7.1.1 Generalized Additive Models Stat 5100: Dr. Bean

Example 1: Baseball Dataset from 4.1.1

Let's see if we can improve upon the penalized linear regression model to predict the log of salary for professional (non-pitcher) baseball players. Note that answers will differ slightly depending on the random seed set.

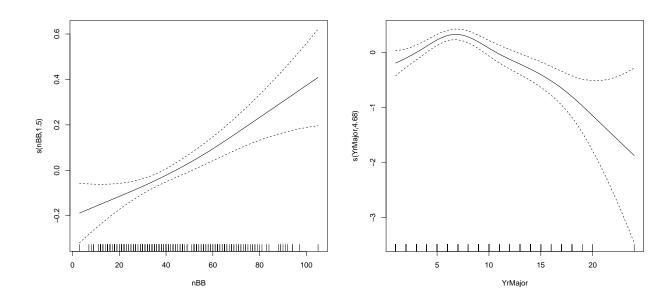
```
# Set a random seed for reproducibility
set.seed(830578)
# Load data
library(stat5100)
library(mgcv)
## Loading required package: nlme
## This is mgcv 1.8-33. For overview type 'help("mgcv-package")'.
data(baseball)
baseball_gam_all <-</pre>
 mgcv::gam(logSalary ~ s(nAtBat) + s(nHits) + s(nHome) +
            s(nRuns) + s(nRBI) + s(nBB) + s(YrMajor) +
            s(CrAtBat) + s(CrHits) + s(CrHome) + s(CrRuns) +
            s(CrRbi) + s(CrBB) + s(nOuts) + s(nAssts) +
            s(nError) + League + Division,
                       data = baseball)
summary(baseball_gam_all)
## Family: gaussian
## Link function: identity
## Formula:
## logSalary ~ s(nAtBat) + s(nHits) + s(nHome) + s(nRuns) + s(nRBI) +
##
      s(nBB) + s(YrMajor) + s(CrAtBat) + s(CrHits) + s(CrHome) +
##
      s(CrRuns) + s(CrRbi) + s(CrBB) + s(nOuts) + s(nAssts) + s(nError) +
      League + Division
##
## Parametric coefficients:
                 Estimate Std. Error t value Pr(>|t|)
##
## (Intercept)
                ## LeagueNational 0.07437
                            0.04414 1.685 0.0936 .
## DivisionWest -0.02042
                            0.04346 -0.470 0.6389
## ---
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
```

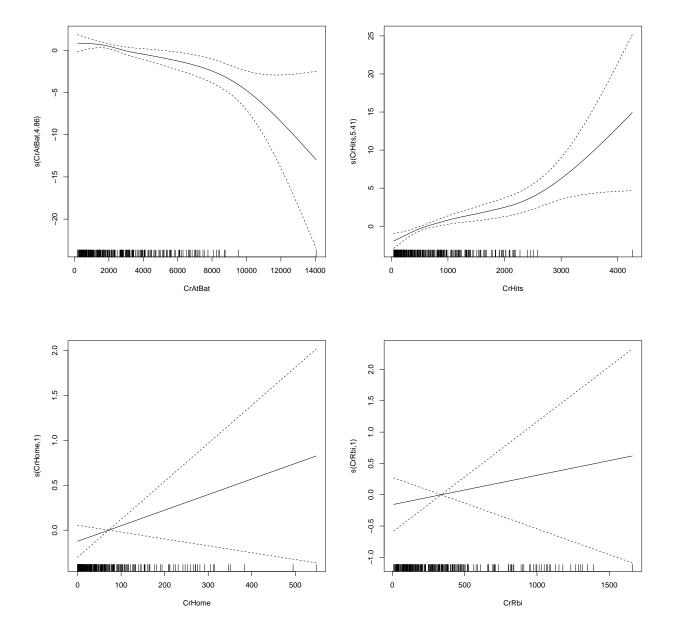
```
## Approximate significance of smooth terms:
##
                edf Ref.df
                               F p-value
## s(nAtBat) 3.470
                    4.373 1.312 0.25485
## s(nHits)
              1.000
                    1.000
                           0.130 0.71892
## s(nHome)
              4.819
                     5.884
                            1.259 0.25922
## s(nRuns)
              4.174
                     5.178
                            1.223 0.29111
## s(nRBI)
              1.971
                     2.551
                           0.975 0.42890
## s(nBB)
              1.245
                     1.445
                           3.885 0.04937 *
## s(YrMajor) 5.315
                     6.270 10.713 < 2e-16 ***
## s(CrAtBat) 7.385
                     8.063
                            2.822 0.00894 **
## s(CrHits)
             3.895
                     4.930
                           2.157 0.05747 .
## s(CrHome)
             8.094
                     8.510
                           3.217 0.00128 **
## s(CrRuns)
              1.000
                     1.000
                            0.079 0.77944
## s(CrRbi)
              8.447
                     8.770
                            2.961 0.00632 **
## s(CrBB)
              1.000
                    1.000
                           0.138 0.71063
## s(nOuts)
              5.957
                     7.070
                           1.982 0.05850
                            0.325 0.56924
## s(nAssts)
             1.000
                    1.000
## s(nError)
             3.601 4.463
                           1.224 0.25678
##
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
## R-sq.(adj) = 0.882
                         Deviance explained = 91.1%
## GCV = 0.12437 Scale est. = 0.093457 n = 263
```

