

Aprendizagem Estatística em Altas Dimensões

[MAE0501/MAE5904/IBI5904]

Ícaro Maia Santos de Castro¹
Rayssa de Carvalho Roberto²
Rodrigo Aoyama Nakahara³
Rodrigo Araujo⁴
Vitor Hugo Vieira de Lima⁵

Dezembro de 2020

Sumário

Leitura dos Dados	2
Análise Descritiva	2
Análise Discriminante	11
Modelagem	11
Testes	19
Regressão Logística	28
Modelagem	28
Teste	34
Random Forest	40
Modelagem	40
Árvores de decisão geradas	46
Teste	48
Support Vector Machine	51
Modelagem	51
Testes	64
Validação	70
AD Quadrática - Dados Removidos	70
SVM Kernel Não Linear - Dados Imputados	72

¹Número USP: 11866921

²Número USP: 10940828

³Número USP: 3510922

⁴Número USP: 9299208

⁵Número USP: 10263886

Leitura dos Dados

```
diabetes <- read_csv("diabetes.csv")

## Parsed with column specification:
## cols(
##   Pregnancies = col_double(),
##   Glucose = col_double(),
##   BloodPressure = col_double(),
##   SkinThickness = col_double(),
##   Insulin = col_double(),
##   BMI = col_double(),
##   DiabetesPedigreeFunction = col_double(),
##   Age = col_double(),
##   Outcome = col_double()
## )

colnames(diabetes)[9] <- "diabetes"
diabetes$diabetes <- as.factor(diabetes$diabetes)
levels(diabetes$diabetes) <- c("No", "Yes")

diabetes[, 2:6][diabetes[, 2:6] == 0] <- NA

head(diabetes) %>% kable(caption="Dados.")
```

Tabela 1: Dados.

Pregnancies	Glucose	BloodPressure	SkinThickness	Insulin	BMI	DiabetesPedigreeFunction	Age	diabetes
6	148	72	35	NA	33,6	0,63	50	Yes
1	85	66	29	NA	26,6	0,35	31	No
8	183	64	NA	NA	23,3	0,67	32	Yes
1	89	66	23	94	28,1	0,17	21	No
0	137	40	35	168	43,1	2,29	33	Yes
5	116	74	NA	NA	25,6	0,20	30	No

Análise Descritiva

```
# OBS: análise descritiva com base no conjunto de treinamento para evitar data snooping

library(dplyr)
library(caret)
library(lattice)
library(ggplot2)
library(GGally)
```

```
## Registered S3 method overwritten by 'GGally':  
##   method from  
##   +.gg   ggplot2
```

```
library(ggcorrplot)
```

```
## Warning: package 'ggcorrplot' was built under R version 4.0.2
```

```
library(scales)
```

```
##  
## Attaching package: 'scales'
```

```
## The following objects are masked from 'package:psych':  
##  
##   alpha, rescale
```

```
## The following object is masked from 'package:purrr':  
##  
##   discard
```

```
## The following object is masked from 'package:readr':  
##  
##   col_factor
```

```
library(pROC)
```

```
## Warning: package 'pROC' was built under R version 4.0.2
```

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

```
##  
## Attaching package: 'pROC'
```

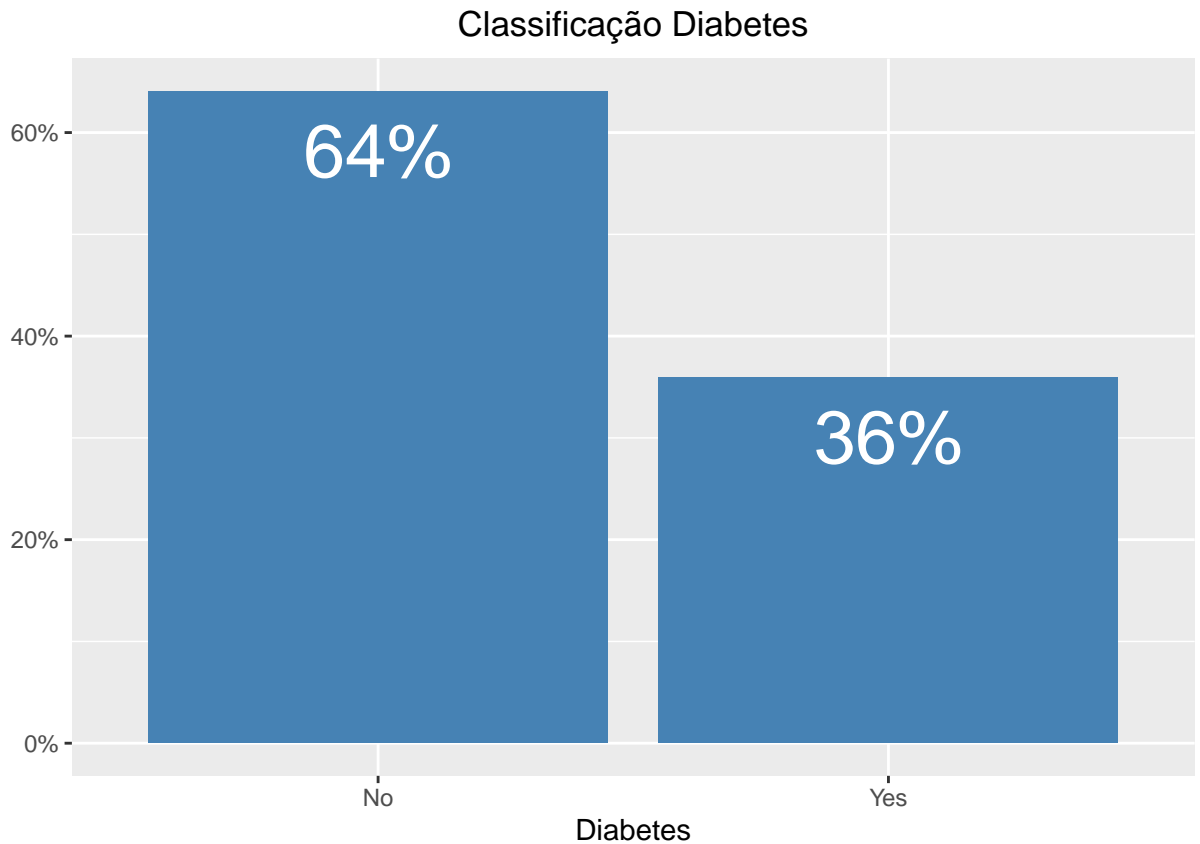
```
## The following object is masked from 'package:colorspace':  
##  
##   coords
```

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

```
# Distribuição da variável resposta
```

```
train %>% count(Diabetes = factor(diabetes)) %>% mutate(pct = prop.table(n)) %>%  
  ggplot(aes(x = Diabetes, y = pct, fill = pct, label = scales::percent(pct))) +
```

```
geom_col(position = 'dodge', fill="steelblue") +
labs(title = "Classificação Diabetes", x = "Diabetes", y = "") +
geom_text(aes(label=scales::percent(pct) ), vjust=1.6, color="white", size=10) +
scale_y_continuous(labels = scales::percent) +
theme(plot.title = element_text(hjust = 0.5), legend.title = element_blank())
```



Matriz completa com dispersão, densidades, correlações, retas de regressão (com IC) e LOESS (com IC)

```
# Função auxiliar para curvas de regressão linear e LOESS nos gráficos abaixo
curvas <- function(data, mapping, ...){
  p <- ggplot(data = data, mapping = mapping) +
    geom_point(size = 0.5) +
    geom_smooth(method = loess, fill = "red", color = "red", ...) +
    geom_smooth(method = lm, fill = "blue", color = "blue", ...)
  p
}
```

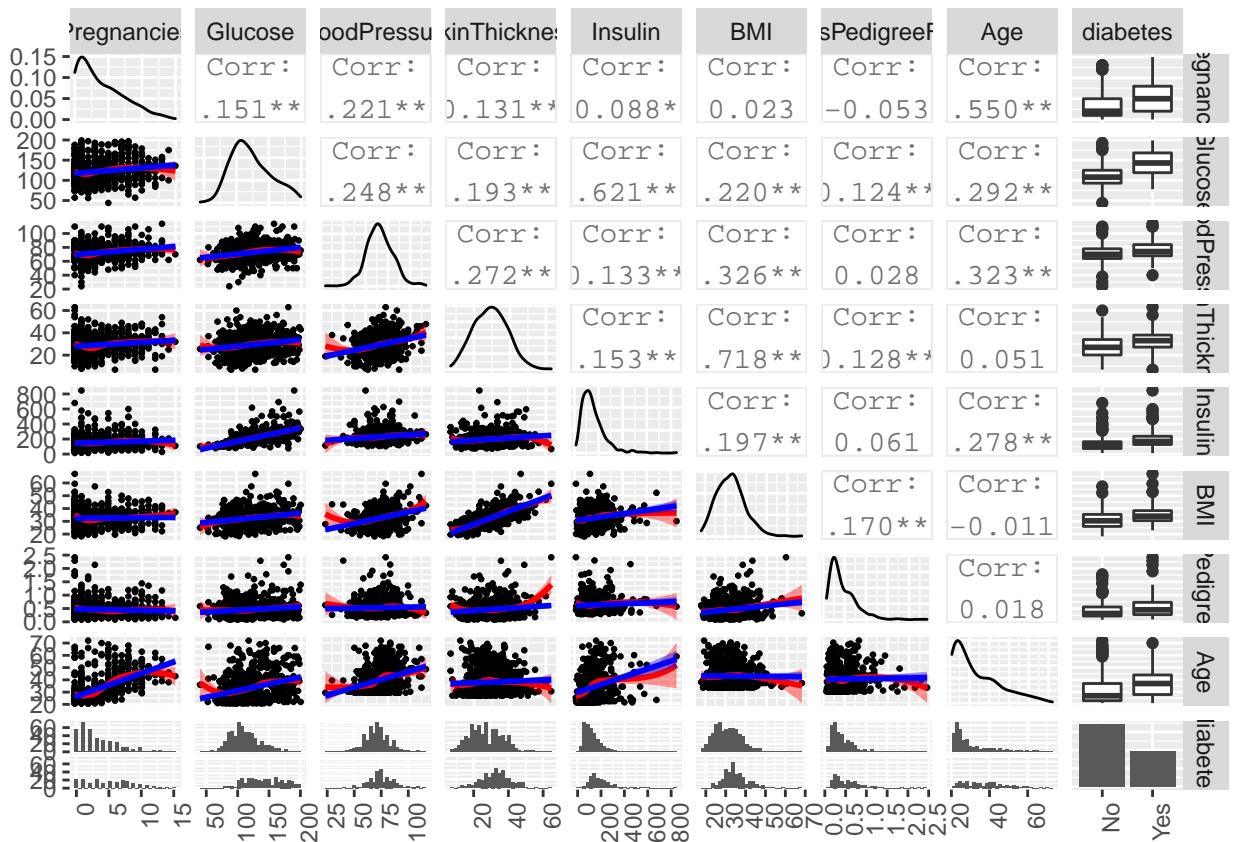
```
ggpairs(train, columns = 1:9, lower = list(continuous = curvas)) + # Obs: pode demorar para montar o gr
theme(axis.text.x = element_text(angle = 90, hjust = 1))
```

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

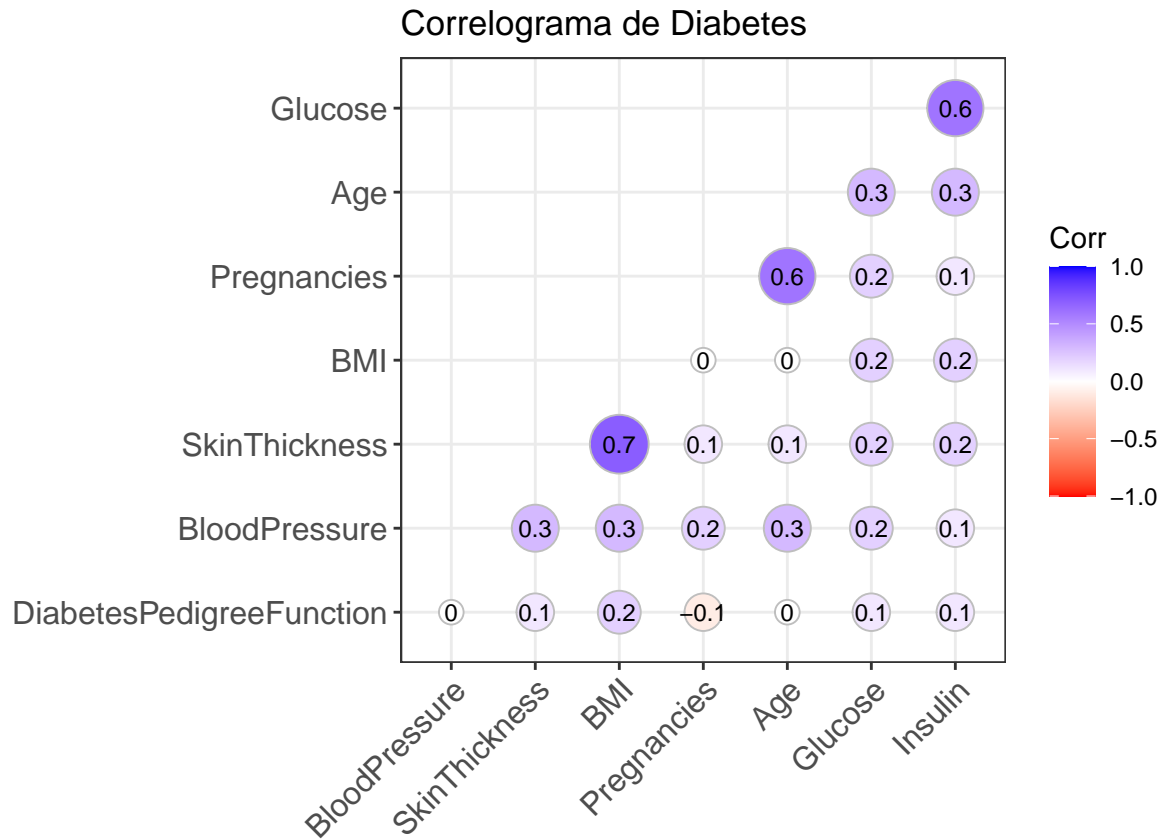
```
## 'geom_smooth()' using formula 'y ~ x'
```

[illegible]

```
## 'stat_bin()' using 'bins = 30'. Pick better value with 'binwidth'.
## 'stat_bin()' using 'bins = 30'. Pick better value with 'binwidth'.
## 'stat_bin()' using 'bins = 30'. Pick better value with 'binwidth'.
## 'stat_bin()' using 'bins = 30'. Pick better value with 'binwidth'.
## 'stat_bin()' using 'bins = 30'. Pick better value with 'binwidth'.
## 'stat_bin()' using 'bins = 30'. Pick better value with 'binwidth'.
## 'stat_bin()' using 'bins = 30'. Pick better value with 'binwidth'.
## 'stat_bin()' using 'bins = 30'. Pick better value with 'binwidth'.
```



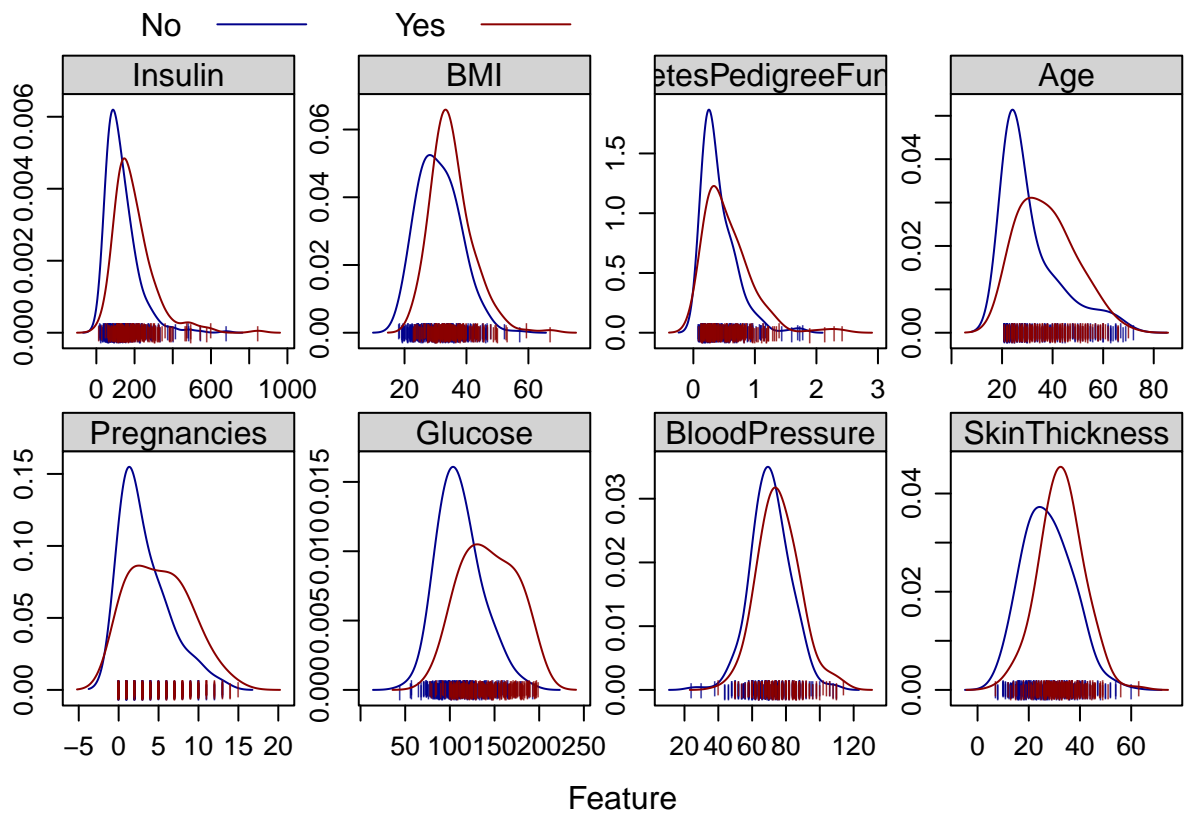
```
# Correlações/Correlograma
corr<-round(cor(train[,-9]),1)
ggcorrplot(corr, hc.order = TRUE,
  type = "lower",
  lab = TRUE,
  lab_size = 3,
  method="circle",
  colors = c("red", "white", "blue"),
  title="Correlograma de Diabetes",
  ggtheme=theme_bw)
```



