# Modele regresji i ich zastosowania Labolatoria 9 i 10

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25 maja 2021

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## 1 Zadanie 1

## 1.1 Podpunkt a

Zaczynamy od podzielenia zbioru na część uczącą i część testową w proporcji 70:30. Sprawdzamy na koniec podobieństwo frakcji STAN w zbiorach.

```
logistyczna <- read.delim("~/Downloads/logistyczna.txt", header=TRUE, comment.char="#")</pre>
head(logistyczna)
##
     STAN RMS10 RMS20 RMS30 PWD
                                      Α
                                          DD
                                                   YDD
## 1
          0.06 0.06 0.08 1.02 112.07 0.22 44.72 0.21 0.19
## 2
        0 0.03 0.06 0.06 1.20 116.90 0.25 93.07 0.17 0.23
## 3
        0 0.05
                0.05 0.08 1.23 155.88 0.47 42.79 0.29 0.85
        0 0.04
                 0.05 0.09 1.33 124.59 0.19 44.78 0.27 0.45
## 5
        0 0.05
                0.06  0.06  1.16  133.42  0.17  40.72  0.35  0.28
        0 0.09 0.13 0.13 1.07 136.54 0.20 55.10 0.17 0.39
## 6
```

```
indices <- sort(sample(1:267, round(0.7 * 267)))

logistyczne_train<-logistyczna[indices,]
logistyczne_test<-logistyczna[-indices,]

sum(logistyczna$STAN)

## [1] 232

sum(logistyczne_train$STAN)

## [1] 164

sum(logistyczne_test$STAN)

## [1] 68</pre>
```

#### 1.2 Podpunkt b

Konstruujemy model regresji logistycznej oparty na wszystkich zmiennych, jakie mamy do dyspozycji.

```
model1<-glm(logistyczne_train$STAN~.,data=logistyczne_train, family = binomial(link = "]</pre>
summary(model1)
##
## Call:
## glm(formula = logistyczne_train$STAN ~ ., family = binomial(link = "logit"),
      data = logistyczne_train)
##
## Deviance Residuals:
       Min
                  1Q
                       Median
                                      30
                                               Max
## -3.10332
            0.08382
                       0.21015
                                0.44111
                                          1.40250
##
## Coefficients:
##
                Estimate Std. Error z value Pr(>|z|)
## (Intercept) -23.898454
                          5.740621 -4.163 3.14e-05 ***
## RMS10
               28.017566 23.368747
                                      1.199
                                              0.2306
## RMS20
              -14.150810 24.794920 -0.571
                                              0.5682
## RMS30
               15.424945 17.206585 0.896
                                              0.3700
## PWD
                          3.207363 4.052 5.09e-05 ***
               12.994665
## A
                0.025891 0.015243 1.699 0.0894 .
               22.046541 9.720716 2.268
## DD
                                              0.0233 *
               -0.001278 0.024409 -0.052
                                              0.9582
## YA
                                    0.117
## YDD
                0.600599 5.114981
                                              0.9065
## AR
                4.670468
                           2.561452
                                    1.823
                                              0.0682 .
## ---
```

```
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
##
## (Dispersion parameter for binomial family taken to be 1)
##
## Null deviance: 139.446 on 186 degrees of freedom
## Residual deviance: 98.544 on 177 degrees of freedom
## AIC: 118.54
##
## Number of Fisher Scoring iterations: 7
```

#### 1.3 Podpunkt c

```
library(lmtest)
## Loading required package: zoo
##
## Attaching package: 'zoo'
## The following objects are masked from 'package:base':
##
##
       as.Date, as.Date.numeric
model2<-glm(logistyczne_train$STAN~1,data=logistyczne_train, family = binomial(link = "]
summary(model2)
##
## Call:
## glm(formula = logistyczne_train$STAN ~ 1, family = binomial(link = "logit"),
     data = logistyczne_train)
##
##
## Deviance Residuals:
     Min 1Q Median
                                  3Q
                                         Max
## -2.0473 0.5123 0.5123 0.5123
                                      0.5123
##
## Coefficients:
              Estimate Std. Error z value Pr(>|z|)
## (Intercept) 1.9644
                         0.2227 8.822
                                          <2e-16 ***
## ---
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
##
## (Dispersion parameter for binomial family taken to be 1)
      Null deviance: 139.45 on 186 degrees of freedom
## Residual deviance: 139.45 on 186 degrees of freedom
## AIC: 141.45
##
## Number of Fisher Scoring iterations: 4
```

```
lrtest(model1,model2)
## Likelihood ratio test
## Model 1: logistyczne_train$STAN ~ RMS10 + RMS20 + RMS30 + PWD + A + DD +
      YA + YDD + AR
## Model 2: logistyczne_train$STAN ~ 1
    #Df LogLik Df Chisq Pr(>Chisq)
## 1 10 -49.272
     1 -69.723 -9 40.901 5.213e-06 ***
## ---
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
# intercept_only <- lm(Y ~ 1, data=mydata_2)</pre>
# #define model with all predictors
# all <- lm(Y ~~., data=mydata_2)
# #perform forward stepwise regression
# forward <- step(model2, direction='forward', scope=formula(model1), trace=0)
# #view results of forward stepwise regression
# forward£anova
#
# #view final model
# forward£coefficients
# summary(forward)
```

