1_Model1
4 Model2 mn_log_loss binary
                               0.490
                                           50 0.00913 Preprocessor1_Model1
mean(train$HeartDisease_FACTORED == "Y")
```

[1] 0.5694823

Model 2 was the better performing model it had the highest accuracy and the lowest loss.

```
final_model <- LR2_wkf %>%
  fit(data = train)

predictions <- predict(final_model, test, type = "class") %>%
  bind_cols(test)

confusionMatrix(
  data = predictions$.pred_class,
  reference = predictions$HeartDisease_FACTORED
)
```

Confusion Matrix and Statistics

Reference

 ${\tt Prediction} \quad {\tt N} \quad {\tt Y}$

N 71 17 Y 23 73

Accuracy : 0.7826

95% CI: (0.716, 0.8399)

No Information Rate : 0.5109 P-Value [Acc > NIR] : 2.577e-14

Kappa : 0.5656

Mcnemar's Test P-Value : 0.4292

Sensitivity : 0.7553
Specificity : 0.8111
Pos Pred Value : 0.8068
Neg Pred Value : 0.7604
Prevalence : 0.5109
Detection Rate : 0.3859
Detection Prevalence : 0.4783

Detection Prevalence : 0.4783 Balanced Accuracy : 0.7832

'Positive' Class : N

Question 3

The model's sensitivity was 0.7553: The model will miss about 24.5% of healthy patients. It will false flag almost 1/4 healthy people as diseased.

The model's specificity was 0.8111: The model correctly identified 81% of actual heart disease cases but will miss 19% of diseased patients.

The model is better at confirming positive cases than it is with ruling out disease.