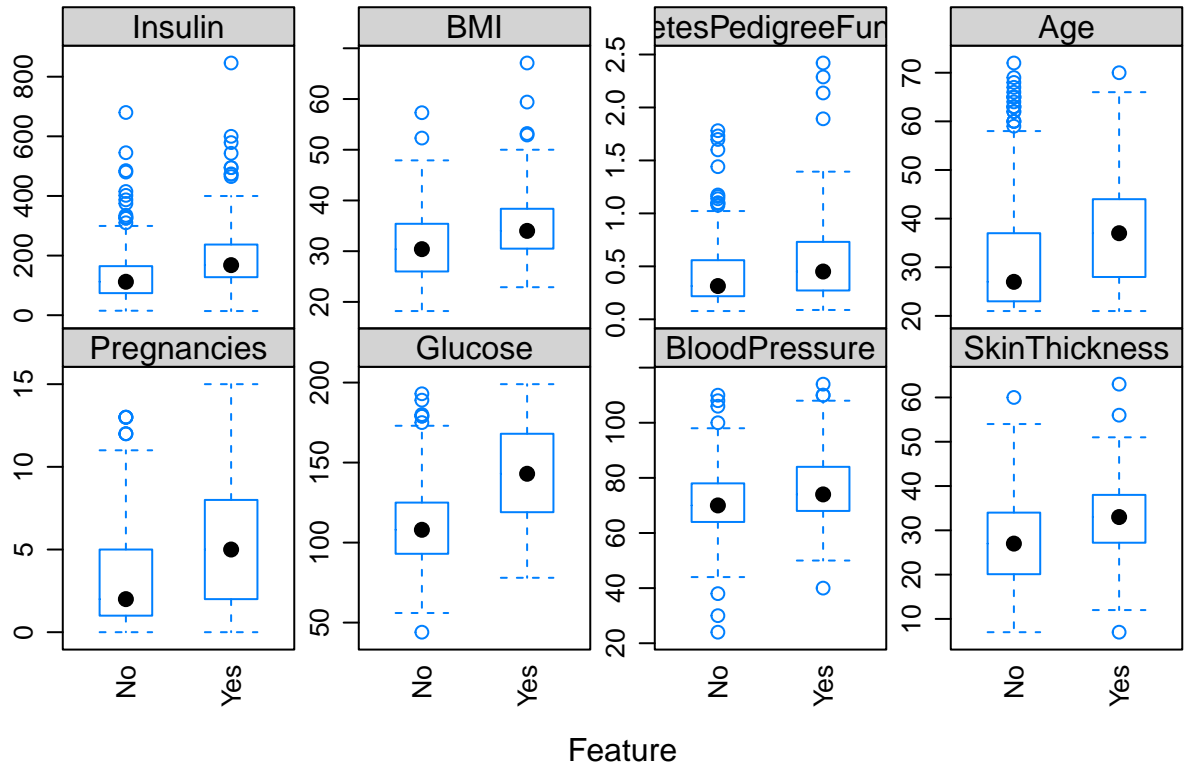
Distribuição da diabetes Yes/No por feature

```
featurePlot(x = train[, 1:8], # Densidades alisadas
            y = train$diabetes,
            plot = "density",
            scales = list(x = list(relation="free"), y = list(relation="free")),
            adjust = 1.5,
            pch = "|",
            layout = c(4, 2),
            auto.key = list(columns = 4),
            par.settings = list(strip.background=list(col="lightgrey"),
                                superpose.line = list(col = c("darkblue","darkred")),
                                superpose.symbol = list(col = c("darkblue","darkred")) ))
```

Warning in draw.key(simpleKey(...), draw = FALSE): not enough rows for columns



```
featurePlot(x = train[, 1:8], # Boxplots
            y = train$diabetes,
            plot = "box",
            scales = list(y = list(relation="free"), x = list(rot = 90)),
            layout = c(4,2 ),
            auto.key = list(columns = 4),
            par.settings = list(strip.background=list(col="lightgrey")))
```

Funções auxiliares

```
library(dplyr)
library(ggraph)
```

```
## Warning: package 'ggraph' was built under R version 4.0.3
```

```
library(igraph)
```

```
## Warning: package 'igraph' was built under R version 4.0.2
```

```
##
## Attaching package: 'igraph'
```

```
## The following objects are masked from 'package:dplyr':
##
##   as_data_frame, groups, union
```

```
## The following objects are masked from 'package:purrr':
##
##   compose, simplify
```

```

## The following object is masked from 'package:tidyr':
##
##     crossing

## The following object is masked from 'package:tibble':
##
##     as_data_frame

## The following objects are masked from 'package:stats':
##
##     decompose, spectrum

## The following object is masked from 'package:base':
##
##     union

plotaroc <- function(rocobj, titulo = "Curva ROC"){
  # Função que plota as curvas roc para os modelos ajustados
  b <- which.max(rocobj$sensitivities + rocobj$specificities)
  best <- round(c(rocobj$thresholds[b],rocobj$specificities[b],rocobj$sensitivities[b]), 3)

  pROC::ggroc(rocobj, col = "red", alpha = 0.5, size = 0.5) +
    theme_gray() +
    ggtitle(titulo) +
    geom_abline(intercept = 1, slope=1, linetype = "dashed") +
    labs(x="Especificidade", y = "Sensibilidade") +
    geom_point(data = tibble(Sensibilidade = best[2],
                             Especificidade = best[3]),
               mapping = aes(x=Sensibilidade, y=Especificidade),
               col = "black") +
    geom_text(mapping = aes(x = best[2] - 0.15,
                           y = best[3] - 0.05),
              label = paste( best[1], "(", best[2], ",", best[3], ")")) +
    geom_text(mapping = aes(x = 0.5,
                           y = 0.01),
              label = paste("AUC: ", round(rocobj$auc,3)))
}

tree_func <- function(final_model,
                      tree_num) {

  # get tree by index
  tree <- randomForest::getTree(final_model,
                                k = tree_num,
                                labelVar = TRUE) %>%
    tibble::rownames_to_column() %>%
    # make leaf split points to NA, so the 0s won't get plotted
    mutate('split point' = ifelse(is.na(prediction), 'split point', NA))

  # prepare data frame for graph
  graph_frame <- data.frame(from = rep(tree$rowname, 2),
                            to = c(tree$'left daughter', tree$'right daughter'))

```

```

# convert to graph and delete the last node that we don't want to plot
graph <- graph_from_data_frame(graph_frame) %>%
  delete_vertices("0")

# set node labels
V(graph)$node_label <- gsub("_", " ", as.character(tree$`split var`))
V(graph)$leaf_label <- as.character(tree$prediction)
V(graph)$split <- as.character(round(tree$`split point`, digits = 2))

# plot
plot <- ggraph(graph, 'dendrogram') +
  theme_bw() +
  geom_edge_link() +
  geom_node_point() +
  geom_node_text(aes(label = node_label), na.rm = TRUE, repel = TRUE) +
  geom_node_label(aes(label = split, vjust = 2, na.rm = TRUE, fill = "white") +
  geom_node_label(aes(label = leaf_label, fill = leaf_label), na.rm = TRUE,
    repel = TRUE, colour = "white", fontface = "bold", show.legend = FALSE) +
  theme(panel.grid.minor = element_blank(),
    panel.grid.major = element_blank(),
    panel.background = element_blank(),
    plot.background = element_rect(fill = "white"),
    panel.border = element_blank(),
    axis.line = element_blank(),
    axis.text.x = element_blank(),
    axis.text.y = element_blank(),
    axis.ticks = element_blank(),
    axis.title.x = element_blank(),
    axis.title.y = element_blank(),
    plot.title = element_text(size = 14))

return(plot)
}

```

Análise Discriminante

Modelagem

```

# Obs: possibilidades de modelos de AD: rda, lda, pda (acurácias iguais) e qda (pior)

train.control <- caret::trainControl(method = "cv", number = 15, classProbs = TRUE) # Cross-validation

## ADL com dados imputados
set.seed(23)
modeloAD1 <- caret::train(diabetes ~ ., data = train, trControl = train.control, method = "lda")
print(modeloAD1)

## Linear Discriminant Analysis

```

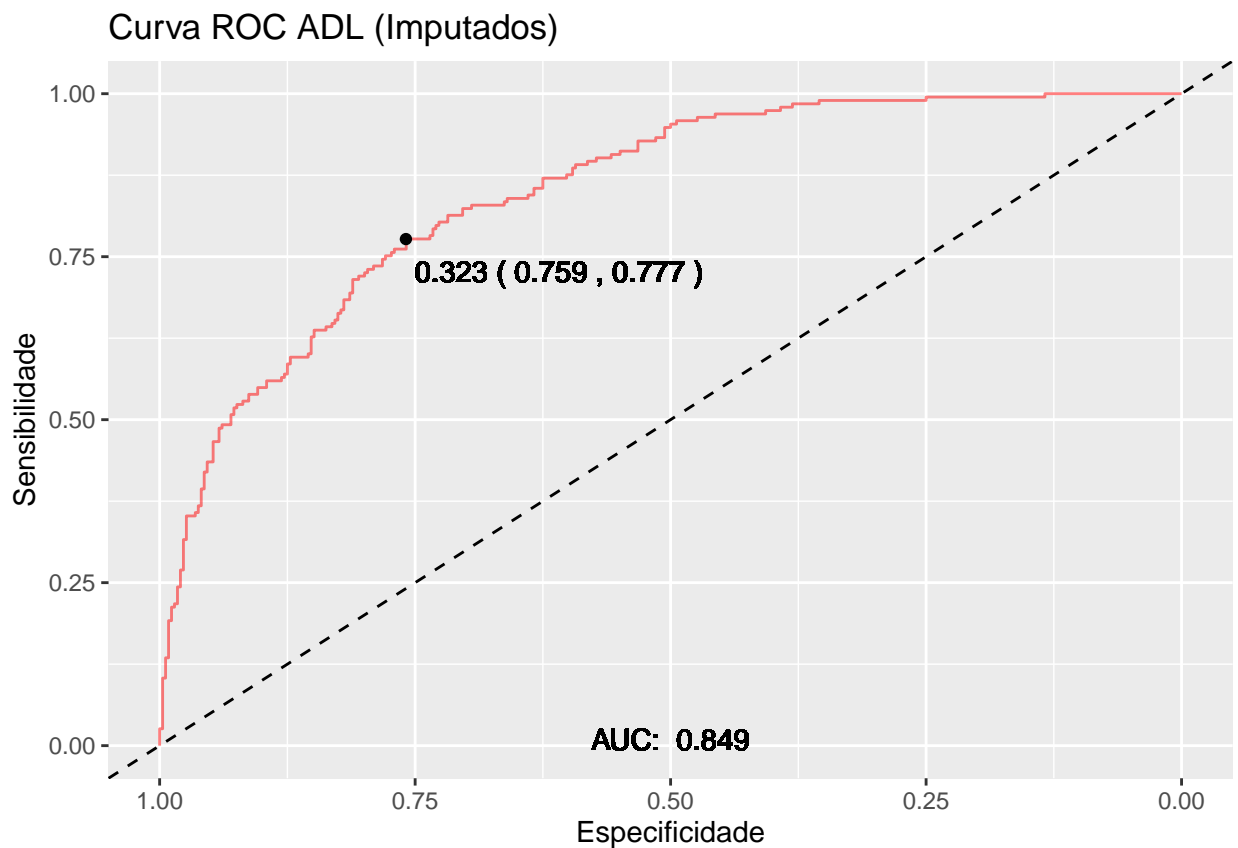
```
##
## 537 samples
## 8 predictor
## 2 classes: 'No', 'Yes'
##
## No pre-processing
## Resampling: Cross-Validated (15 fold)
## Summary of sample sizes: 501, 501, 501, 501, 501, 501, ...
## Resampling results:
##
## Accuracy   Kappa
## 0.7636508  0.4650271

rocAD1 <- roc(response = train$diabetes, predictor = predict(modeloAD1, train, type = "prob")[,2])

## Setting levels: control = No, case = Yes

## Setting direction: controls < cases

plotaroc(rocAD1, titulo = "Curva ROC ADL (Imputados)")
```



```

## ADL sem missing
set.seed(23)
modeloAD2 <- caret::train(diabetes ~ ., data = train_without_NAs, trControl = train.control, method = "
print(modeloAD2)

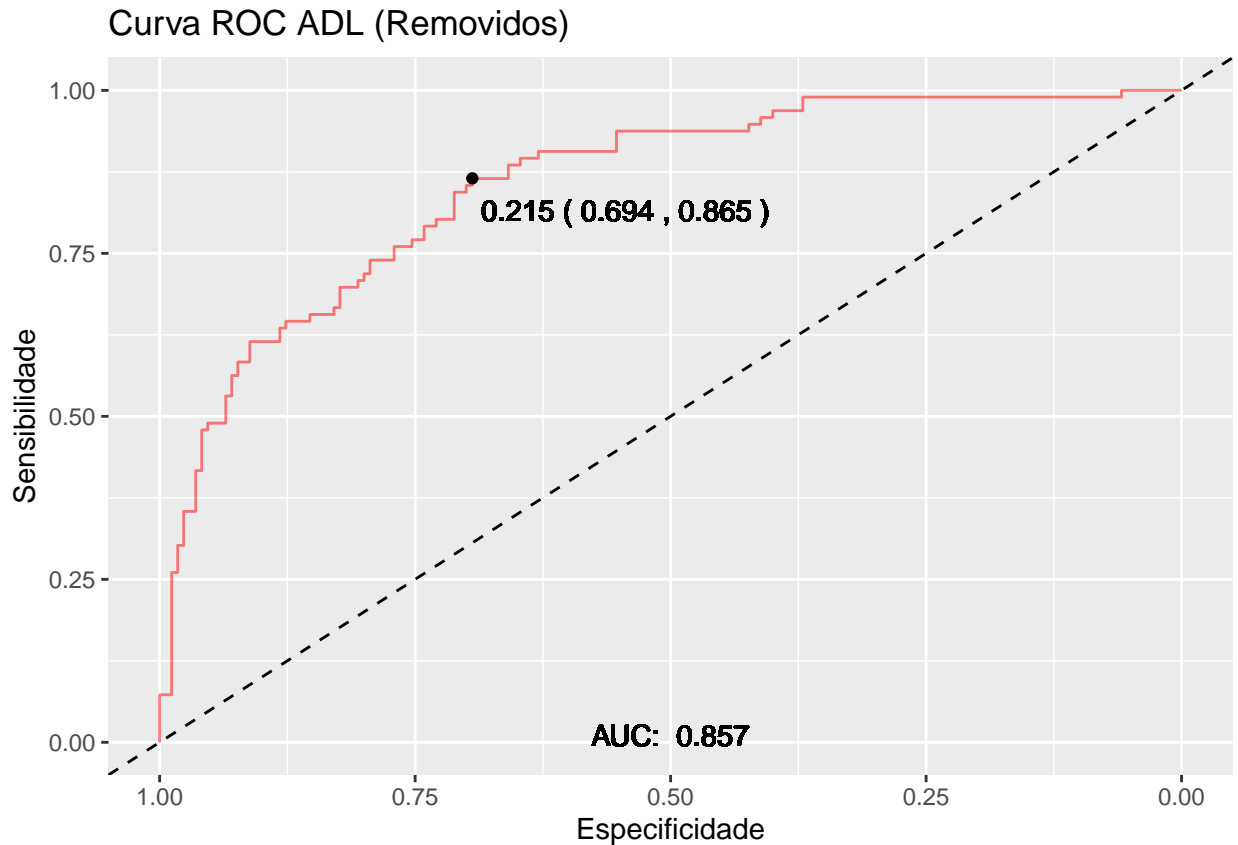
## Linear Discriminant Analysis
##
## 266 samples
## 8 predictor
## 2 classes: 'No', 'Yes'
##
## No pre-processing
## Resampling: Cross-Validated (15 fold)
## Summary of sample sizes: 248, 248, 249, 248, 247, 249, ...
## Resampling results:
##
## Accuracy   Kappa
## 0.7784199  0.5053595

rocAD2 <- roc(response = train_without_NAs$diabetes, predictor = predict(modeloAD2, train_without_NAs,

## Setting levels: control = No, case = Yes
## Setting direction: controls < cases

plotaroc(rocAD2, titulo = "Curva ROC ADL (Removidos)")

```



```
## AD Flexível com dados imputados
set.seed(23)
modeloAD3 <- caret::train(diabetes ~ ., data = train, trControl = train.control, method = "fda")
```

```
## Loading required package: earth
```

```
## Warning: package 'earth' was built under R version 4.0.3
```

```
## Loading required package: plotmo
```

```
## Warning: package 'plotmo' was built under R version 4.0.3
```

```
## Loading required package: plotrix
```

```
##
```

```
## Attaching package: 'plotrix'
```

```
## The following object is masked from 'package:scales':
```

```
##
```

```
## rescale
```

```
## The following object is masked from 'package:psych':  
##  
##     rescale
```

```
## Loading required package: TeachingDemos
```

```
## Warning: package 'TeachingDemos' was built under R version 4.0.3
```

```
##  
## Attaching package: 'TeachingDemos'
```

```
## The following objects are masked from 'package:Hmisc':  
##  
##     cnvrt.coords, subplot
```

```
print(modeloAD3)
```