- P-value wynosi 0.0004606 i jest ona mniejsza od poziomu istotności na poziomie 0.05
- Poprzez oszacowanie statystyki jesteśmy w stanie odrzucić hipotezę zerową.
- Odrzucenie testu ma sens, widzieliśmy duże znaczenie na zmienną STAN przez atrybut PWD.

Poprzez oszacowanie statystyki jesteśmy w stanie odrzucić hipotezę zerową.

#### 1.4 Podpunkt d

Liczymy wartości estymatorów i standardowych błędów predyktorów z wyjściowego modelu.

```
## (Intercept) RMS10 RMS20 RMS30 PWD
## -23.898453997 28.017565506 -14.150810109 15.424944737 12.994664971
## A DD YA YDD AR
## 0.025891494 22.046541090 -0.001278148 0.600598986 4.670468404

sqrt(diag(summary(model1)$cov.unscaled)*summary(model1)$dispersion)
```

```
## (Intercept) RMS10 RMS20 RMS30 PWD A
## 5.74062068 23.36874667 24.79491970 17.20658543 3.20736292 0.01524286
## DD YA YDD AR
## 9.72071648 0.02440914 5.11498087 2.56145217
```

Największe wartości błędów dotyczą zmiennych RMS10, RMS20 i RMS30.

```
confint(model1, level = 0.95)
## Waiting for profiling to be done...
##
                     2.5 %
                                97.5 %
## (Intercept) -36.455414711 -13.70139645
## RMS10
             -16.853960385 76.03649746
             -66.368850297 33.56924761
## RMS20
## RMS30
             -16.620839528 52.97215903
## PWD
               7.257806160 19.95021953
## A
              -0.003180568 0.05732016
## DD
               4.944941093 42.86830994
## YA
              -0.048136357 0.04859228
## YDD
              -9.381964968 10.88636616
## AR
              -0.006619474 10.09161433
summary(model1)
##
## Call:
## glm(formula = logistyczne_train$STAN ~ ., family = binomial(link = "logit"),
     data = logistyczne_train)
##
## Deviance Residuals:
       Min
                 1Q
                       Median
                                    3Q
                                             Max
                      0.21015
                                         1.40250
## -3.10332
             0.08382
                               0.44111
##
## Coefficients:
                Estimate Std. Error z value Pr(>|z|)
## (Intercept) -23.898454 5.740621 -4.163 3.14e-05 ***
## RMS10
              28.017566 23.368747
                                   1.199
                                          0.2306
## RMS20
              -14.150810 24.794920 -0.571
                                            0.5682
## RMS30
              15.424945 17.206585 0.896 0.3700
## PWD
              12.994665 3.207363 4.052 5.09e-05 ***
## A
               0.025891 0.015243 1.699 0.0894 .
## DD
               22.046541 9.720716 2.268 0.0233 *
## YA
              ## YDD
               0.600599
                        5.114981 0.117
                                          0.9065
## AR.
               4.670468 2.561452 1.823 0.0682 .
## ---
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
##
```

```
## (Dispersion parameter for binomial family taken to be 1)
##

## Null deviance: 139.446 on 186 degrees of freedom
## Residual deviance: 98.544 on 177 degrees of freedom
## AIC: 118.54
##

## Number of Fisher Scoring iterations: 7
```