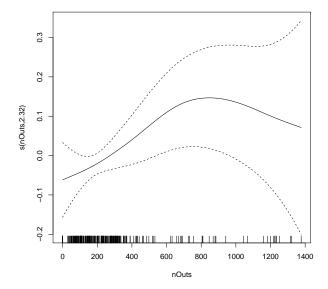
Now, let's refit the models but only using the significant terms:

We can take a look at the estimated spline functions for each of the predictor variables. In each of the below plots, the x-axis contains the various levels of the predictor variables. On the y-axis, we see the estimated spline function (keep in mind that these are multiple different polynomial functions being concatenated together). Along the x-axis you will see little notches: these each indicate the unique points that went into creating the spline.

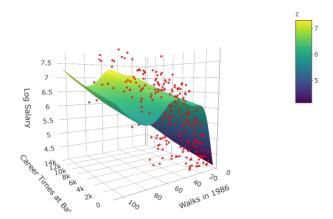
```
mgcv::plot.gam(baseball_gam, scale = 0)
```







For simplicity, if we fit a GAM with just CrAtBat and nBB (like we did in the LOESS example), then we get the following surface plot:



This plot is comparable to the plot from the LOESS example in 4.3.1. Note that this plot uses an interactive plot feature available in the plotly package of R.

Example 2: Diabetes Dataset

The Pima Indians Diabetes dataset is a dataset from the National Institute of Diabetes and Digestive and Kidney Diseases. Our goal here is to predict whether or not a patient has diabetes. In this dataset, all patients are females that are at least 21 and are of Pima Indian heritage.

Let's split our data into a training and testing dataset and see how well we do on the testing dataset by training on the training dataset.

```
data("diabetes")
head(diabetes)
## Pregnancies Glucose BloodPressure SkinThickness Insulin BMI
```

```
## 1
               6
                     148
                                     72
                                                    35
                                                             0 33.6
## 2
                                                    29
               1
                      85
                                     66
                                                             0 26.6
                                                    0
## 3
               8
                     183
                                     64
                                                             0 23.3
## 4
               1
                      89
                                     66
                                                   23
                                                            94 28.1
## 5
               0
                     137
                                     40
                                                   35
                                                          168 43.1
                                     74
                                                            0 25.6
## 6
               5
                     116
                                                    0
## DiabetesPedigreeFunction Age Outcome
                        0.627 50
                        0.351 31
## 2
                                         \cap
## 3
                        0.672 32
                                         1
## 4
                        0.167 21
                                         \cap
## 5
                        2.288 33
                        0.201 30
## 6
                                         \cap
# How many observations are there?
nrow(diabetes)
## [1] 768
# Create a training and testing split with 80% training data
train_index <- sample(1:nrow(diabetes), size = 0.80*nrow(diabetes))</pre>
diabetes_train <- diabetes[train_index, ]</pre>
diabetes_test <- diabetes[-train_index, ]</pre>
diabetes_gam <- mgcv::gam(Outcome ~ s(Pregnancies) + s(Glucose) + s(BloodPressure) +
                            s(SkinThickness) + s(Insulin) + s(BMI) +
                            s(DiabetesPedigreeFunction) + s(Age), family = "binomial",
                         data = diabetes_train)
```

Now let's see how accurate we are on the testing dataset:

```
# Here are the predicted class probabilities
test_class_prob <- predict(diabetes_gam, diabetes_test, type = "response")
# If the probability is higher than 50% of having diabetes, mark it as a 1.
pred_class <- rep(0, nrow(diabetes_test))
pred_class[test_class_prob > 0.50] <- 1

# Now that we have our predicted class, let's get some statistics on our accuracy.
total_test <- nrow(diabetes_test)
total_correct <- sum(pred_class == diabetes_test$Outcome)

# Error rate
(total_test - total_correct) / total_test

## [1] 0.1948052

# Successful prediction rate
total_correct / total_test
## [1] 0.8051948</pre>
```