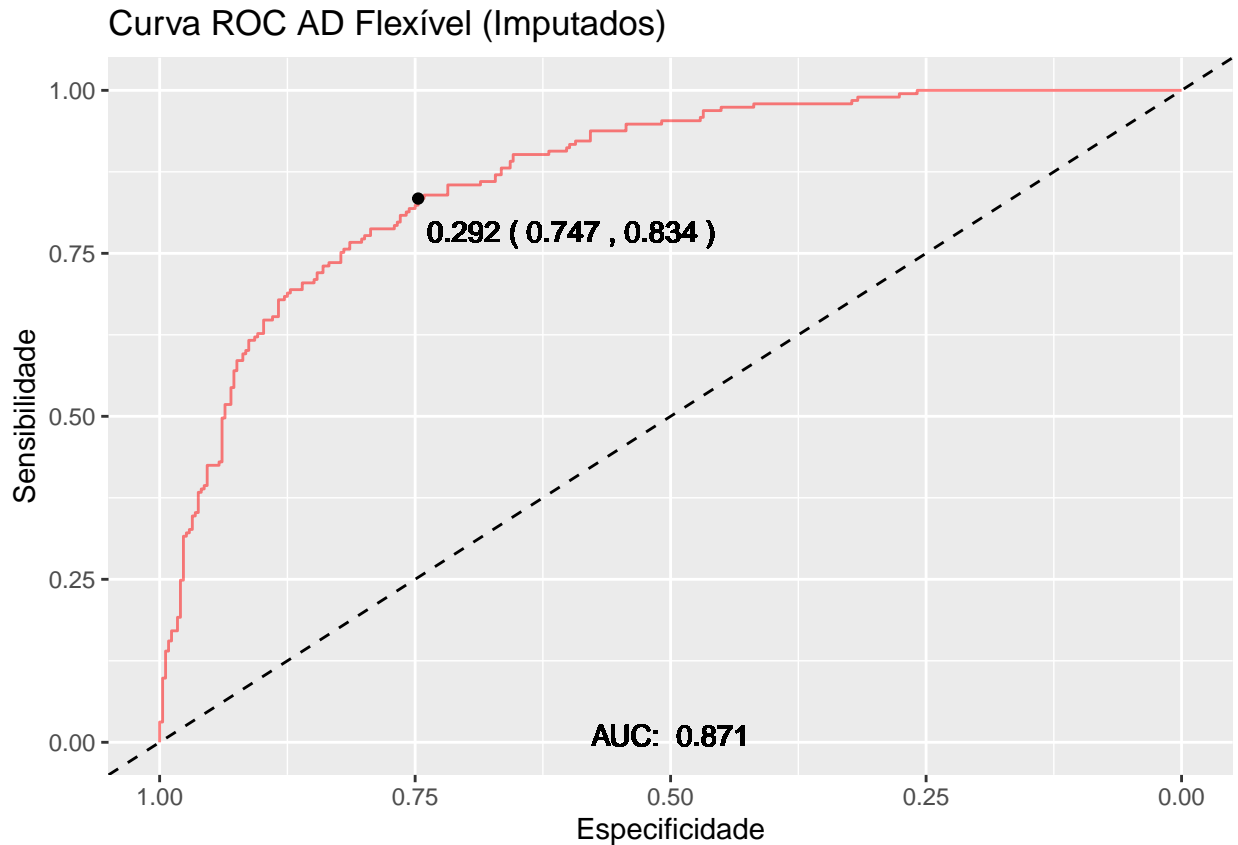
```
## Flexible Discriminant Analysis  
##  
## 537 samples  
## 8 predictor  
## 2 classes: 'No', 'Yes'  
##  
## No pre-processing  
## Resampling: Cross-Validated (15 fold)  
## Summary of sample sizes: 501, 501, 501, 501, 501, 501, ...  
## Resampling results across tuning parameters:  
##  
##   nprune  Accuracy   Kappa  
##    2      0.7450794 0.4129759  
##    8      0.7823280 0.5204850  
##   15      0.7823280 0.5192155  
##  
## Tuning parameter 'degree' was held constant at a value of 1  
## Accuracy was used to select the optimal model using the largest value.  
## The final values used for the model were degree = 1 and nprune = 8.
```

```
rocAD3 <- roc(response = train$diabetes, predictor = predict(modeloAD3, train, type = "prob"),[,2])
```

```
## Setting levels: control = No, case = Yes
```

```
## Setting direction: controls < cases
```

```
plotaroc(rocAD3, titulo = "Curva ROC AD Flexível (Imputados)")
```



```
## AD Flexível sem missing
set.seed(23)
modeloAD4 <- caret::train(diabetes ~ ., data = train_without_NAs, trControl = train.control, method = "
print(modeloAD4)
```

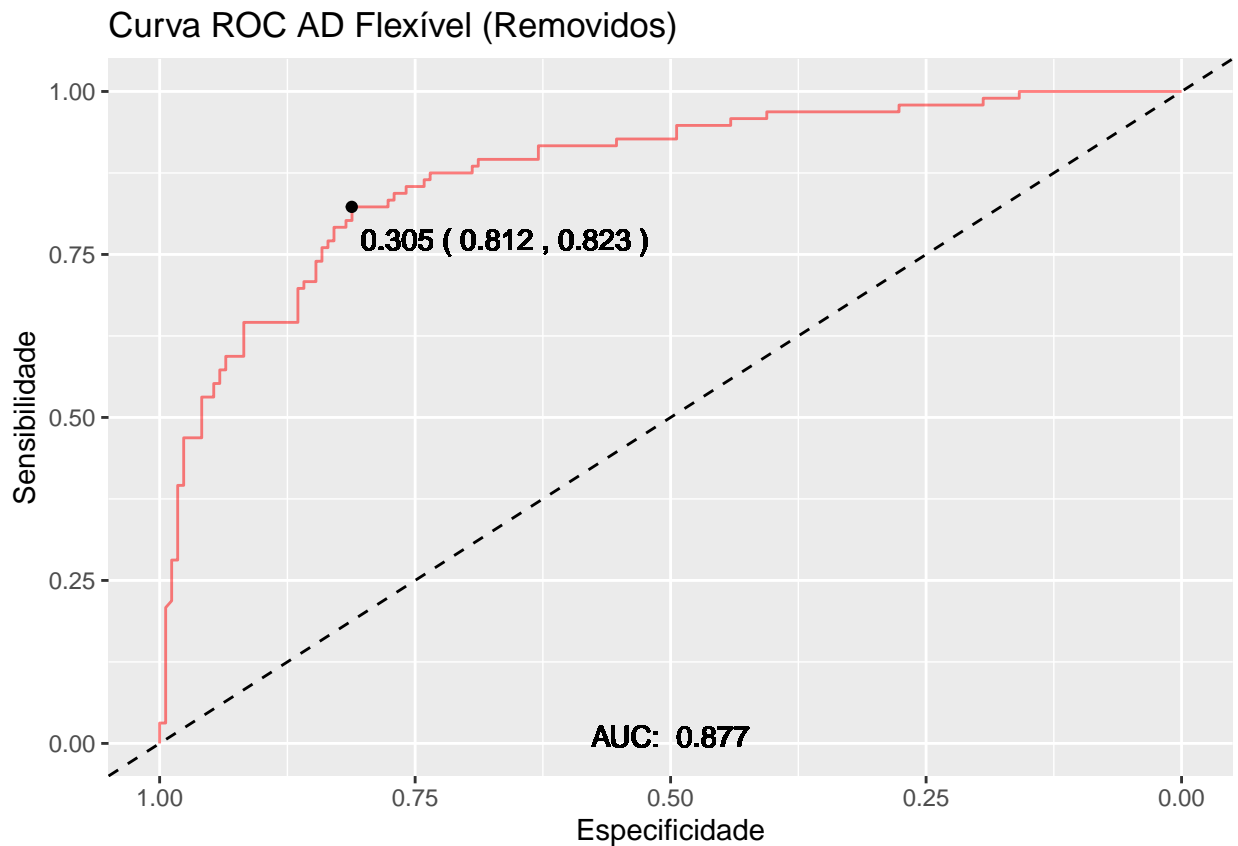
```
## Flexible Discriminant Analysis
##
## 266 samples
## 8 predictor
## 2 classes: 'No', 'Yes'
##
## No pre-processing
## Resampling: Cross-Validated (15 fold)
## Summary of sample sizes: 248, 248, 249, 248, 247, 249, ...
## Resampling results across tuning parameters:
##
##  nprune  Accuracy  Kappa
##    2      0.7481137 0.4233678
##    8      0.7516455 0.4422724
##   15      0.7446279 0.4240573
##
## Tuning parameter 'degree' was held constant at a value of 1
## Accuracy was used to select the optimal model using the largest value.
## The final values used for the model were degree = 1 and nprune = 8.
```



```
rocAD4 <- roc(response = train_without_NAs$diabetes, predictor = predict(modeloAD4, train_without_NAs,

## Setting levels: control = No, case = Yes
## Setting direction: controls < cases

plotaroc(rocAD4, titulo = "Curva ROC AD Flexível (Removidos)")
```



```
## AD Quadrática com dados imputados
set.seed(23)
modeloAD5 <- caret::train(diabetes ~ ., data = train, trControl = train.control, method = "qda")
print(modeloAD5)

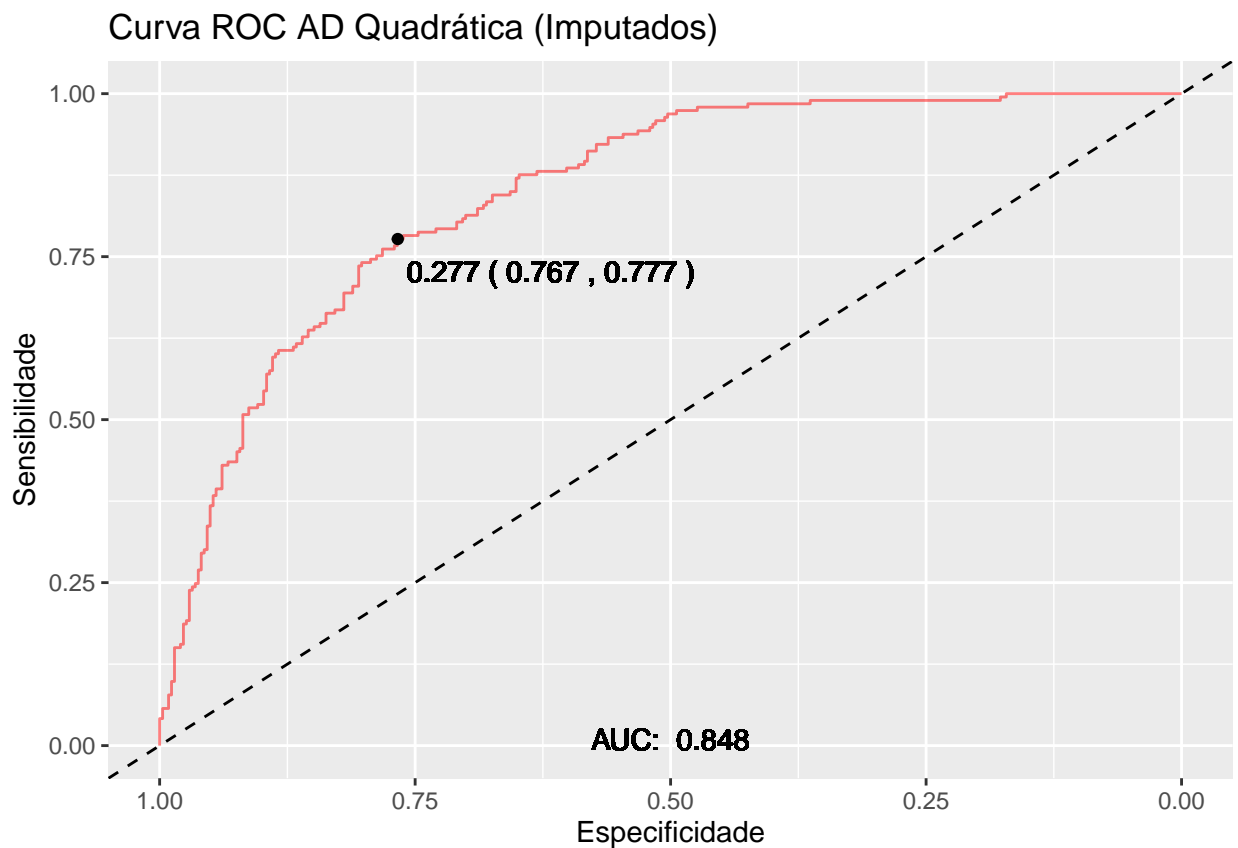
## Quadratic Discriminant Analysis
##
## 537 samples
## 8 predictor
## 2 classes: 'No', 'Yes'
##
## No pre-processing
## Resampling: Cross-Validated (15 fold)
## Summary of sample sizes: 501, 501, 501, 501, 501, 501, ...
## Resampling results:
##
```

```
## Accuracy Kappa
## 0.7431217 0.4201363
```

```
rocAD5 <- roc(response = train$diabetes, predictor = predict(modeloAD5, train, type = "prob")[,2])
```

```
## Setting levels: control = No, case = Yes
## Setting direction: controls < cases
```

```
plotaroc(rocAD5, titulo = "Curva ROC AD Quadrática (Imputados)")
```



```
## AD Quadrática sem missing
```

```
set.seed(23)
```

```
modeloAD6 <- caret::train(diabetes ~ ., data = train_without_NAs, trControl = train.control, method = "qda")
```

```
print(modeloAD6)
```

```
## Quadratic Discriminant Analysis
```

```
##
```

```
## 266 samples
```

```
## 8 predictor
```

```
## 2 classes: 'No', 'Yes'
```

```
##
```

```
## No pre-processing
```

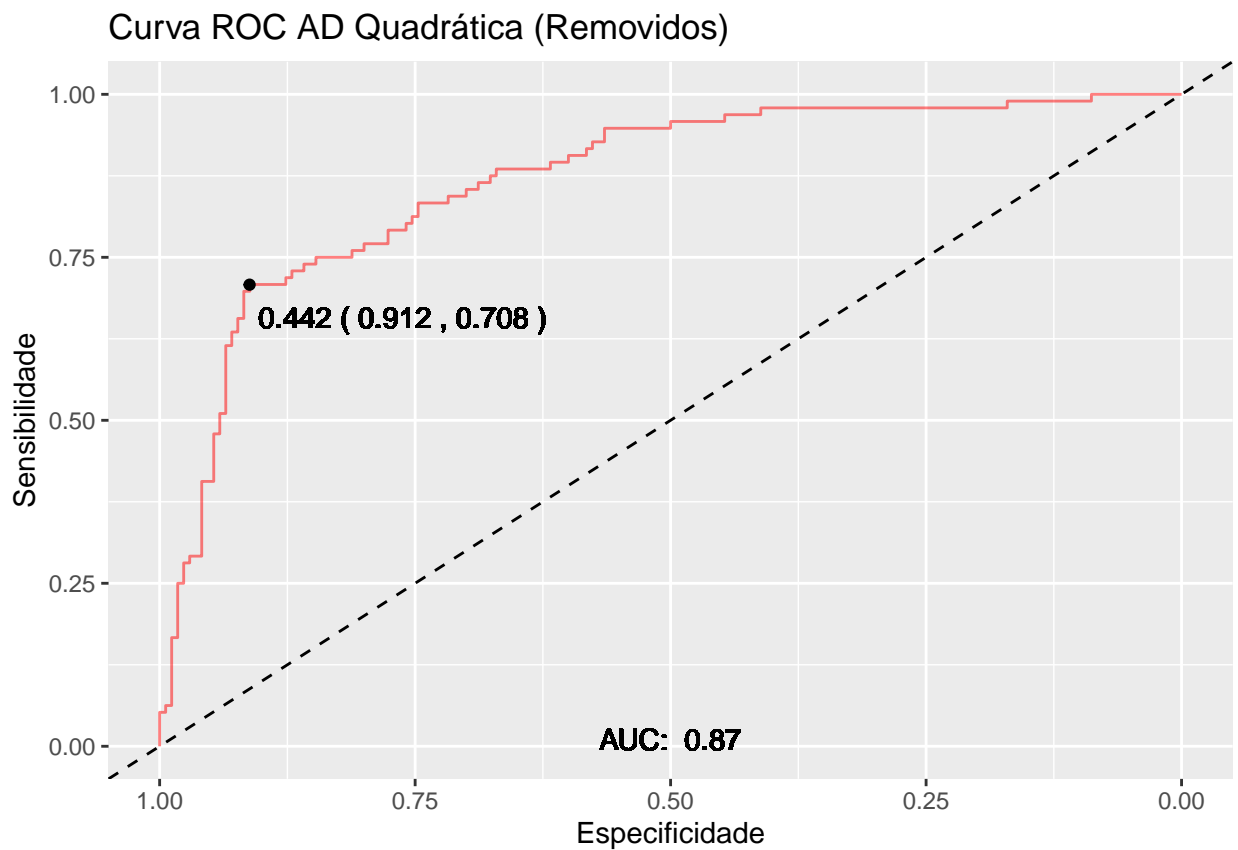
```
## Resampling: Cross-Validated (15 fold)
```

```
## Summary of sample sizes: 248, 248, 249, 248, 247, 249, ...
## Resampling results:
##
## Accuracy Kappa
## 0.7860681 0.5158632
```

```
rocAD6 <- roc(response = train_without_NAs$diabetes, predictor = predict(modeloAD6, train_without_NAs,
```

```
## Setting levels: control = No, case = Yes
## Setting direction: controls < cases
```

```
plotaroc(rocAD6, titulo = "Curva ROC AD Quadrática (Removidos)")
```



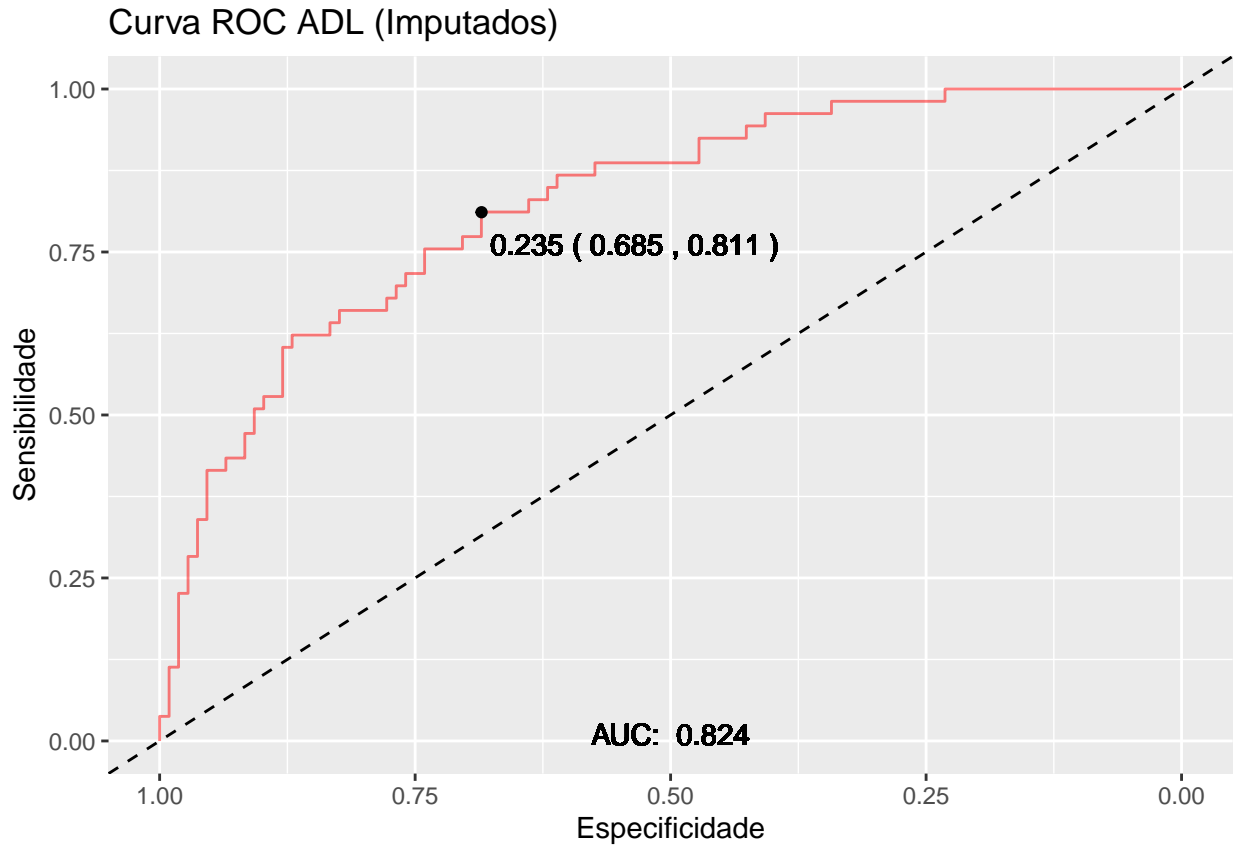
Testes

```
## ADL com dados imputados
rocAD1 <- roc(response = test$diabetes, predictor = predict(modeloAD1, test, type = "prob")[,2])
```

```
## Setting levels: control = No, case = Yes
```

```
## Setting direction: controls < cases
```

```
plotaroc(rocAD1, titulo = "Curva ROC ADL (Imputados)")
```



```
predictAD1 <- predict(modeloAD1, newdata = test)
confusionMatrix(predictAD1, test$diabetes)
```

```
## Confusion Matrix and Statistics
##
##           Reference
## Prediction No Yes
##      No  99  28
##      Yes   9  25
##
##           Accuracy : 0.7702
##           95% CI : (0.6974, 0.8327)
##      No Information Rate : 0.6708
##      P-Value [Acc > NIR] : 0.003815
##
##           Kappa : 0.4274
##
##  McNemar's Test P-Value : 0.003085
##
##           Sensitivity : 0.9167
##           Specificity : 0.4717
##           Pos Pred Value : 0.7795
```

```
##      Neg Pred Value : 0.7353
##      Prevalence : 0.6708
##      Detection Rate : 0.6149
##      Detection Prevalence : 0.7888
##      Balanced Accuracy : 0.6942
##
##      'Positive' Class : No
##
```