- Hipoteza zerowa  $\beta_i = 0$  jest odrzucana tylko dla i = 5
- Zmienna PWD ma zdecydowanie największy wpływ na zmienną STAN, dla pozostałych zmiennych hipoteza zerowa mogłaby być przyjęta. Poziom istotności byl wyższy od 0.05

#### 1.5 Podpunkt e

Konstruujemy *classification table* i wyznaczamy parametry charakteryzujące zdolności predykcyjne modelu na dwóch poziomach odcięcia 0.5 i 0.7

```
library(knitr)
n=length(logistyczne_test$STAN)
real_values <- logistyczne_test$STAN
pred_values <- predict(model1, newdata = logistyczne_test)</pre>
class_tab <- table(real_values, pred_values>0.5)
true_negative <- class_tab[1,1]</pre>
false_positive <- class_tab[1,2]</pre>
false_negative <- class_tab[2,1]</pre>
true_positive <- class_tab[2,2]</pre>
pasteO("Sensivity for pi 0.5: ", round ((true_positive/(true_positive+false_negative)) >
## [1] "Sensivity for pi 0.5: 89.70588"
paste0("Specifity for pi 0.5: ", round ((true_negative/(true_negative+false_positive)) >
## [1] "Specifity for pi 0.5: 8.33333"
pasteO("False positive rate for pi 0.5: ", round (false_positive/(false_positive+true_ne
## [1] "False positive rate for pi 0.5: 0.91667"
pasteO("False negative rate for pi 0.5: ", round (false_negative/(false_negative+true_negative)
## [1] "False negative rate for pi 0.5: 0.875"
 pasteO("Overall proportion of correct classifications for pi 0.5: ", round ((true_negations))
## [1] "Overall proportion of correct classifications for pi 0.5: 0.775"
kable(class_tab)
```

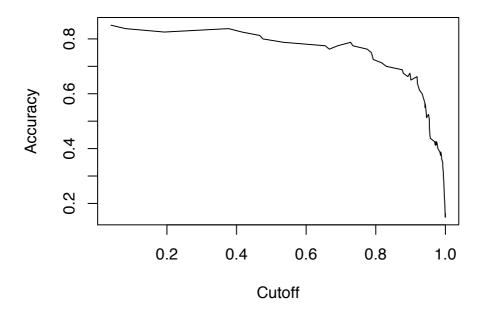
	FALSE	TRUE
0	1	11
1	7	61

```
n=length(logistyczne_test$STAN)
real_values <- logistyczne_test$STAN
pred_values <- predict(model1, newdata = logistyczne_test)</pre>
class_tab <- table(real_values, pred_values>0.7)
true_negative <- class_tab[1,1]</pre>
false_positive <- class_tab[1,2]</pre>
false_negative <- class_tab[2,1]</pre>
true_positive <- class_tab[2,2]</pre>
paste0("Sensivity for pi 0.7: ", round ((true_positive/(true_positive+false_negative)) >
## [1] "Sensivity for pi 0.7: 88.23529"
paste0("Specifity for pi 0.7: ", round ((true_negative/(true_negative+false_positive)) >
## [1] "Specifity for pi 0.7: 16.66667"
pasteO("False positive rate for pi 0.7: ", round (false_positive/(false_positive+true_ne
## [1] "False positive rate for pi 0.7: 0.83333"
pasteO("False negative rate for pi 0.7: ", round (false_negative/(false_negative+true_negative)
## [1] "False negative rate for pi 0.7: 0.8"
pasteO("Overall proportion of correct classifications for pi 0.7: ", round ((true_negations))
## [1] "Overall proportion of correct classifications for pi 0.7: 0.775"
kable(class_tab)
```

