# Logistic Regression on the Plasma Data

## Karen Mazidi

Logistic regression example using the plasma data set in package HSAUR.

### Data exploration

library(HSAUR)

We can read more about the plasma data set by typing "?plasma" at the console, after package HSAUR is loaded. We want to learn to predict ESR > 20 or not, based on the levels of the plasma proteins fibrinogen and globulin. ESR stands for erythrocyte sedimentation rate, the rate at which red blood cells settle in blood plasma. Values > 20 indicate some possible associations with various health conditions.

```
## Loading required package: tools
attach(plasma)
str(plasma)
                    32 obs. of 3 variables:
  'data.frame':
   $ fibrinogen: num 2.52 2.56 2.19 2.18 3.41 2.46 3.22 2.21 3.15 2.6 ...
  $ globulin : int 38 31 33 31 37 36 38 37 39 41 ...
                : Factor w/ 2 levels "ESR < 20", "ESR > 20": 1 1 1 1 1 1 1 1 1 1 ...
   $ ESR
head(plasma)
##
     fibrinogen globulin
                              ESR
## 1
           2.52
                      38 ESR < 20
## 2
           2.56
                      31 ESR < 20
                      33 ESR < 20
## 3
           2.19
           2.18
                      31 ESR < 20
## 4
## 5
           3.41
                      37 ESR < 20
                      36 ESR < 20
           2.46
attach(plasma)
## The following objects are masked from plasma (pos = 3):
##
##
       ESR, fibrinogen, globulin
```

#### Plot the data

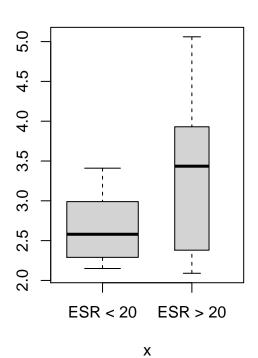
The first pair of plots show us that observations where ESR>20 are rarer. This is indicated by the thinner boxes because we set varwidth=TRUE in the bloxplot call. More importantly, the boxplots show that ESR>20 observations are associated with slightly higher levels of globulin and significantly higher levels of fibronogen.

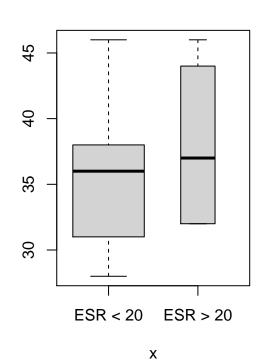
The second set of pots are conditional density plots. We can make the same observations as the box plots. Here they are just visualized differently. The total probability space is the rectangle, with the lighter grey indicating ESR>20.

```
par(mfrow=c(1,2))
plot(ESR, fibrinogen, main="Fibrinogen", ylab="", varwidth=TRUE)
plot(ESR, globulin, main="Globulin", ylab="", varwidth=TRUE)
```

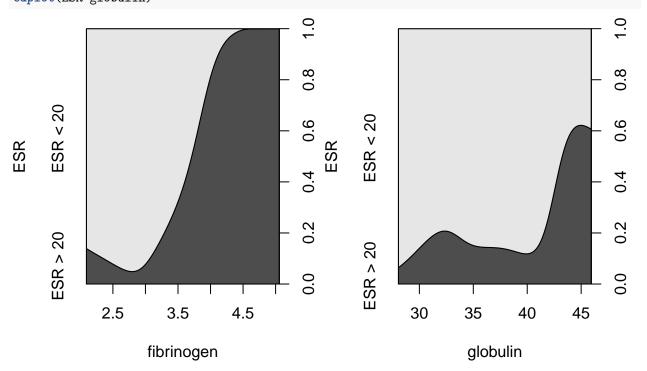


## Globulin





par(mfrow=c(1,2))
cdplot(ESR~fibrinogen)
cdplot(ESR~globulin)



### Train and test sets

Even though our data is small, we will go ahead and divide it into train and test sets.

```
set.seed(3)
i <- sample(1:nrow(plasma), 0.75*nrow(plasma), replace=FALSE)
train <- plasma[i,]
test <- plasma[-i,]</pre>
```

#### Build a logistic regression model

Our first model uses only fibronogen as a predictor. The glm() function is used for logistic regression, with parameter family=binomial