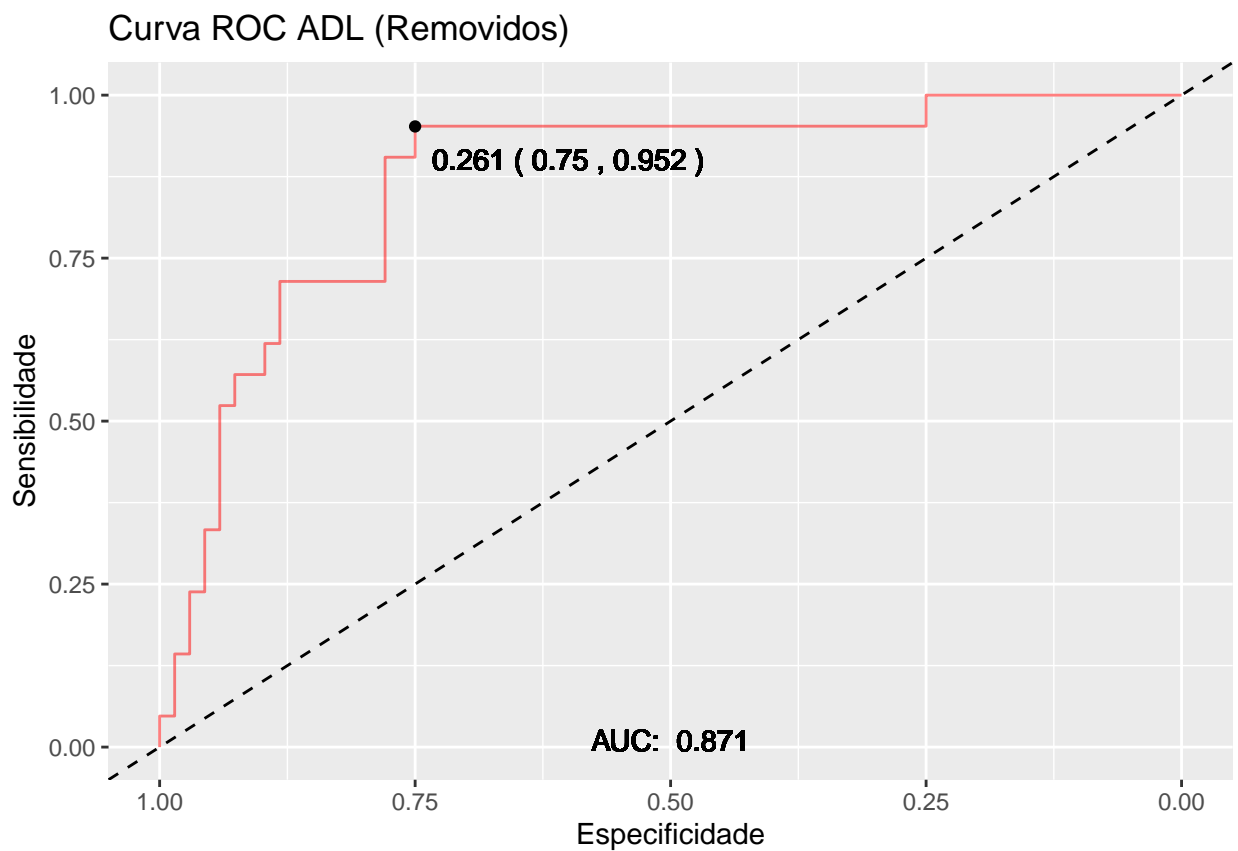
```
## ADL sem missing
```

```
rocAD2 <- roc(response = test_without_NAs$diabetes, predictor = predict(modeloAD2, test_without_NAs, type = "prob"))
```

```
## Setting levels: control = No, case = Yes
```

```
## Setting direction: controls < cases
```

```
plotaroc(rocAD2, titulo = "Curva ROC ADL (Removidos)")
```



```
predictAD2 <- predict(modeloAD2, newdata = test_without_NAs)
confusionMatrix(predictAD2, test_without_NAs$diabetes)
```

```
## Confusion Matrix and Statistics
```

```
##
```

```
##      Reference
```

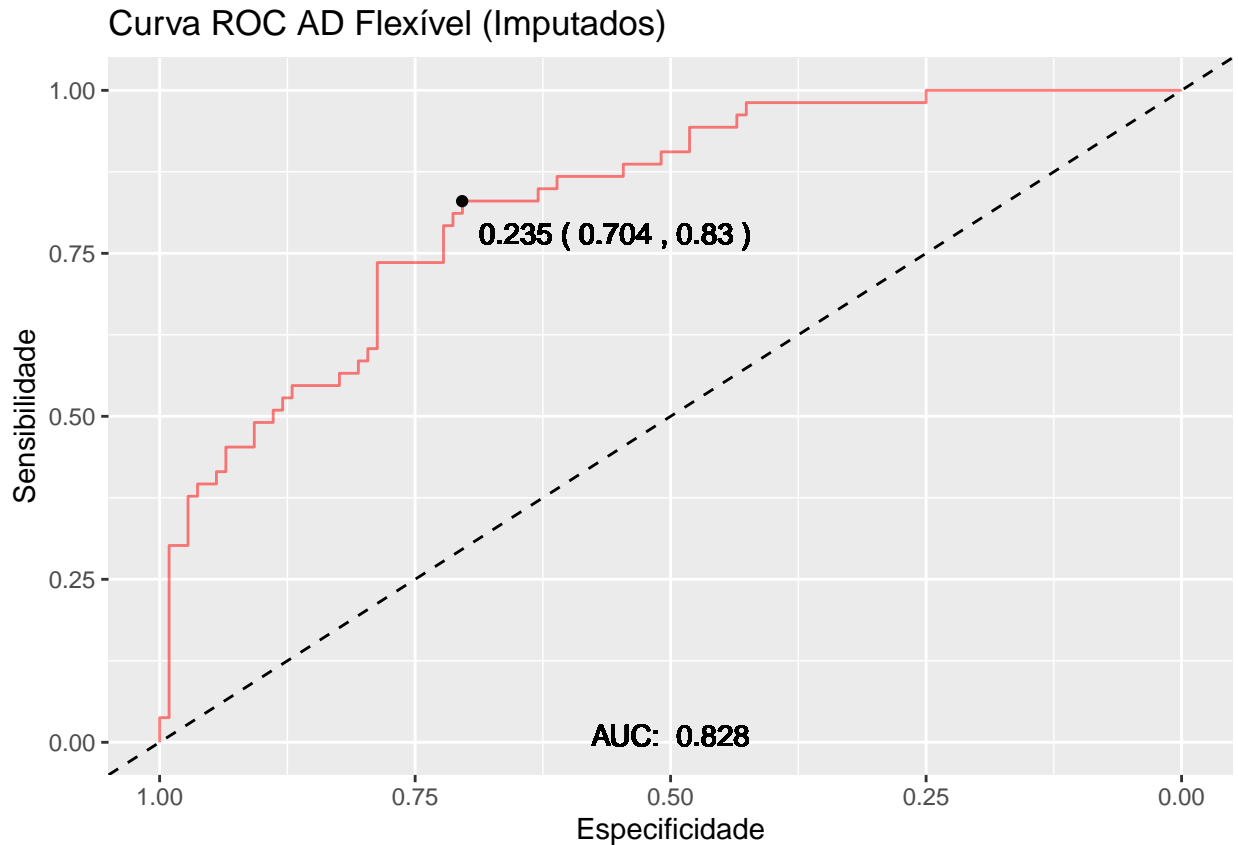
```

## Prediction No Yes
##      No  64  12
##      Yes  4   9
##
##              Accuracy : 0.8202
##              95% CI : (0.7245, 0.8936)
##      No Information Rate : 0.764
##      P-Value [Acc > NIR] : 0.12910
##
##              Kappa : 0.4258
##
##      McNemar's Test P-Value : 0.08012
##
##      Sensitivity : 0.9412
##      Specificity : 0.4286
##      Pos Pred Value : 0.8421
##      Neg Pred Value : 0.6923
##      Prevalence : 0.7640
##      Detection Rate : 0.7191
##      Detection Prevalence : 0.8539
##      Balanced Accuracy : 0.6849
##
##      'Positive' Class : No
##
## ADL Flexível com dados imputados
rocAD3 <- roc(response = test$diabetes, predictor = predict(modeloAD3, test, type = "prob"),[,2])

## Setting levels: control = No, case = Yes
## Setting direction: controls < cases

plotaroc(rocAD3, titulo = "Curva ROC AD Flexível (Imputados)")

```



```
predictAD3 <- predict(modeloAD3, newdata = test)
confusionMatrix(predictAD3, test$diabetes)
```

```
## Confusion Matrix and Statistics
##
##           Reference
## Prediction No Yes
##      No  95  26
##      Yes  13  27
##
##           Accuracy : 0.7578
##           95% CI : (0.6841, 0.8217)
##      No Information Rate : 0.6708
##      P-Value [Acc > NIR] : 0.01039
##
##           Kappa : 0.415
##
##  Mcnemar's Test P-Value : 0.05466
##
##           Sensitivity : 0.8796
##           Specificity : 0.5094
##      Pos Pred Value : 0.7851
##      Neg Pred Value : 0.6750
##           Prevalence : 0.6708
##      Detection Rate : 0.5901
```

```
## Detection Prevalence : 0.7516
## Balanced Accuracy : 0.6945
##
## 'Positive' Class : No
##
```

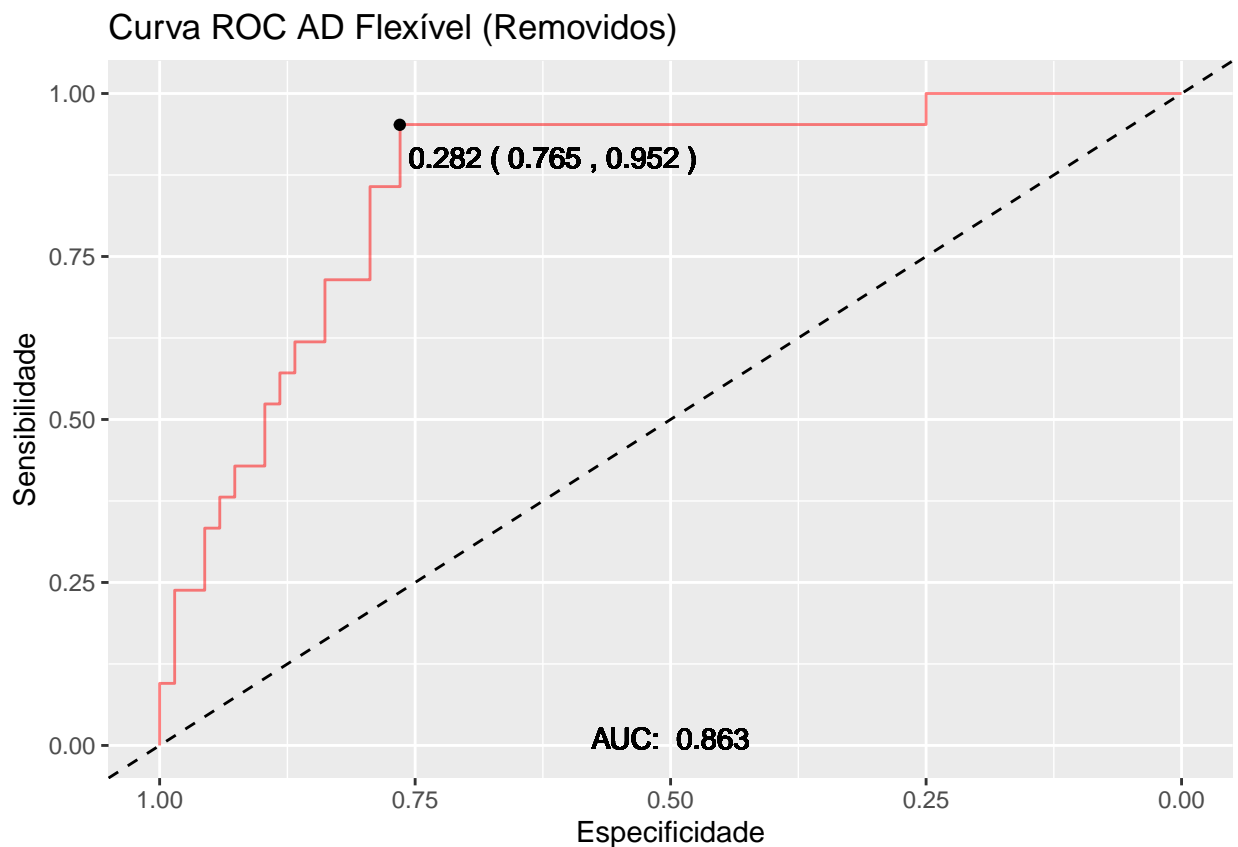
```
## ADL Flexível sem missing
```

```
rocAD4 <- roc(response = test_without_NAs$diabetes, predictor = predict(modeloAD4, test_without_NAs, type = "prob"))
```

```
## Setting levels: control = No, case = Yes
```

```
## Setting direction: controls < cases
```

```
plotaroc(rocAD4, titulo = "Curva ROC AD Flexível (Removidos)")
```



```
predictAD4 <- predict(modeloAD4, newdata = test_without_NAs)
confusionMatrix(predictAD4, test_without_NAs$diabetes)
```

```
## Confusion Matrix and Statistics
```

```
##
```

```
## Reference
```

```
## Prediction No Yes
```

```
## No 61 11
```

```
## Yes 7 10
```



```

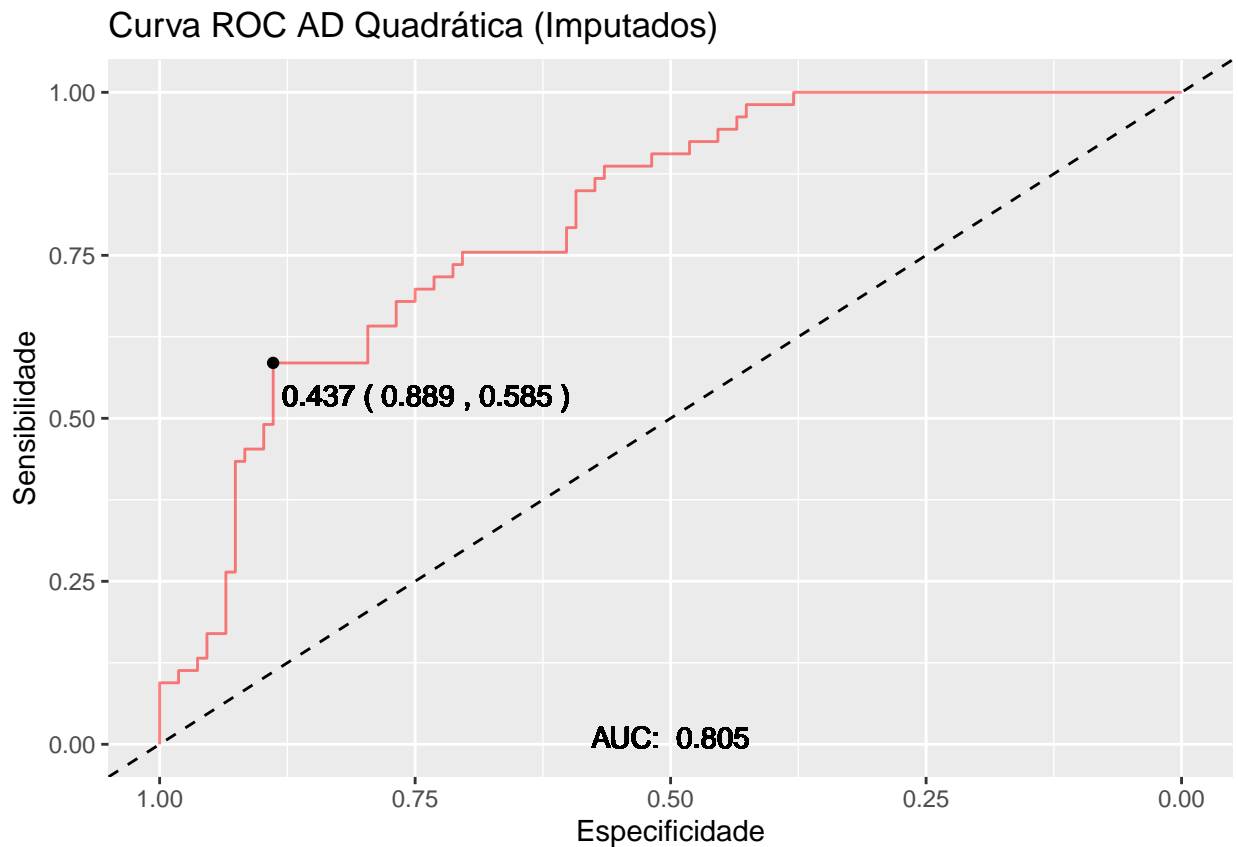
##
##           Accuracy : 0.7978
##           95% CI   : (0.6993, 0.8755)
##    No Information Rate : 0.764
##    P-Value [Acc > NIR] : 0.2709
##
##           Kappa : 0.3996
##
##    McNemar's Test P-Value : 0.4795
##
##           Sensitivity : 0.8971
##           Specificity : 0.4762
##           Pos Pred Value : 0.8472
##           Neg Pred Value : 0.5882
##           Prevalence : 0.7640
##           Detection Rate : 0.6854
##           Detection Prevalence : 0.8090
##           Balanced Accuracy : 0.6866
##
##           'Positive' Class : No
##

## AD Quadrática com dados imputados
rocAD5 <- roc(response = test$diabetes, predictor = predict(modeloAD5, test, type = "prob"),[,2])

## Setting levels: control = No, case = Yes
## Setting direction: controls < cases

plotaroc(rocAD5, titulo = "Curva ROC AD Quadrática (Imputados)")