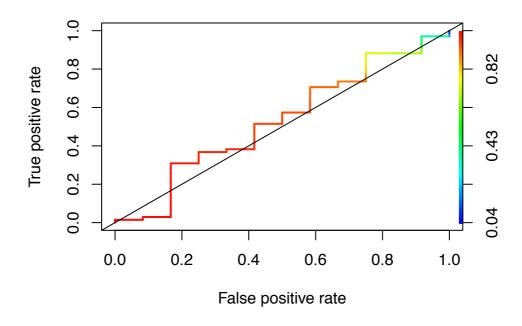
	FALSE	TRUE
0	2	10
1	8	60

### 1.6 Podpunkt f

```
library(ROCR)
pred = predict(model1,logistyczne_test,type="response")
pred = prediction(pred, logistyczne_test$STAN)
perf = performance(pred, "acc")
plot(perf)
```



```
roc = performance(pred, "tpr", "fpr")
plot(roc, colorize = T, lwd = 2)
abline(a = 0, b = 1)
```



```
library(pROC)
## Type 'citation("pROC")' for a citation.
```

```
##
## Attaching package: 'pROC'
## The following objects are masked from 'package:stats':
##
## cov, smooth, var

pred = predict(model1,logistyczne_test,type="response")
auc<-auc(logistyczne_test$STAN, pred)
## Setting levels: control = 0, case = 1
## Setting direction: controls < cases
auc
## Area under the curve: 0.5306
2*auc-1
## [1] 0.06127451</pre>
```

- Rysunek krzywej ROC wskazuje, że model na prawie każdym poziomie przewyższa skuteczność 0.5 co oznacza, że ma lepszą skuteczność dopasowania niż klasyfikator losowy.
- Pole które jest zakreślone pod krzywą ROC jest nazywane parametrem AUC. Wynosi on 0.6299 i jest powyżej 0.5 stąd lepsze skuteczność niż klasyfikatora losowego, daleko jesdnak do klasyfikatora idealnego -poziom AUC=1
- Współczynnik Giniego: GC = 2\*AUC-1 (wyższość klasyfikatora nad losowym)
- Współczynnik Giniego:0.2598039

#### 2 Zadanie 2

#### 2.1 Podpunkt a

```
library(readxl)
cancer <- read_excel("/Users/jansolarz/Downloads/cancer remission.xlsx")

## New names:
## * '' -> ...8
## * '' -> ...9
## * '' -> ...10
## * '' -> ...11

cancer<-cancer[,1:7]</pre>
```

#### 2.2 Podpunkt b

Zgodnie z metodą forward zaczynamy do przyłączania pojedynczych zmiennych niezależnych sprawdzając wysokość poziomu istotności zgodnie z zadanymi poziomami w zadaniu.

```
model_1<-glm(cancer$remission~1,data=cancer, family = binomial(link = "logit"))</pre>
model_2<-glm(cancer$remission~cancer$cell,data=cancer, family = binomial(link = "logit")</pre>
lrtest(model_1,model_2)
## Likelihood ratio test
## Model 1: cancer$remission ~ 1
## Model 2: cancer$remission ~ cancer$cell
## #Df LogLik Df Chisq Pr(>Chisq)
## 1 1 -17.186
## 2 2 -15.896 1 2.58
                              0.1082
model_3<-glm(cancer$remission~cancer$smear,data=cancer, family = binomial(link = "logit")</pre>
lrtest(model_1,model_3)
## Likelihood ratio test
##
## Model 1: cancer$remission ~ 1
## Model 2: cancer$remission ~ cancer$smear
    #Df LogLik Df Chisq Pr(>Chisq)
## 1 1 -17.186
## 2
      2 -16.640 1 1.0921
model_4<-glm(cancer$remission~cancer$infil,data=cancer, family = binomial(link = "logit")</pre>
lrtest(model_1,model_4)
## Likelihood ratio test
##
## Model 1: cancer$remission ~ 1
## Model 2: cancer$remission ~ cancer$infil
## #Df LogLik Df Chisq Pr(>Chisq)
## 1 1 -17.186
## 2 2 -16.201 1 1.9698
                               0.1605
model_5<-glm(cancer$remission~cancer$li,data=cancer, family = binomial(link = "logit"))</pre>
lrtest(model_1,model_5)
## Likelihood ratio test
## Model 1: cancer$remission ~ 1
## Model 2: cancer$remission ~ cancer$li
    #Df LogLik Df Chisq Pr(>Chisq)
## 1 1 -17.186
       2 -13.037 1 8.2988
## 2
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
```

```
model_6<-glm(cancer$remission~cancer$blast,data=cancer, family = binomial(link = "logit")</pre>
lrtest(model_1,model_6)
## Likelihood ratio test
## Model 1: cancer$remission ~ 1
## Model 2: cancer$remission ~ cancer$blast
    #Df LogLik Df Chisq Pr(>Chisq)
## 1 1 -17.186
## 2 2 -15.410 1 3.5513
                               0.0595 .
## ---
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
model_7<-glm(cancer$remission~cancer$temp,data=cancer, family = binomial(link = "logit")</pre>
lrtest(model_1,model_7)
## Likelihood ratio test
## Model 1: cancer$remission ~ 1
## Model 2: cancer$remission ~ cancer$temp
## #Df LogLik Df Chisq Pr(>Chisq)
## 1 1 -17.186
       2 -16.837 1 0.698
## 2
                              0.4035
# #define intercept-only model
# intercept_only <- lm(Y ~ 1, data=mydata_2) #define model with all predictors
# all <- lm(Y ~ \tilde{} ., data=mydata_2) #perform forward stepwise regression
# forward <- step(intercept_only, direction='forward', scope=formula(all), trace=0)</pre>
```

