# ST558\_HW5

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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 : 0.887	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.

#### Question 2

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

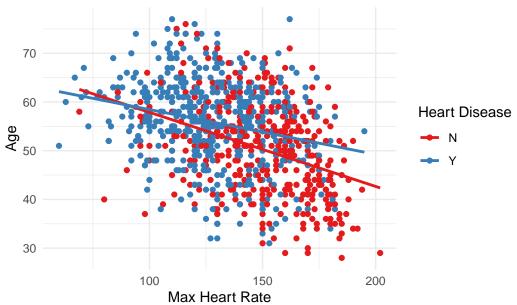
<sup>#</sup> i 3 more variables: ExerciseAngina <chr>, Oldpeak <dbl>,

<sup>#</sup> 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()
```

# Age vs Max Heart Rate by Heart Disease Status



<sup>`</sup>geom\_smooth()` using formula = 'y ~ x'

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

```
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
                                     Max
-22.7703 -5.7966 0.4516 5.7772 20.6378
Coefficients:
                          Estimate Std. Error t value Pr(>|t|)
                          75.58896 3.07510 24.581 < 2e-16 ***
(Intercept)
                          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 '**' 0.05 '.' 0.1 ' ' 1
Residual standard error: 8.478 on 730 degrees of freedom
                            Adjusted R-squared: 0.1806
Multiple R-squared: 0.1839,
F-statistic: 54.84 on 3 and 730 DF, p-value: < 2.2e-16
```

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

#### Question 3

[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
```

- -- Recipe ------
- -- Inputs

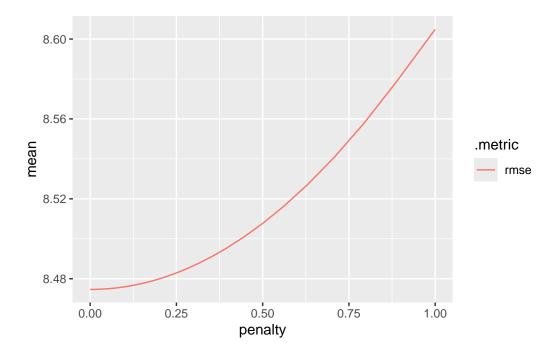
Number of variables by role

outcome: 1
predictor: 2

- -- Operations
- \* Dummy variables from: all\_nominal\_predictors()
- \* Centering and scaling for: all\_predictors()
- \* Interactions with: MaxHR:starts\_with("HeartDisease\_FACTORED")

```
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()
-- Model -----
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
                      id
                              .metrics
                                                   .notes
   st>
                      <chr> <chr>> <chr>>
                                                    st>
 1 <split [660/74] > Fold01 <tibble [400 \times 5] > <tibble [0 \times 3] >
 2 < \text{split} [660/74] > \text{Fold02} < \text{tibble} [400 x 5] > < \text{tibble} [0 x 3] >
 3 \left(\frac{660}{74}\right) Fold03 \left(\frac{400 \times 5}{5}\right) \left(\frac{660}{74}\right) Fold03 \left(\frac{400 \times 5}{5}\right)
 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 \left(\frac{661}{73}\right) Fold06 \left(\frac{400 \times 5}{5}\right) \left(\frac{661}{73}\right)
 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 x 5] > <tibble [0 x 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>
                      <chr> <dbl> <int>
                                                   <dbl> <chr>
       e-10 rmse
                                   8.47
                                                  0.124 Preprocessor1_Model001
 1 1
                      standard
                                             10
 2 1.12e-10 rmse
                      standard
                                 8.47
                                             10
                                                  0.124 Preprocessor1_Model002
                                 8.47
 3 1.26e-10 rmse
                      standard
                                                  0.124 Preprocessor1_Model003
                                             10
 4 1.41e-10 rmse
                      standard
                                 8.47
                                             10
                                                  0.124 Preprocessor1_Model004
 5 1.59e-10 rmse
                      standard
                                 8.47
                                             10
                                                  0.124 Preprocessor1_Model005
 6 1.78e-10 rmse
                      standard
                                   8.47
                                             10
                                                  0.124 Preprocessor1_Model006
 7 2.00e-10 rmse
                                 8.47
                                             10
                                                  0.124 Preprocessor1_Model007
                      standard
 8 2.25e-10 rmse
                                   8.47
                                                  0.124 Preprocessor1_Model008
                      standard
                                             10
 9 2.52e-10 rmse
                                    8.47
                                                  0.124 Preprocessor1_Model009
                      standard
                                             10
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
```

```
lasso_workflow %>%
  finalize_workflow(lowest_rmse)
```

Preprocessor: Recipe
Model: linear\_reg()

- -- Preprocessor ------
- 3 Recipe Steps
- \* step\_dummy()
- \* step\_normalize()

#	A tibble: 4 x 3		
	term	${\tt estimate}$	penalty
	<chr></chr>	<dbl></dbl>	<dbl></dbl>
1	(Intercept)	54.0	0.000000001
2	MaxHR	-3.08	0.000000001
3	<pre>HeartDisease_FACTORED_Y</pre>	1.36	0.000000001
4	MaxHR_x_HeartDisease_FACTORED_Y	1.03	0.000000001

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.

```
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)

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> <chr>
                  <chr> <dbl> <int> <dbl> <chr>
1 Model1 accuracy binary
                            0.677 50 0.00670 Preprocessor1_Model1
2 Model1 mn_log_loss binary
                                       50 0.00550 Preprocessor1_Model1
                              0.607
3 Model2 accuracy
                                       50 0.00676 Preprocessor1_Model1
                   binary
                              0.776
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

Prediction N 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

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.