```



```
predictAD5 <- predict(modeloAD5, newdata = test)
confusionMatrix(predictAD5, test$diabetes)
```

```
## Confusion Matrix and Statistics
##
##           Reference
## Prediction No Yes
##      No  96  25
##      Yes  12  28
##
##           Accuracy : 0.7702
##           95% CI : (0.6974, 0.8327)
##      No Information Rate : 0.6708
##      P-Value [Acc > NIR] : 0.003815
##
##           Kappa : 0.445
##
##  Mcnemar's Test P-Value : 0.048520
##
##           Sensitivity : 0.8889
##           Specificity : 0.5283
##      Pos Pred Value : 0.7934
##      Neg Pred Value : 0.7000
##           Prevalence : 0.6708
##      Detection Rate : 0.5963
```

```
## Detection Prevalence : 0.7516
## Balanced Accuracy : 0.7086
##
## 'Positive' Class : No
##
```

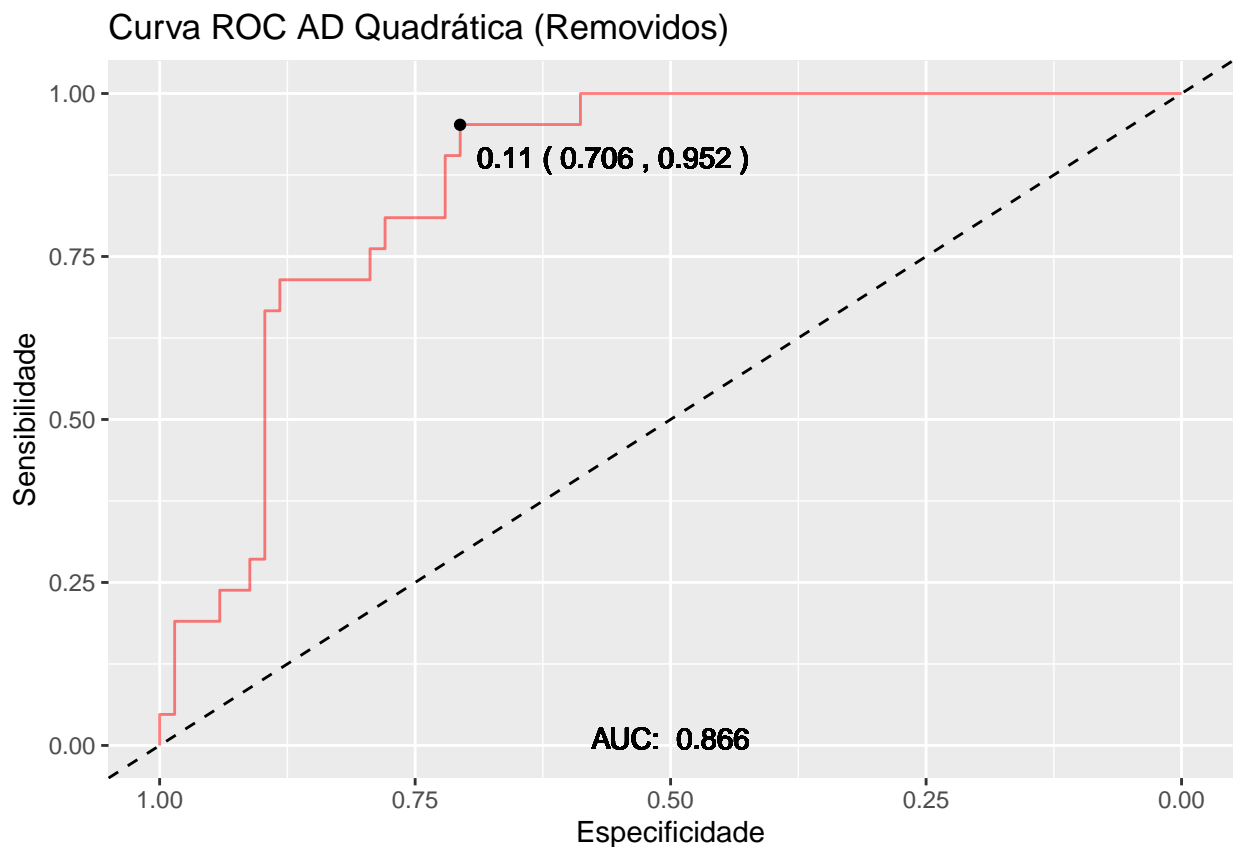
```
## AD Quadrática sem missing
```

```
rocAD6 <- roc(response = test_without_NAs$diabetes, predictor = predict(modeloAD6, test_without_NAs, type = "prob"))
```

```
## Setting levels: control = No, case = Yes
```

```
## Setting direction: controls < cases
```

```
plotaroc(rocAD6, titulo = "Curva ROC AD Quadrática (Removidos)")
```



```
predictAD6 <- predict(modeloAD6, newdata = test_without_NAs)
confusionMatrix(predictAD6, test_without_NAs$diabetes)
```

```
## Confusion Matrix and Statistics
```

```
##
```

```
## Reference
```

```
## Prediction No Yes
```

```
## No 60 6
```

```
## Yes 8 15
```

```
##
##           Accuracy : 0.8427
##           95% CI : (0.7502, 0.9112)
##      No Information Rate : 0.764
##      P-Value [Acc > NIR] : 0.04778
##
##           Kappa : 0.5776
##
##  McNemar's Test P-Value : 0.78927
##
##           Sensitivity : 0.8824
##           Specificity : 0.7143
##      Pos Pred Value : 0.9091
##      Neg Pred Value : 0.6522
##           Prevalence : 0.7640
##      Detection Rate : 0.6742
##      Detection Prevalence : 0.7416
##      Balanced Accuracy : 0.7983
##
##      'Positive' Class : No
##
```

Regressão Logística

Modelagem

```
library(caret)

train.control <- caret::trainControl(method = "cv", number = 10) # Cross-validation com k=10

## Logística simples com ligação logit / dados imputados
set.seed(23)
modeloRL1 <- caret::train(diabetes ~ ., data = train, trControl = train.control, method = "glm", family=
print(modeloRL1)

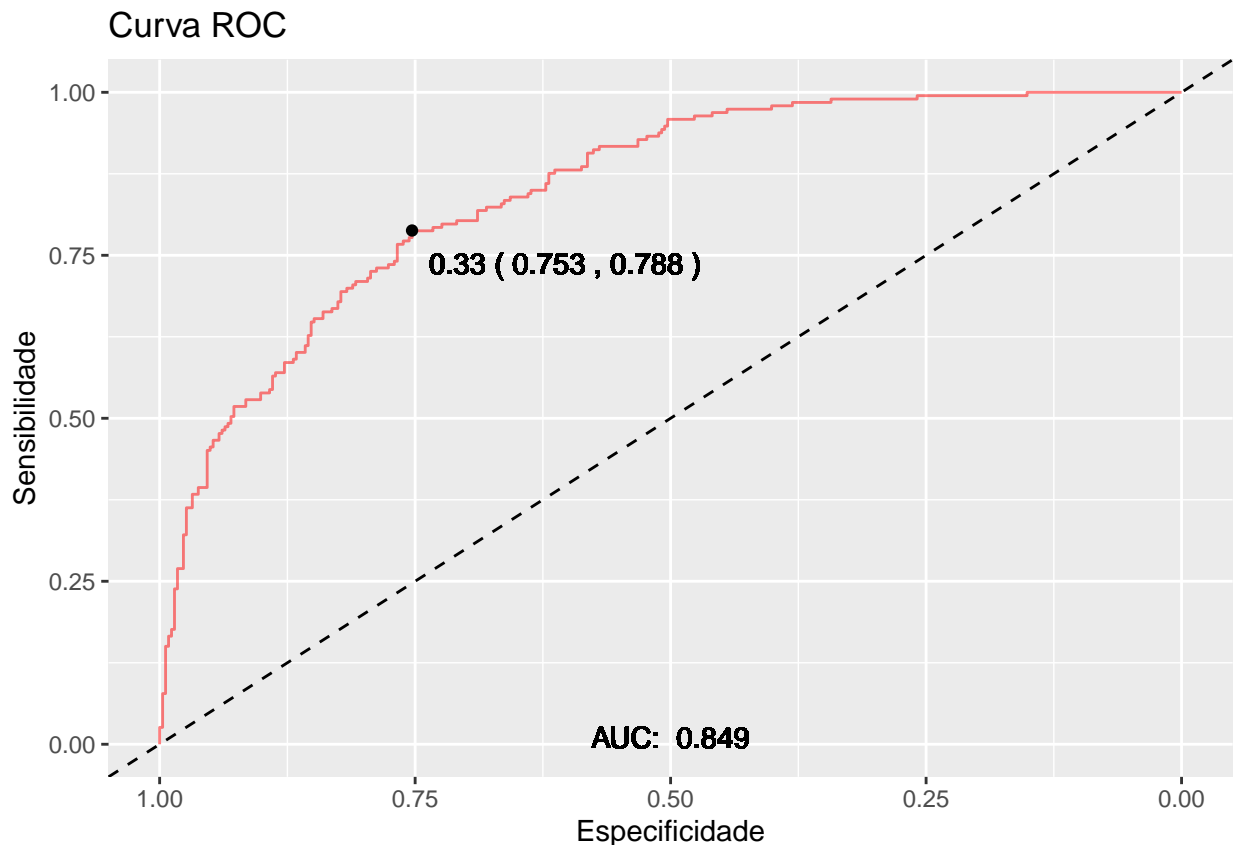
## Generalized Linear Model
##
## 537 samples
## 8 predictor
## 2 classes: 'No', 'Yes'
##
## No pre-processing
## Resampling: Cross-Validated (10 fold)
## Summary of sample sizes: 483, 483, 484, 483, 482, 484, ...
## Resampling results:
##
## Accuracy      Kappa
## 0.7595623     0.4586088
```

```
rocRL1 <- roc(response = train$diabetes, predictor = predict(modeloRL1, train, type = "prob")[,2])
```

```
## Setting levels: control = No, case = Yes
```

```
## Setting direction: controls < cases
```

```
plotaroc(rocRL1)
```



```
## Logística simples com ligação logit / dados sem missing
```

```
set.seed(23)
```

```
modeloRL2 <- caret::train(diabetes ~ ., data = train_without_NAs, trControl = train.control, method = "glm",
print(modeloRL2))
```

```
## Generalized Linear Model
```

```
##
```

```
## 266 samples
```

```
## 8 predictor
```

```
## 2 classes: 'No', 'Yes'
```

```
##
```

```
## No pre-processing
```

```
## Resampling: Cross-Validated (10 fold)
```

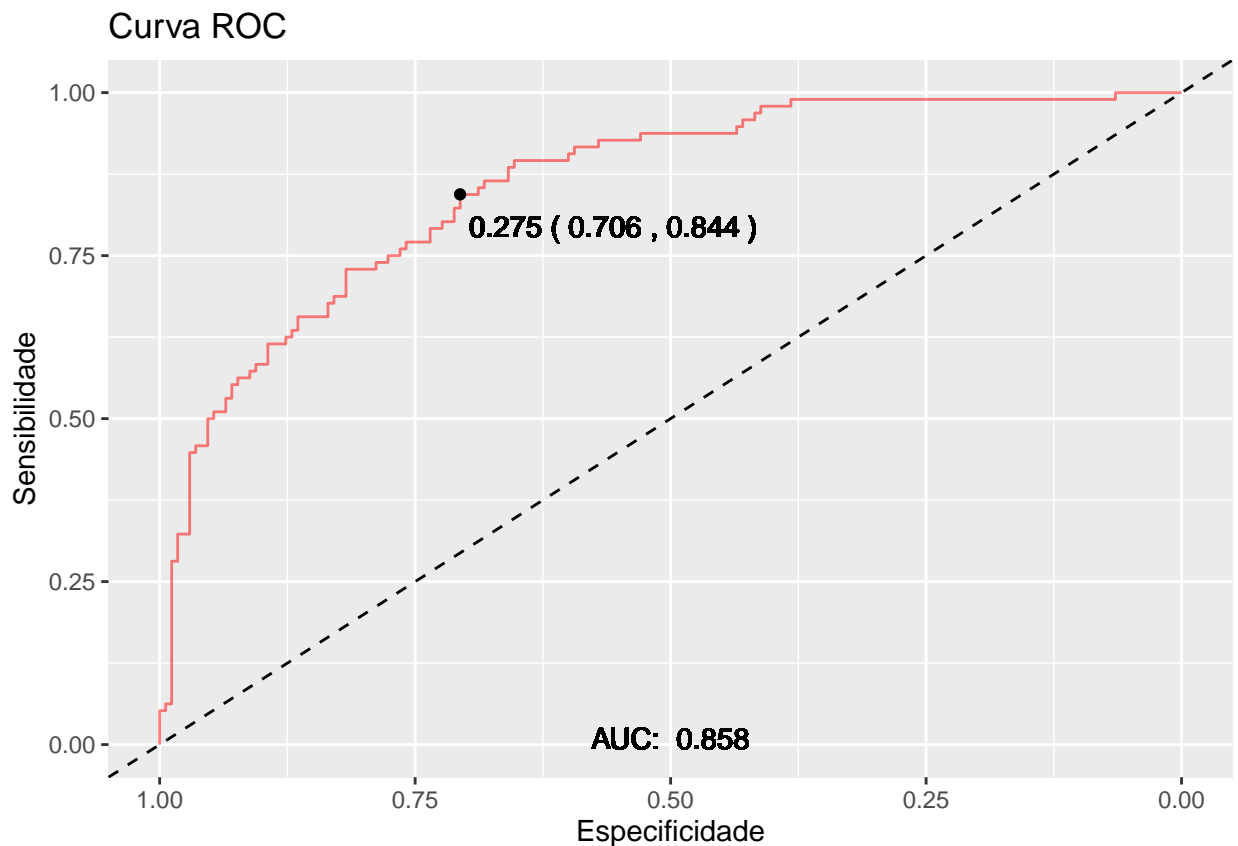
```
## Summary of sample sizes: 240, 240, 239, 239, 239, 239, ...
```

```
## Resampling results:
##
##   Accuracy   Kappa
##   0.7673789  0.4828079
```

```
rocRL2 <- roc(response = train_without_NAs$diabetes, predictor = predict(modeloRL2, train_without_NAs,
```

```
## Setting levels: control = No, case = Yes
## Setting direction: controls < cases
```

```
plotaroc(rocRL2)
```



```
## Logística regularizada / dados imputados
```

```
set.seed(23)
```

```
modeloRL3 <- caret::train(diabetes ~ ., data = train, trControl = train.control, method = "regLogistic",
```

```
print(modeloRL3)
```

```
## Regularized Logistic Regression
```

```
##
```

```
## 537 samples
```

```
## 8 predictor
```

```
## 2 classes: 'No', 'Yes'
```

```
##
```

```

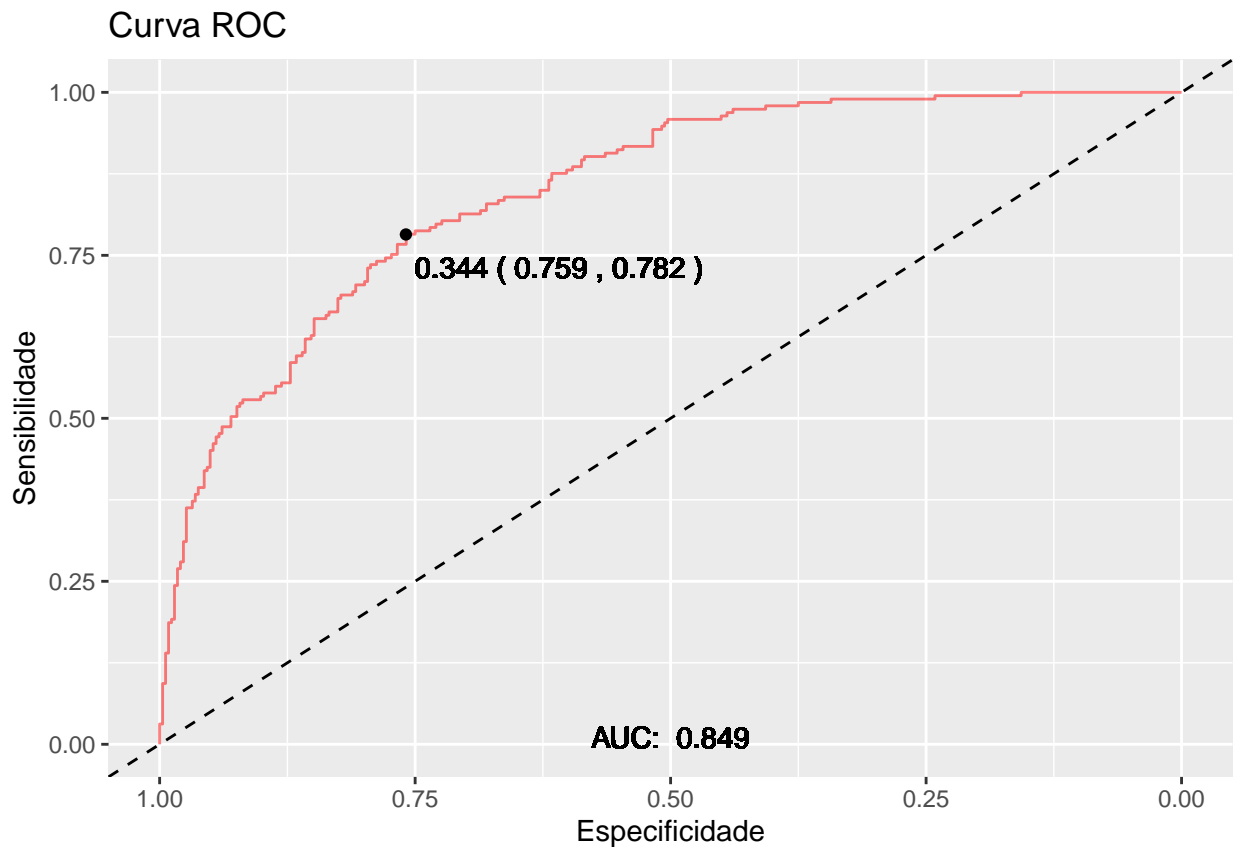
## No pre-processing
## Resampling: Cross-Validated (10 fold)
## Summary of sample sizes: 483, 483, 484, 483, 482, 484, ...
## Resampling results across tuning parameters:
##
##   cost  loss      epsilon  Accuracy  Kappa
##   0.5   L1       0.001    0.7482765 0.4294592
##   0.5   L1       0.010    0.7426847 0.4156713
##   0.5   L1       0.100    0.7482053 0.4254680
##   0.5   L2_dual  0.001    0.6961737 0.2507155
##   0.5   L2_dual  0.010    0.6813925 0.2298081
##   0.5   L2_dual  0.100    0.6217813 0.1749702
##   0.5   L2_primal 0.001    0.7445715 0.4210244
##   0.5   L2_primal 0.010    0.6998126 0.3193385
##   0.5   L2_primal 0.100    0.6964195 0.3018360
##   1.0   L1       0.001    0.7558236 0.4469887
##   1.0   L1       0.010    0.7539369 0.4431172
##   1.0   L1       0.100    0.7426161 0.4055951
##   1.0   L2_dual  0.001    0.6039839 0.1296527
##   1.0   L2_dual  0.010    0.6144038 0.2065002
##   1.0   L2_dual  0.100    0.6311416 0.1155712
##   1.0   L2_primal 0.001    0.7464583 0.4229977
##   1.0   L2_primal 0.010    0.6998126 0.3193385
##   1.0   L2_primal 0.100    0.6964195 0.3018360
##   2.0   L1       0.001    0.7595623 0.4554919
##   2.0   L1       0.010    0.7558236 0.4468964
##   2.0   L1       0.100    0.7538670 0.4388071
##   2.0   L2_dual  0.001    0.6149863 0.1897020
##   2.0   L2_dual  0.010    0.5702999 0.1900379
##   2.0   L2_dual  0.100    0.5902649 0.1230814
##   2.0   L2_primal 0.001    0.7500934 0.4318992
##   2.0   L2_primal 0.010    0.7070802 0.3426133
##   2.0   L2_primal 0.100    0.6964195 0.3018360
##
## Accuracy was used to select the optimal model using the largest value.
## The final values used for the model were cost = 2, loss = L1 and epsilon
##   = 0.001.

rocRL3 <- roc(response = train$diabetes, predictor = predict(modeloRL3, train, type = "prob"),[,2])

## Setting levels: control = No, case = Yes
## Setting direction: controls < cases

plotaroc(rocRL3)