The summary is a little different for logistic regression compared to linear regression: \* the residual are deviance residuals - measures of deviance contributed from each observation \* the coefficients represent changes in the log odds of y \* model metrics

```
glm1 <- glm(ESR~fibrinogen, data=train, family=binomial)
summary(glm1)</pre>
```

```
##
## Call:
## glm(formula = ESR ~ fibrinogen, family = binomial, data = train)
##
## Deviance Residuals:
                      Median
##
      Min
                 1Q
                                   3Q
                                           Max
  -0.8197 -0.5334 -0.3672 -0.2551
                                        2.7268
##
## Coefficients:
##
              Estimate Std. Error z value Pr(>|z|)
                 -8.087
## (Intercept)
                             3.663
                                   -2.208
                                             0.0273 *
## fibrinogen
                  2.102
                             1.133
                                     1.855
                                             0.0635 .
## ---
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
##
##
  (Dispersion parameter for binomial family taken to be 1)
##
##
       Null deviance: 21.627
                             on 23
                                    degrees of freedom
## Residual deviance: 15.740 on 22 degrees of freedom
## AIC: 19.74
## Number of Fisher Scoring iterations: 5
```

#### **Evaluate**

Our first model uses only fibronogen as a predictor. On our small test data we got about 88% accuracy. The table shows that all test observations were predicted as not ESR>20 and one of the 8 observations actually was ESR>20. Internally, the ESR>20 factor is coded as 1 for not >20 and 2 for ESR>20. This is why we compare them as integer().

```
probs <- predict(glm1, newdata=test, type="response")
pred <- ifelse(probs>0.5, 2, 1)
acc1 <- mean(pred==as.integer(test$ESR))
print(paste("glm1 accuracy = ", acc1))</pre>
```

```
## [1] "glm1 accuracy = 0.75"
```

```
table(pred, as.integer(test$ESR))
##
## pred 1 2
## 1 6 2
```

#### What does it mean?

Let's explore the meaning of the coefficient.

```
fibro <- glm1$coefficients[2]
intercept <- glm1$coefficients[1]

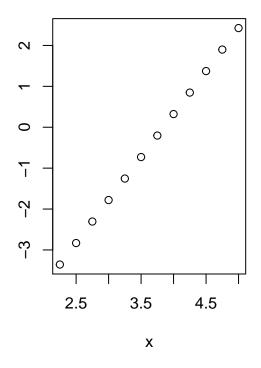
log_odds <- function(x, fibro, intercept){
  intercept + fibro * x
}

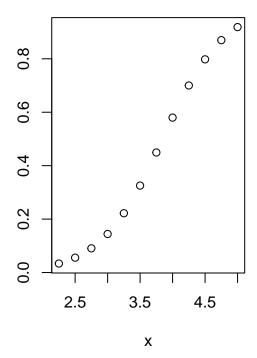
x <- seq(from=2.25, to=5.0, by=0.25)
y <- log_odds(x, fibro, intercept)
par(mfrow=c(1,2))
plot(x,y, main="log odds", ylab="")

prob <- exp(y) / (1 + exp(y))
plot(x, prob, main="probabilities", ylab="")</pre>
```

# log odds

## probabilities





## Build another model

This model uses both predictors. On the test set we got the same accuracy.

```
glm2 <- glm(ESR~fibrinogen+globulin, data=train, family=binomial)
summary(glm2)</pre>
```

```
##
## Call:
## glm(formula = ESR ~ fibrinogen + globulin, family = binomial,
##
       data = train)
##
## Deviance Residuals:
                         Median
                   10
                                       30
                                                 Max
## -0.99591 -0.50791 -0.17843 -0.08106
                                             2.26810
##
## Coefficients:
               Estimate Std. Error z value Pr(>|z|)
## (Intercept) -20.4969
                           10.2436 -2.001
                                             0.0454 *
## fibrinogen
                 2.6476
                            1.5488
                                     1.709
                                             0.0874 .
## globulin
                                     1.562
                 0.2834
                            0.1814
                                             0.1182
## ---
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
##
## (Dispersion parameter for binomial family taken to be 1)
##
##
       Null deviance: 21.627 on 23 degrees of freedom
## Residual deviance: 12.308 on 21 degrees of freedom
## AIC: 18.308
##
## Number of Fisher Scoring iterations: 6
probs <- predict(glm1, newdata=test, type="response")</pre>
pred <- ifelse(probs>0.5, 2, 1)
acc2 <- mean(pred==as.integer(test$ESR))</pre>
print(paste("glm2 accuracy = ", acc2))
## [1] "glm2 accuracy = 0.75"
table(pred, as.integer(test$ESR))
##
## pred 1 2
      1 6 2
```

### Compare the models with anova()

The second model is only slightly better than the first. The residuals dropped by only 2 points, and the p-value is not small.

```
anova(glm1, glm2)

## Analysis of Deviance Table

##
## Model 1: ESR ~ fibrinogen

## Model 2: ESR ~ fibrinogen + globulin

## Resid. Df Resid. Dev Df Deviance

## 1 22 15.740

## 2 21 12.308 1 3.4322
```