Przeprowadzone testy wskazują na najniższą wartość  $Pr(i_i)$  przy zmiennej  $i_i$  0.003967 dlatego przyłączamy ją do naszego modelu.

```
model_1<-glm(cancer$remission~cancer$li,data=cancer, family = binomial(link = "logit"))
model_2<-glm(cancer$remission~cancer$li+cancer$cell,data=cancer, family = binomial(link

lrtest(model_1,model_2)

## Likelihood ratio test

## # Model 1: cancer$remission ~ cancer$li

## Model 2: cancer$remission ~ cancer$li + cancer$cell

## #Df LogLik Df Chisq Pr(>Chisq)

## 1 2 -13.037

## 2 3 -12.170 1 1.7322 0.1881

model_3<-glm(cancer$remission~cancer$li+cancer$smear,data=cancer, family = binomial(link)

lrtest(model_1,model_3)

## Likelihood ratio test</pre>
```

```
##
## Model 1: cancer$remission ~ cancer$li
## Model 2: cancer$remission ~ cancer$li + cancer$smear
## #Df LogLik Df Chisq Pr(>Chisq)
## 1 2 -13.037
      3 -12.969 1 0.1356
## 2
                              0.7127
model_4<-glm(cancer$remission~cancer$li+cancer$infil,data=cancer, family = binomial(link
lrtest(model_1,model_4)
## Likelihood ratio test
##
## Model 1: cancer$remission ~ cancer$li
## Model 2: cancer$remission ~ cancer$li + cancer$infil
## #Df LogLik Df Chisq Pr(>Chisq)
     2 -13.037
## 1
## 2
      3 -12.745 1 0.5825
model_6<-glm(cancer$remission~cancer$li+cancer$blast,data=cancer, family = binomial(link
lrtest(model_1,model_6)
## Likelihood ratio test
##
## Model 1: cancer$remission ~ cancer$li
## Model 2: cancer$remission ~ cancer$li + cancer$blast
## #Df LogLik Df Chisq Pr(>Chisq)
## 1 2 -13.037
      3 -12.991 1 0.0918
## 2
                              0.7619
model_7<-glm(cancer$remission~cancer$li+cancer$temp,data=cancer, family = binomial(link
lrtest(model_1,model_7)
## Likelihood ratio test
## Model 1: cancer$remission ~ cancer$li
## Model 2: cancer$remission ~ cancer$li + cancer$temp
   #Df LogLik Df Chisq Pr(>Chisq)
## 1 2 -13.037
## 2 3 -12.324 1 1.4251 0.2326
```

Przeprowadzone testy wskazują na najniższą wartość Pr(i) przy zmiennej cell 0.1881 dlatego przyłączamy ją do naszego modelu.

```
model_1<-glm(cancer$remission~cancer$li+cancer$cell,data=cancer, family = binomial(link
model_2<-glm(cancer$remission~cancer$li+cancer$cell+cancer$smear,data=cancer, family = b
lrtest(model_1,model_2)

## Likelihood ratio test
##</pre>
```

```
## Model 1: cancer$remission ~ cancer$li + cancer$cell
## Model 2: cancer$remission ~ cancer$li + cancer$cell + cancer$smear
## #Df LogLik Df Chisq Pr(>Chisq)
      3 -12.170
## 1
## 2
      4 -12.146 1 0.0483
                              0.8261
model_4<-glm(cancer$remission~cancer$li+cancer$cell+cancer$infil,data=cancer, family = h
lrtest(model_1,model_4)
## Likelihood ratio test
## Model 1: cancer$remission ~ cancer$li + cancer$cell
## Model 2: cancer$remission ~ cancer$li + cancer$cell + cancer$infil
## #Df LogLik Df Chisq Pr(>Chisq)
## 1 3 -12.170
      4 -12.151 1 0.0391
## 2
                              0.8433
model_6<-glm(cancer$remission~cancer$li+cancer$cell+cancer$blast,data=cancer, family = h
lrtest(model_1,model_6)
## Likelihood ratio test
## Model 1: cancer$remission ~ cancer$li + cancer$cell
## Model 2: cancer$remission ~ cancer$li + cancer$cell + cancer$blast
## #Df LogLik Df Chisq Pr(>Chisq)
## 1 3 -12.170
## 2 4 -12.093 1 0.1545
                              0.6943
model_7<-glm(cancer$remission~cancer$li+cancer$cell+cancer$temp,data=cancer, family = b
lrtest(model_1,model_7)
## Likelihood ratio test
## Model 1: cancer$remission ~ cancer$li + cancer$cell
## Model 2: cancer$remission ~ cancer$li + cancer$cell + cancer$temp
## #Df LogLik Df Chisq Pr(>Chisq)
## 1 3 -12.170
## 2 4 -10.977 1 2.3874 0.1223
```