```



```
## Logística regularizada / dados sem missing
set.seed(23)
modeloRL4 <- caret::train(diabetes ~ ., data = train_without_NAs, trControl = train.control, method = "glmnet")
print(modeloRL4)
```

```
## Regularized Logistic Regression
##
## 266 samples
## 8 predictor
## 2 classes: 'No', 'Yes'
##
## No pre-processing
## Resampling: Cross-Validated (10 fold)
## Summary of sample sizes: 240, 240, 239, 239, 239, 239, ...
## Resampling results across tuning parameters:
##
## cost loss epsilon Accuracy Kappa
## 0.5 L1 0.001 0.7598291 0.4596795
## 0.5 L1 0.010 0.7636752 0.4669602
## 0.5 L1 0.100 0.7488604 0.4299137
## 0.5 L2_dual 0.001 0.6394587 0.1554546
## 0.5 L2_dual 0.010 0.6065527 0.1369186
## 0.5 L2_dual 0.100 0.6501425 0.1759711
## 0.5 L2_primal 0.001 0.7263533 0.3848223
## 0.5 L2_primal 0.010 0.7149573 0.3616567
```



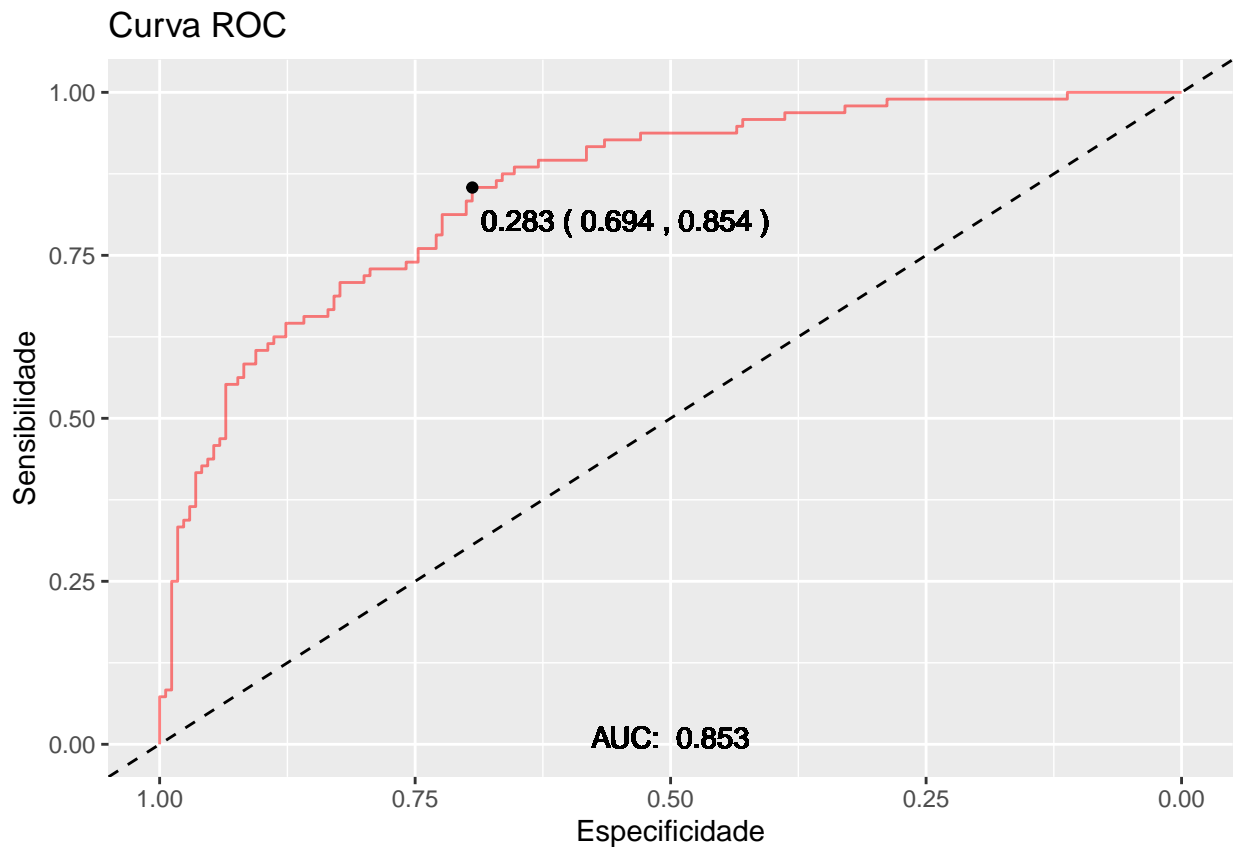
```
## 0.5 L2_primal 0.100 0.6807692 0.2740010
## 1.0 L1 0.001 0.7709402 0.4834174
## 1.0 L1 0.010 0.7747863 0.4903988
## 1.0 L1 0.100 0.7522792 0.4352616
## 1.0 L2_dual 0.001 0.6249288 0.1842344
## 1.0 L2_dual 0.010 0.5915954 0.1786622
## 1.0 L2_dual 0.100 0.6173789 0.1770171
## 1.0 L2_primal 0.001 0.7565527 0.4527102
## 1.0 L2_primal 0.010 0.7339031 0.4037206
## 1.0 L2_primal 0.100 0.6807692 0.2740010
## 2.0 L1 0.001 0.7710826 0.4902196
## 2.0 L1 0.010 0.7747863 0.4970394
## 2.0 L1 0.100 0.7712251 0.4816945
## 2.0 L2_dual 0.001 0.5992877 0.1660801
## 2.0 L2_dual 0.010 0.6019943 0.2116899
## 2.0 L2_dual 0.100 0.6427350 0.1862283
## 2.0 L2_primal 0.001 0.7601140 0.4623563
## 2.0 L2_primal 0.010 0.7487179 0.4343284
## 2.0 L2_primal 0.100 0.6807692 0.2740010
##
```

```
## Accuracy was used to select the optimal model using the largest value.
## The final values used for the model were cost = 1, loss = L1 and epsilon = 0.01.
```

```
rocRL4 <- roc(response = train_without_NAs$diabetes, predictor = predict(modeloRL4, train_without_NAs, "
```

```
## Setting levels: control = No, case = Yes
## Setting direction: controls < cases
```

```
plotaroc(rocRL4)
```



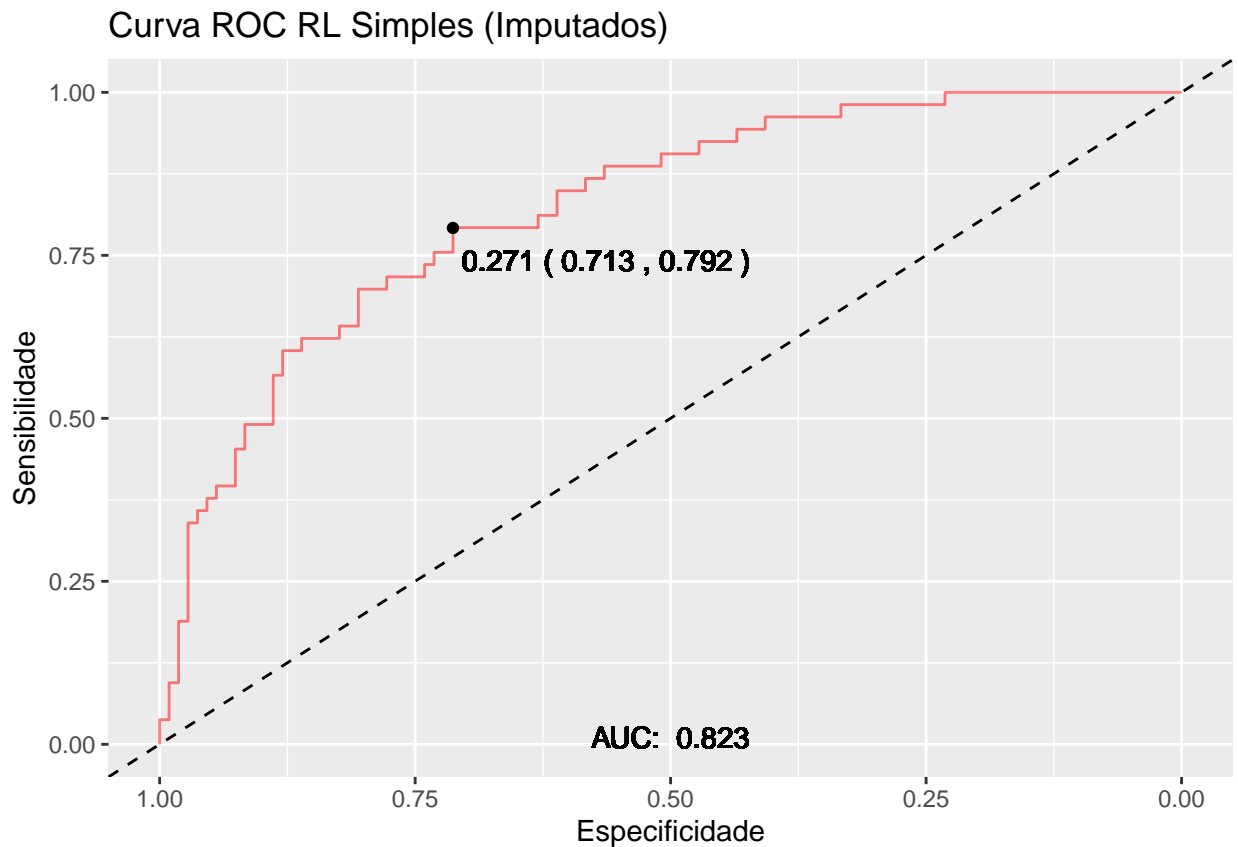
Teste

```
## Regressão Logística Simples com dados imputados
rocRL1 <- roc(response = test$diabetes, predictor = predict(modeloRL1, test, type = "prob"),[2])

## Setting levels: control = No, case = Yes

## Setting direction: controls < cases

plotaroc(rocRL1, titulo = "Curva ROC RL Simples (Imputados)")
```



```
predictRL1 <- predict(modeloRL1, newdata = test)
confusionMatrix(predictRL1, test$diabetes)
```

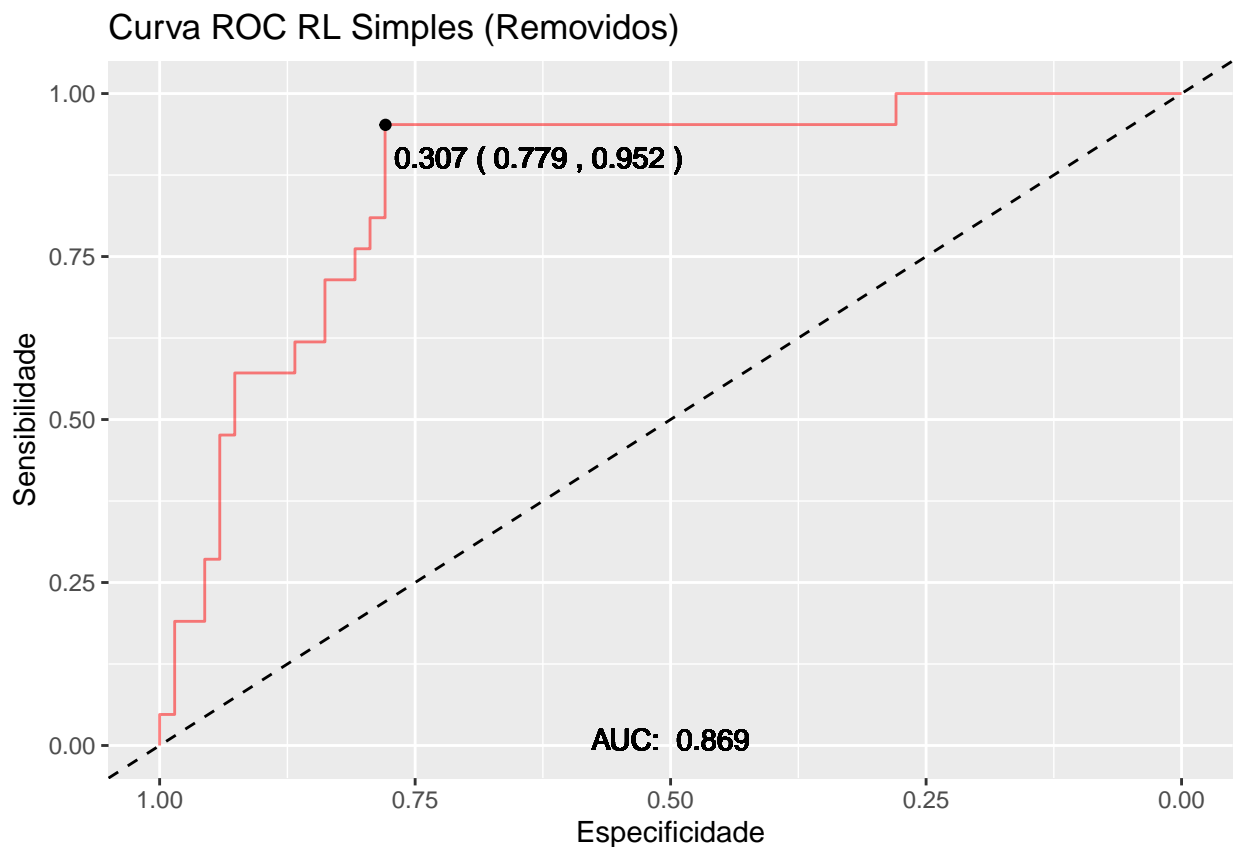
```
## Confusion Matrix and Statistics
##
##           Reference
## Prediction No Yes
##      No  99  27
##      Yes   9  26
##
##           Accuracy : 0.7764
##           95% CI : (0.7041, 0.8382)
##      No Information Rate : 0.6708
##      P-Value [Acc > NIR] : 0.002203
##
##           Kappa : 0.4458
##
##  Mcnemar's Test P-Value : 0.004607
##
##           Sensitivity : 0.9167
##           Specificity : 0.4906
##      Pos Pred Value : 0.7857
##      Neg Pred Value : 0.7429
##           Prevalence : 0.6708
##      Detection Rate : 0.6149
```

```
## Detection Prevalence : 0.7826
## Balanced Accuracy : 0.7036
##
## 'Positive' Class : No
##
```

```
## Regressão Logística Simples sem missing
rocRL2 <- roc(response = test_without_NAs$diabetes, predictor = predict(modeloRL2, test_without_NAs, type = "probabilities"))
```

```
## Setting levels: control = No, case = Yes
## Setting direction: controls < cases
```

```
plotaroc(rocRL2, titulo = "Curva ROC RL Simples (Removidos)")
```



```
predictRL2 <- predict(modeloRL2, newdata = test_without_NAs)
confusionMatrix(predictRL2, test_without_NAs$diabetes)
```

```
## Confusion Matrix and Statistics
##
##           Reference
## Prediction No Yes
##           No 64 11
##           Yes 4 10
```

```

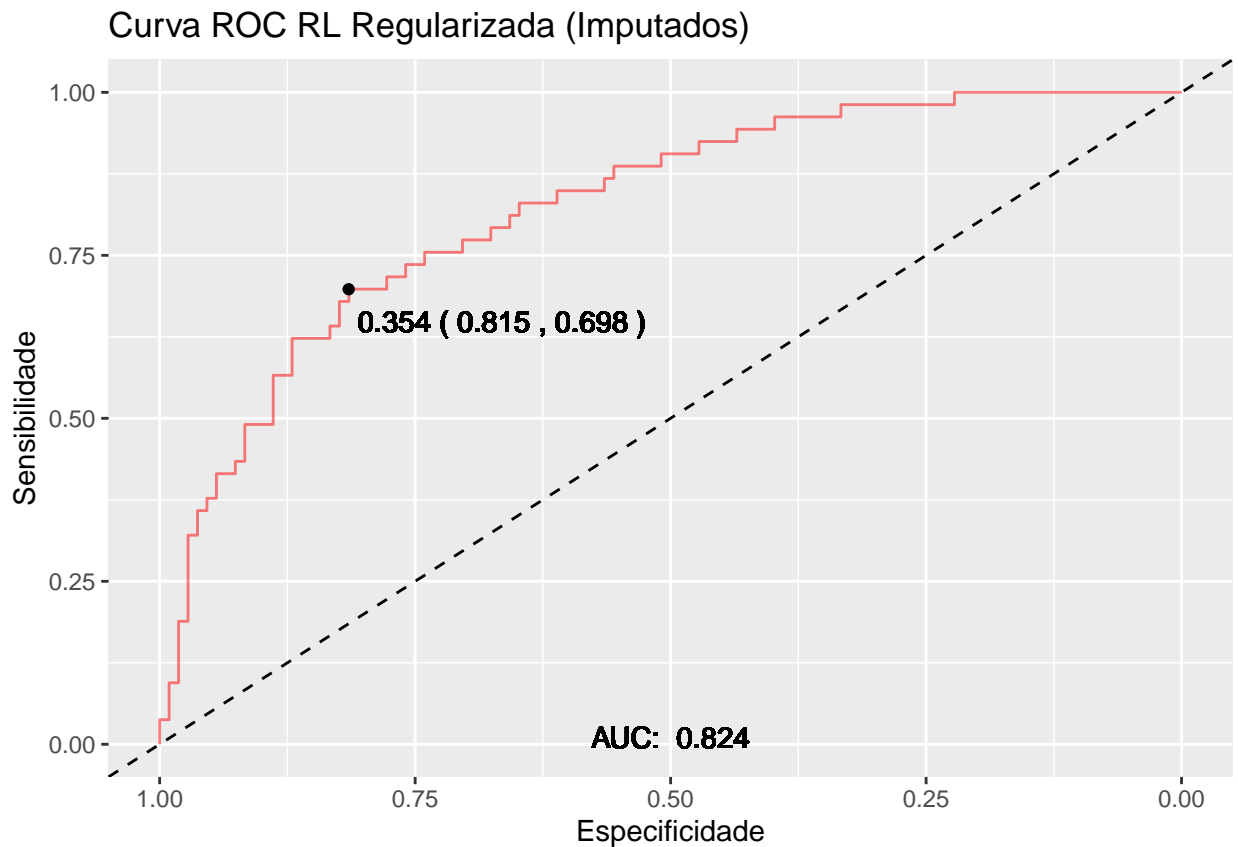
##
##           Accuracy : 0.8315
##           95% CI : (0.7373, 0.9025)
##      No Information Rate : 0.764
##      P-Value [Acc > NIR] : 0.08127
##
##           Kappa : 0.4717
##
##      McNemar's Test P-Value : 0.12134
##
##           Sensitivity : 0.9412
##           Specificity : 0.4762
##      Pos Pred Value : 0.8533
##      Neg Pred Value : 0.7143
##           Prevalence : 0.7640
##      Detection Rate : 0.7191
##      Detection Prevalence : 0.8427
##      Balanced Accuracy : 0.7087
##
##      'Positive' Class : No
##

## Regressão Logística Regularizada com dados imputados
rocRL3 <- roc(response = test$diabetes, predictor = predict(modeloRL3, test, type = "prob"),2)

## Setting levels: control = No, case = Yes
## Setting direction: controls < cases

plotaroc(rocRL3, titulo = "Curva ROC RL Regularizada (Imputados)")

```



```
predictRL3 <- predict(modeloRL3, newdata = test)
confusionMatrix(predictRL3, test$diabetes)
```

```
## Confusion Matrix and Statistics
##
##           Reference
## Prediction No Yes
##      No  99  28
##      Yes   9  25
##
##           Accuracy : 0.7702
##           95% CI : (0.6974, 0.8327)
##      No Information Rate : 0.6708
##      P-Value [Acc > NIR] : 0.003815
##
##           Kappa : 0.4274
##
##  Mcnemar's Test P-Value : 0.003085
##
##           Sensitivity : 0.9167
##           Specificity : 0.4717
##      Pos Pred Value : 0.7795
##      Neg Pred Value : 0.7353
##           Prevalence : 0.6708
##      Detection Rate : 0.6149
```

```
## Detection Prevalence : 0.7888
## Balanced Accuracy : 0.6942
##
## 'Positive' Class : No
##
```

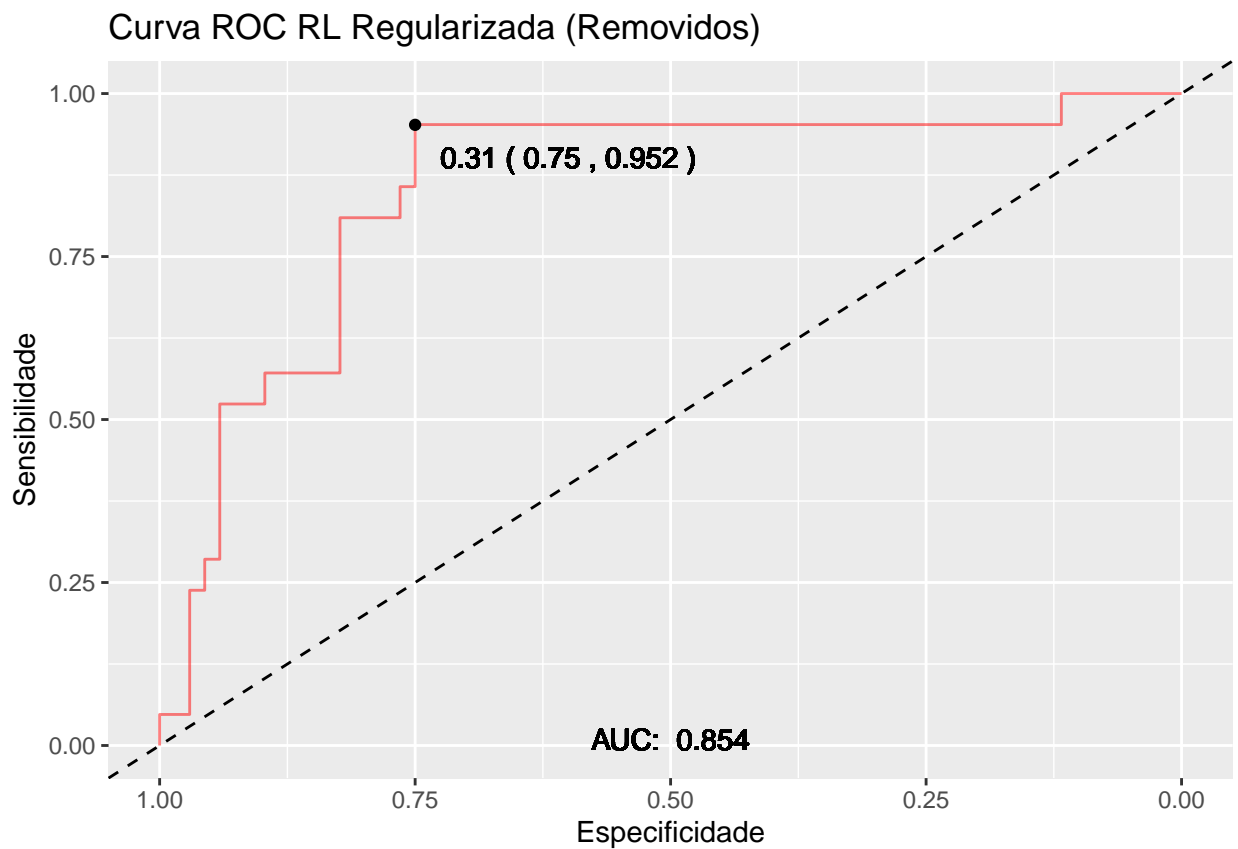
```
## Regressão Logística Regularizada sem missing
```

```
rocRL4 <- roc(response = test_without_NAs$diabetes, predictor = predict(modeloRL4, test_without_NAs, type = "prob"))
```

```
## Setting levels: control = No, case = Yes
```

```
## Setting direction: controls < cases
```

```
plotaroc(rocRL4, titulo = "Curva ROC RL Regularizada (Removidos)")
```



```
predictRL4 <- predict(modeloRL4, newdata = test_without_NAs)
confusionMatrix(predictRL4, test_without_NAs$diabetes)
```

```
## Confusion Matrix and Statistics
```

```
##
```

```
## Reference
```

```
## Prediction No Yes
```

```
## No 64 11
```

```
## Yes 4 10
```

```
##
##           Accuracy : 0.8315
##           95% CI : (0.7373, 0.9025)
##      No Information Rate : 0.764
##      P-Value [Acc > NIR] : 0.08127
##
##           Kappa : 0.4717
##
##  Mcnemar's Test P-Value : 0.12134
##
##      Sensitivity : 0.9412
##      Specificity : 0.4762
##      Pos Pred Value : 0.8533
##      Neg Pred Value : 0.7143
##      Prevalence : 0.7640
##      Detection Rate : 0.7191
##      Detection Prevalence : 0.8427
##      Balanced Accuracy : 0.7087
##
##      'Positive' Class : No
##
```

Random Forest

Modelagem

```
library(caret)

train.control <- caret::trainControl(method = "cv", number = 10) # Cross-validation com k=10

## Random Forest / dados imputados
set.seed(23)
tuneGrid <- expand.grid(.mtry = c(1: 10))
modeloRF1 <- train(diabetes~.,
  train,
  method = "rf",
  metric = "Accuracy",
  tuneGrid = tuneGrid,
  trControl = train.control,
  importance = TRUE,
  ntree = 1000)

## Warning in randomForest.default(x, y, mtry = param$mtry, ...): invalid mtry:
## reset to within valid range

## Warning in randomForest.default(x, y, mtry = param$mtry, ...): invalid mtry:
## reset to within valid range

## Warning in randomForest.default(x, y, mtry = param$mtry, ...): invalid mtry:
## reset to within valid range
```



```

## Random Forest
##
## 537 samples
## 8 predictor
## 2 classes: 'No', 'Yes'
##
## No pre-processing
## Resampling: Cross-Validated (10 fold)
## Summary of sample sizes: 483, 483, 484, 483, 482, 484, ...
## Resampling results across tuning parameters:
##
##  mtry  Accuracy  Kappa
##  1     0.7506099 0.4361204
##  2     0.7562004 0.4559871
##  3     0.7468001 0.4344252
##  4     0.7486869 0.4382093
##  5     0.7487218 0.4409065
##  6     0.7505387 0.4457852
##  7     0.7523569 0.4506149
##  8     0.7468001 0.4377380
##  9     0.7431300 0.4295050
## 10     0.7487567 0.4392488
##
## Accuracy was used to select the optimal model using the largest value.
## The final value used for the model was mtry = 2.

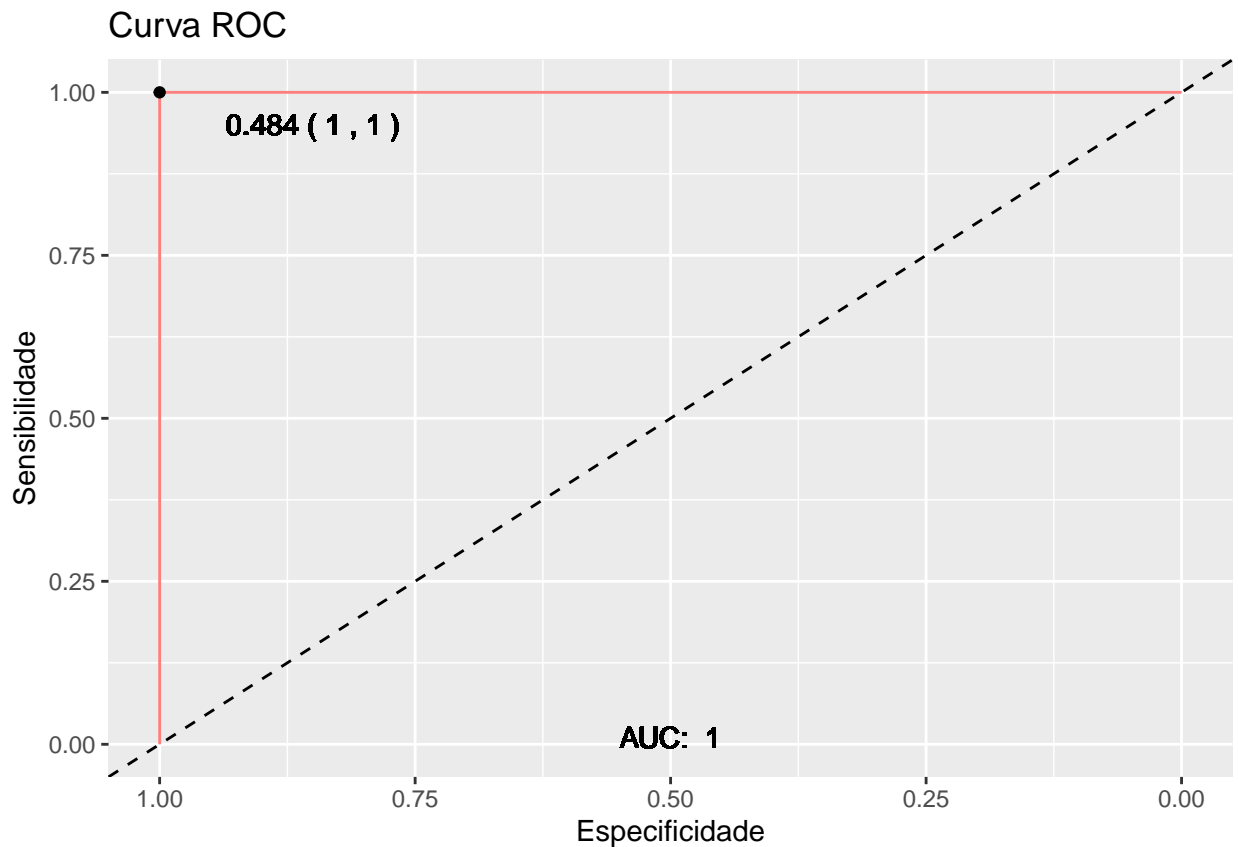
rocRF1 <- roc(response = train$diabetes, predictor = predict(modeloRF1, train, type = "prob"),[,2])

## Setting levels: control = No, case = Yes

## Setting direction: controls < cases

plotaroc(rocRF1)

```



```
## Random Forest / sem missing
set.seed(23)
tuneGrid <- expand.grid(.mtry = c(1: 10))
modeloRF2 <- train(diabetes~.,
  train_without_NAs,
  method = "rf",
  metric = "Accuracy",
  tuneGrid = tuneGrid,
  trControl = train.control,
  importance = TRUE,
  ntree = 1000)
```

```
## Warning in randomForest.default(x, y, mtry = param$mtry, ...): invalid mtry:
## reset to within valid range
```

```
## Warning in randomForest.default(x, y, mtry = param$mtry, ...): invalid mtry:
## reset to within valid range
```

```
## Warning in randomForest.default(x, y, mtry = param$mtry, ...): invalid mtry:
## reset to within valid range
```

```
## Warning in randomForest.default(x, y, mtry = param$mtry, ...): invalid mtry:
## reset to within valid range
```

```
## Warning in randomForest.default(x, y, mtry = param$mtry, ...): invalid mtry:
```



```
## 2 classes: 'No', 'Yes'
##
## No pre-processing
## Resampling: Cross-Validated (10 fold)
## Summary of sample sizes: 240, 240, 239, 239, 239, 239, ...
## Resampling results across tuning parameters:
```

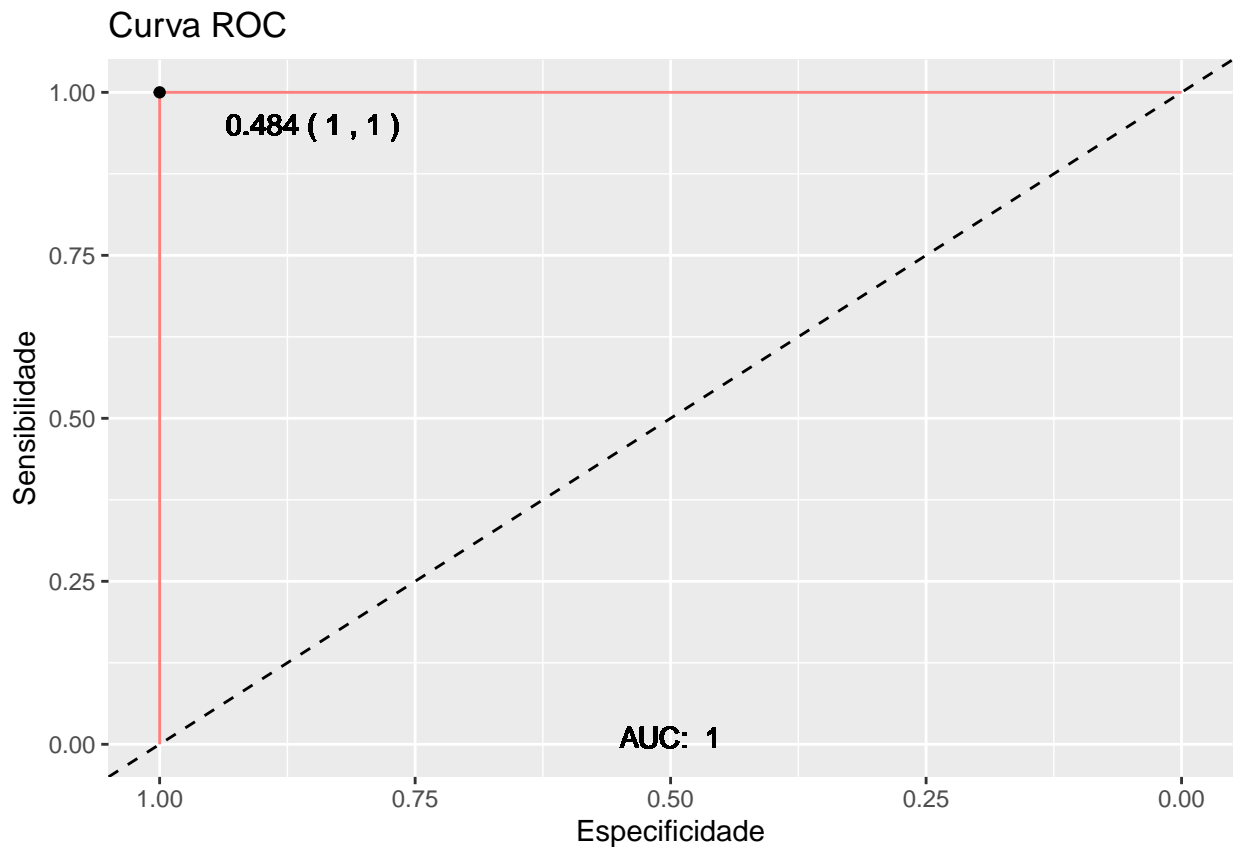
```
##
## mtry Accuracy Kappa
## 1 0.7750712 0.4927747
## 2 0.7787749 0.5063822
## 3 0.7903134 0.5363471
## 4 0.7938746 0.5468630
## 5 0.7826211 0.5182975
## 6 0.7827635 0.5166230
## 7 0.7864672 0.5255357
## 8 0.7827635 0.5201542
## 9 0.7904558 0.5372170
## 10 0.7903134 0.5347168
```

```
##
## Accuracy was used to select the optimal model using the largest value.
## The final value used for the model was mtry = 4.
```

```
rocRF2 <- roc(response = train_without_NAs$diabetes, predictor = predict(modeloRF1, train_without_NAs, "
```

```
## Setting levels: control = No, case = Yes
## Setting direction: controls < cases
```

```
plotaroc(rocRF2)
```