Przeprowadzone testy wskazują na najniższą wartość  $\Pr(\mathsf{¿Chisq})$  przy zmiennej temp~0.1223 dlatego przyłączamy ją do naszego modelu.

```
model_1<-glm(cancer$remission~cancer$li+cancer$cell+cancer$temp,data=cancer, family = boundel_2<-glm(cancer$remission~cancer$li+cancer$cell+cancer$temp+cancer$smear,data=cancer
lrtest(model_1,model_2)

## Likelihood ratio test
##
## Model 1: cancer$remission~ cancer$li + cancer$cell + cancer$temp</pre>
```

```
## Model 2: cancer$remission ~ cancer$li + cancer$cell + cancer$temp + cancer$smear
    #Df LogLik Df Chisq Pr(>Chisq)
## 1
      4 -10.977
## 2
      5 -10.929 1 0.0954
model_4<-glm(cancer$remission~cancer$li+cancer$cell+cancer$temp+cancer$infil,data=cancer
lrtest(model_1,model_4)
## Likelihood ratio test
##
## Model 1: cancer$remission ~ cancer$li + cancer$cell + cancer$temp
## Model 2: cancer$remission ~ cancer$li + cancer$cell + cancer$temp + cancer$infil
    #Df LogLik Df Chisq Pr(>Chisq)
      4 -10.977
## 2
      5 -10.935 1 0.0842
model_6<-glm(cancer$remission~cancer$li+cancer$cell+cancer$temp+cancer$blast,data=cancer
lrtest(model_1,model_6)
## Likelihood ratio test
##
## Model 1: cancer$remission ~ cancer$li + cancer$cell + cancer$temp
## Model 2: cancer$remission ~ cancer$li + cancer$cell + cancer$temp + cancer$blast
    #Df LogLik Df Chisq Pr(>Chisq)
     4 -10.977
## 1
## 2 5 -10.966 1 0.0209 0.8852
```

Przeprowadzone testy wskazują na najniższą wartość  $Pr(\xi Chisq)$  przy zmiennej *smear* 0.7574 nie przyłączamy jej jednak do modelu ze względu na to że jest wyższa od przyjętego poziomi istotności.

Nasz ostateczny model

```
summary(model_1)
##
## Call:
## glm(formula = cancer$remission ~ cancer$li + cancer$cell + cancer$temp,
      family = binomial(link = "logit"), data = cancer)
##
##
## Deviance Residuals:
                  1Q
                        Median
                                      30
                                               Max
## -2.02043 -0.66313 -0.08323 0.81282
                                           1.65887
##
## Coefficients:
              Estimate Std. Error z value Pr(>|z|)
                          56.888 1.189
## (Intercept) 67.634
                                           0.2345
## cancer$li
                 3.867
                           1.778 2.175
                                          0.0297 *
## cancer$cell
                 9.652
                            7.751 1.245
                                          0.2130
## cancer$temp -82.074
                       61.712 -1.330 0.1835
```

```
## ---
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
##
## (Dispersion parameter for binomial family taken to be 1)
##
## Null deviance: 34.372 on 26 degrees of freedom
## Residual deviance: 21.953 on 23 degrees of freedom
## AIC: 29.953
##
## Number of Fisher Scoring iterations: 7

sqrt(diag(summary(model_1)$cov.unscaled)*summary(model_1)$dispersion)
## (Intercept) cancer$li cancer$cell cancer$temp
## 56.887547 1.778278 7.751076 61.712382
```

- Predyktorami zostają zmienne *li,cell,temp*
- Wartości estymatorów wynoszą:  $\hat{\beta}_1 = 67.634$ ,  $\hat{\beta}_2 = 3.867$ ,  $\hat{\beta}_3 = 9.652$ ,  $\hat{\beta}_4 = -82.074$
- Wartości SE odpowiednio 56.887547, 1.778278, 7.751076, 61.712382
- AIC: 29.953

#### 2.3 Podpunkt c

• Dodajemy do tabeli cancer remission  $\pi(x) := P(remiss = 1 \mid X = x)$  oraz lewy i prawy koniec przedziału ufności na poziomie 95 procent.

```
func = function(x){ exp(x)/(1+exp(x))}
tabela <- data.frame(cancer)
predicted_values <- predict(model_1, data = cancer, type='link', se.fit=TRUE)
pi <-func(predicted_values$fit)
high <- func(predicted_values$fit + (predicted_values$se.fit*qnorm(0.95)))
low <- func(predicted_values$fit - (predicted_values$se.fit*qnorm(0.95)))
tabela<-cbind(tabela,pi,low,high)
kable(tabela)</pre>
```

remission	cell	smear	infil	li	blast	temp	pi	low	high
1	0.80	0.83	0.66	1.9	1.100	0.996	0.7226489	0.2344817	0.9568289
1	0.90	0.36	0.32	1.4	0.740	0.992	0.5787391	0.3115861	0.8065760
0	0.80	0.88	0.70	0.8	0.176	0.982	0.1045990	0.0119993	0.5291077
0	1.00	0.87	0.87	0.7	1.053	0.986	0.2825773	0.0946241	0.5974888
1	0.90	0.75	0.68	1.3	0.519	0.980	0.7141804	0.3175562	0.9306408
0	1.00	0.65	0.65	0.6	0.519	0.982	0.2708868	0.0765172	0.6248959
1	0.95	0.97	0.92	1.0	1.230	0.992	0.3215554	0.1549074	0.5506637
0	0.95	0.87	0.83	1.9	1.354	1.020	0.6072319	0.1516246	0.9304290
0	1.00	0.45	0.45	0.8	0.322	0.999	0.1663164	0.0402579	0.4868618
0	0.95	0.36	0.34	0.5	0.000	1.038	0.0015693	0.0000036	0.4090487
0	0.85	0.39	0.33	0.7	0.279	0.988	0.0728520	0.0092136	0.3990210
0	0.70	0.76	0.53	1.2	0.146	0.982	0.1728570	0.0110953	0.7956046
0	0.80	0.46	0.37	0.4	0.380	1.006	0.0034575	0.0000336	0.2636162
0	0.20	0.39	0.08	0.8	0.114	0.990	0.0001850	0.0000000	0.8017311
0	1.00	0.90	0.90	1.1	1.037	0.990	0.5712204	0.2968420	0.8078416
1	1.00	0.84	0.84	1.9	2.064	1.020	0.7146954	0.2167842	0.9577549
0	0.65	0.42	0.27	0.5	0.114	1.014	0.0006223	0.0000008	0.3203593
0	1.00	0.75	0.75	1.0	1.322	1.004	0.2228888	0.0590836	0.5671126
0	0.50	0.44	0.22	0.6	0.114	0.990	0.0015425	0.0000022	0.5260312
1	1.00	0.63	0.63	1.1	1.072	0.986	0.6491095	0.3174187	0.8803675
0	1.00	0.33	0.33	0.4	0.176	1.010	0.0169297	0.0005605	0.3459119
0	0.90	0.93	0.84	0.6	1.591	1.020	0.0062175	0.0000721	0.3517630
1	1.00	0.58	0.58	1.0	0.531	1.002	0.2526057	0.0784679	0.5729315
0	0.95	0.32	0.30	1.6	0.886	0.988	0.8701089	0.4993059	0.9782600
1	1.00	0.60	0.60	1.7	0.964	0.990	0.9313166	0.5549406	0.9932640
1	1.00	0.69	0.69	0.9	0.398	0.986	0.4605092	0.2011016	0.7432325
0	1.00	0.73	0.73	0.7	0.398	0.986	0.2825773	0.0946241	0.5974888