Árvores de decisão geradas

```
tree_num <- which(modeloRF1$finalModel$forest$ndbigtree == min(modeloRF1$finalModel$forest$ndbigtree))
print(paste("Menor árvore com imputação", tree_num))
```

```
## [1] "Menor árvore com imputação 876"
```

```
tree_func(final_model = modeloRF1$finalModel, tree_num[1])
```

```
## Warning: Duplicated aesthetics after name standardisation: na.rm
```

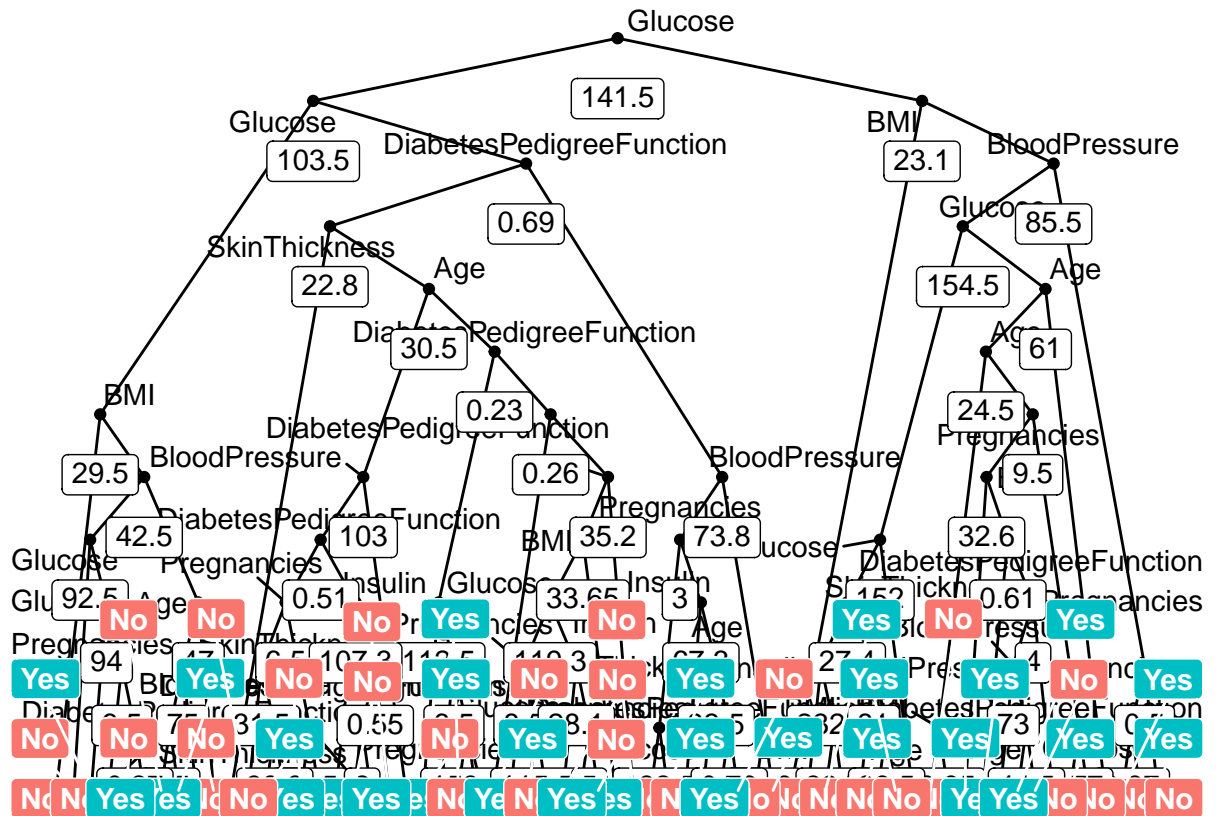
```
## Warning: Duplicated aesthetics after name standardisation: na.rm
```

```
## Warning: Duplicated aesthetics after name standardisation: na.rm
```

```
## Warning: Removed 64 rows containing missing values (geom_text_repel).
```

```
## Warning: Removed 64 rows containing missing values (geom_label).
```

```
## Warning: Removed 63 rows containing missing values (geom_label_repel).
```



```
tree_num <- which(modeloRF2$finalModel$forest$ndbigtree == min(modeloRF2$finalModel$forest$ndbigtree))
print(paste("Menor árvore sem imputação:", tree_num))
```

```
## [1] "Menor árvore sem imputação: 250"
```

```
tree_func(final_model = modeloRF2$finalModel, tree_num[1])
```

```
## Warning: Duplicated aesthetics after name standardisation: na.rm
```

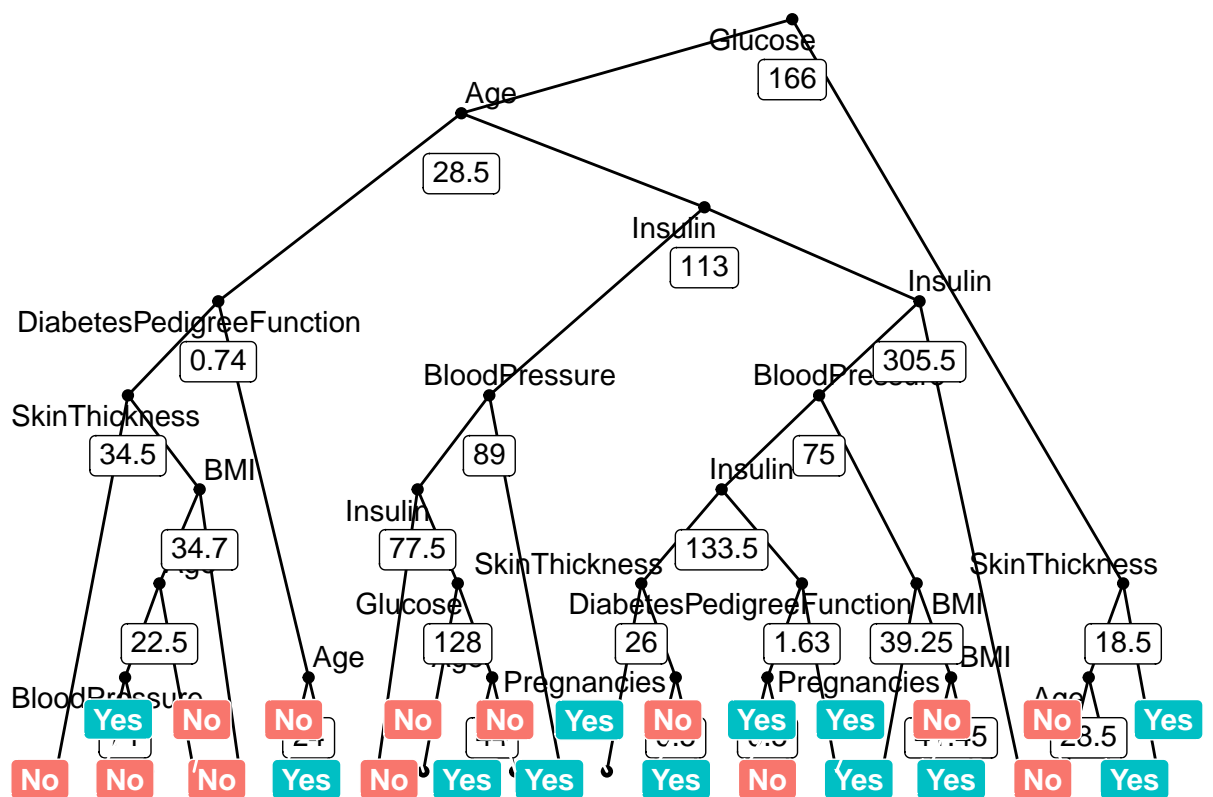
```
## Warning: Duplicated aesthetics after name standardisation: na.rm
```

```
## Warning: Duplicated aesthetics after name standardisation: na.rm
```

```
## Warning: Removed 25 rows containing missing values (geom_text_repel).
```

```
## Warning: Removed 25 rows containing missing values (geom_label).
```

```
## Warning: Removed 24 rows containing missing values (geom_label_repel).
```



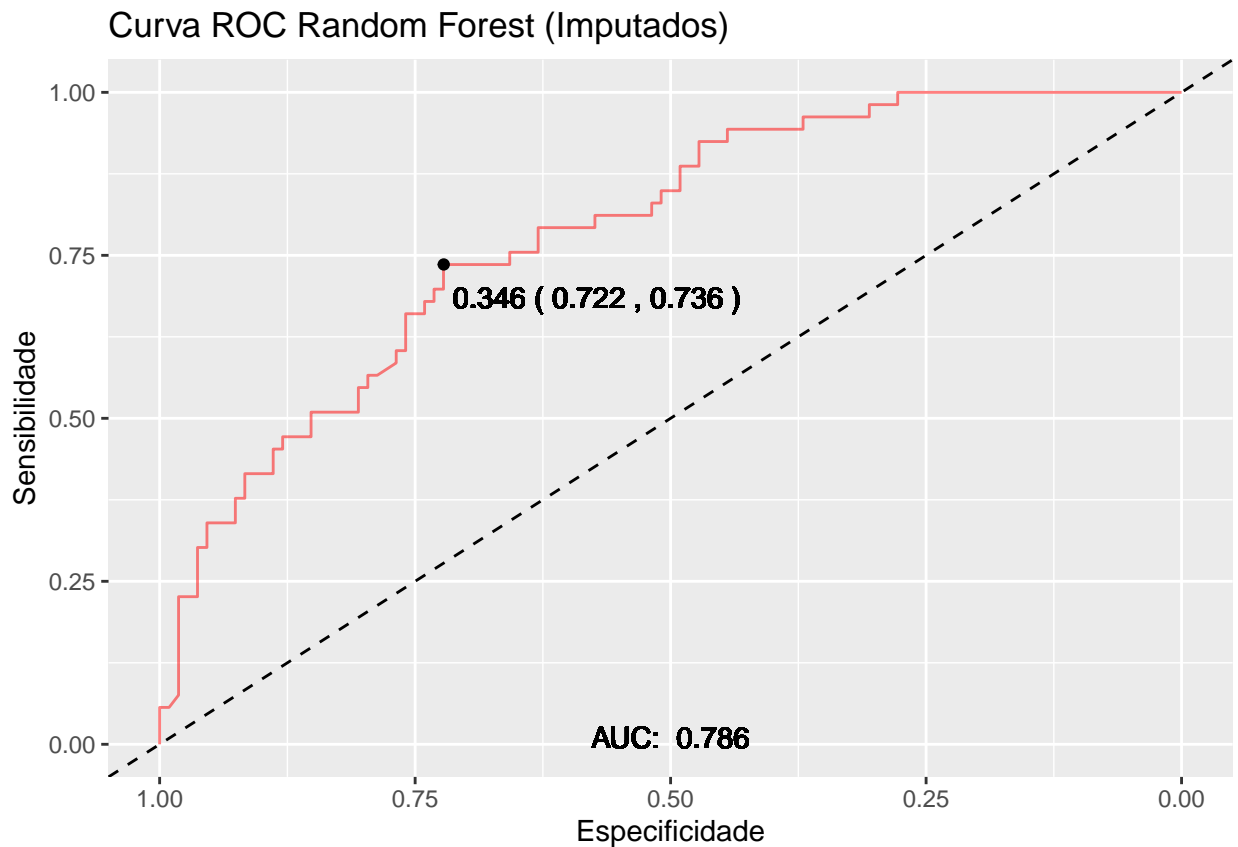
Teste

```
## Random Forest com dados imputados
rocRF1 <- roc(response = test$diabetes, predictor = predict(modeloRF1, test, type = "prob")[,2])

## Setting levels: control = No, case = Yes

## Setting direction: controls < cases

plotaroc(rocRF1, titulo = "Curva ROC Random Forest (Imputados)")
```

```
predictRF1 <- predict(modeloRF1, newdata = test)
print("Dados imputados:")
```

```
## [1] "Dados imputados:"
```

```
confusionMatrix(predictRF1, test$diabetes)
```

```
## Confusion Matrix and Statistics
```

```
##
```

```
##           Reference
```

```
## Prediction No Yes
```

```
##           No  93  28
```

```
##           Yes  15  25
```

```
##
```

```
##           Accuracy : 0.7329
```

```
##           95% CI : (0.6576, 0.7995)
```

```
## No Information Rate : 0.6708
```

```
## P-Value [Acc > NIR] : 0.05368
```

```
##
```

```
##           Kappa : 0.355
```

```
##
```

```
## McNemar's Test P-Value : 0.06725
```

```
##
```

```
##           Sensitivity : 0.8611
```

```
##          Specificity : 0.4717
##          Pos Pred Value : 0.7686
##          Neg Pred Value : 0.6250
##          Prevalence : 0.6708
##          Detection Rate : 0.5776
##          Detection Prevalence : 0.7516
##          Balanced Accuracy : 0.6664
##
##          'Positive' Class : No
##
```

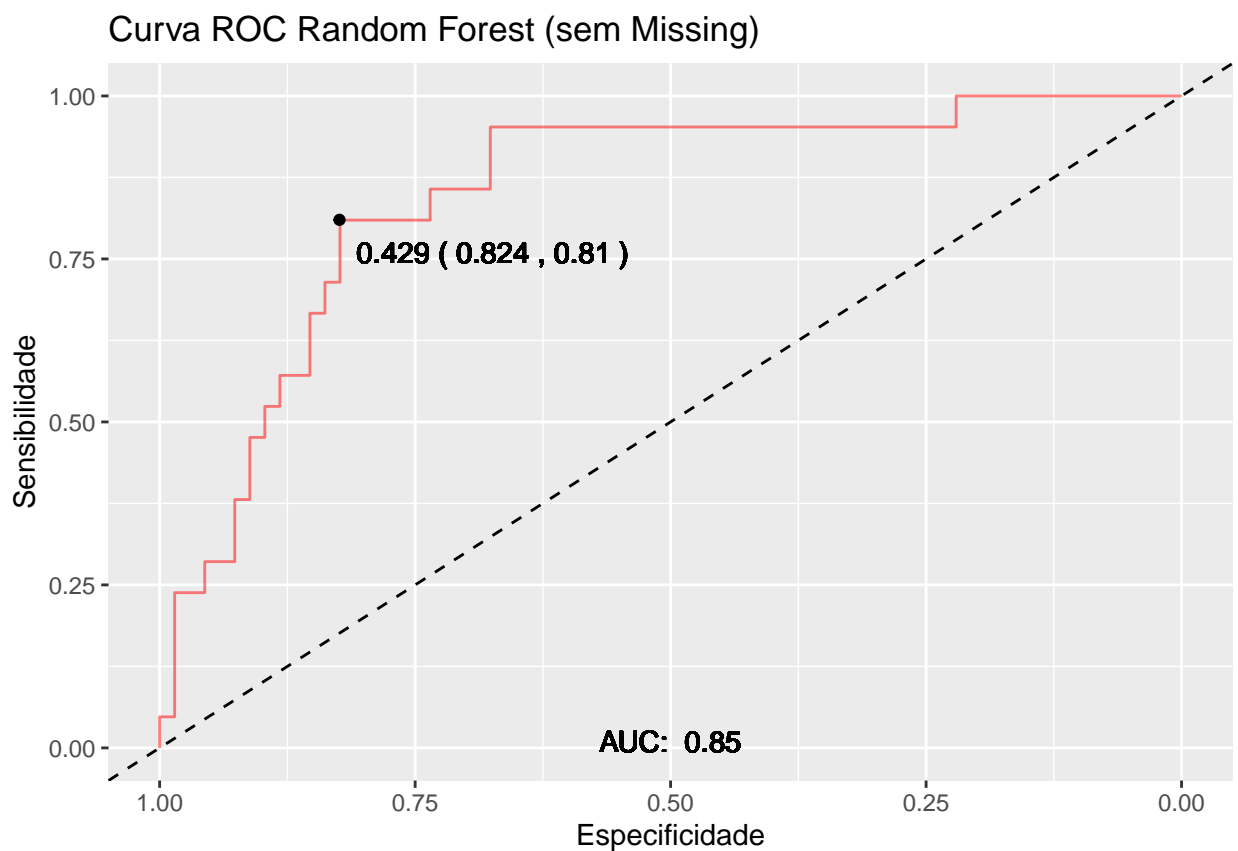
```
## Random Forest com dados imputados
```

```
rocRF2 <- roc(response = test_without_NAs$diabetes, predictor = predict(modeloRF2, test_without_NAs, type = "probabilities"))
```

```
## Setting levels: control = No, case = Yes
```

```
## Setting direction: controls < cases
```

```
plotaroc(rocRF2, titulo = "Curva ROC Random Forest (sem Missing)")
```



```
predictRF2 <- predict(modeloRF2, newdata = test_without_NAs)
print("Sem Missing:")
```

```
## [1] "Sem Missing:"
```

```
confusionMatrix(predictRF2, test_without_NAs$diabetes)
```

```
## Confusion Matrix and Statistics
##
##              Reference
## Prediction No Yes
##           No  59   9
##           Yes   9  12
##
##              Accuracy : 0.7978
##              95% CI : (0.6993, 0.8755)
##           No Information Rate : 0.764
##           P-Value [Acc > NIR] : 0.2709
##
##              Kappa : 0.4391
##
##  Mcnemar's Test P-Value : 1.0000
##
##           Sensitivity : 0.8676
##           Specificity : 0.5714
##           Pos Pred Value : 0.8676
##           Neg Pred Value : 0.5714
##           Prevalence : 0.7640
##           Detection Rate : 0.6629
##           Detection Prevalence : 0.7640
##           Balanced Accuracy : 0.7195
##
##           'Positive' Class : No
##
```

Support Vector Machine

Support Vector Machine (SVM) estabelece um limite de decisão ideal que separa os pontos de dados de diferentes classes e, em seguida, prevê a classe de novas observações com base nesse limite de separação. Os diferentes grupos podem ser separáveis por uma linha reta linear ou por uma linha limite não linear.

Modelagem

SVM Kernel Linear com dados imputados

```
train.control <- caret::trainControl(method = "cv", number = 10, savePred=T, classProb=T) # Cross-validation
#library(e1071) #SVM
## SVM Kernel Linear com dados imputados
set.seed(23)
modeloSVM1 <- caret::train(
  diabetes ~., data = train, method = "svmLinear",
  probability = T,
```

```

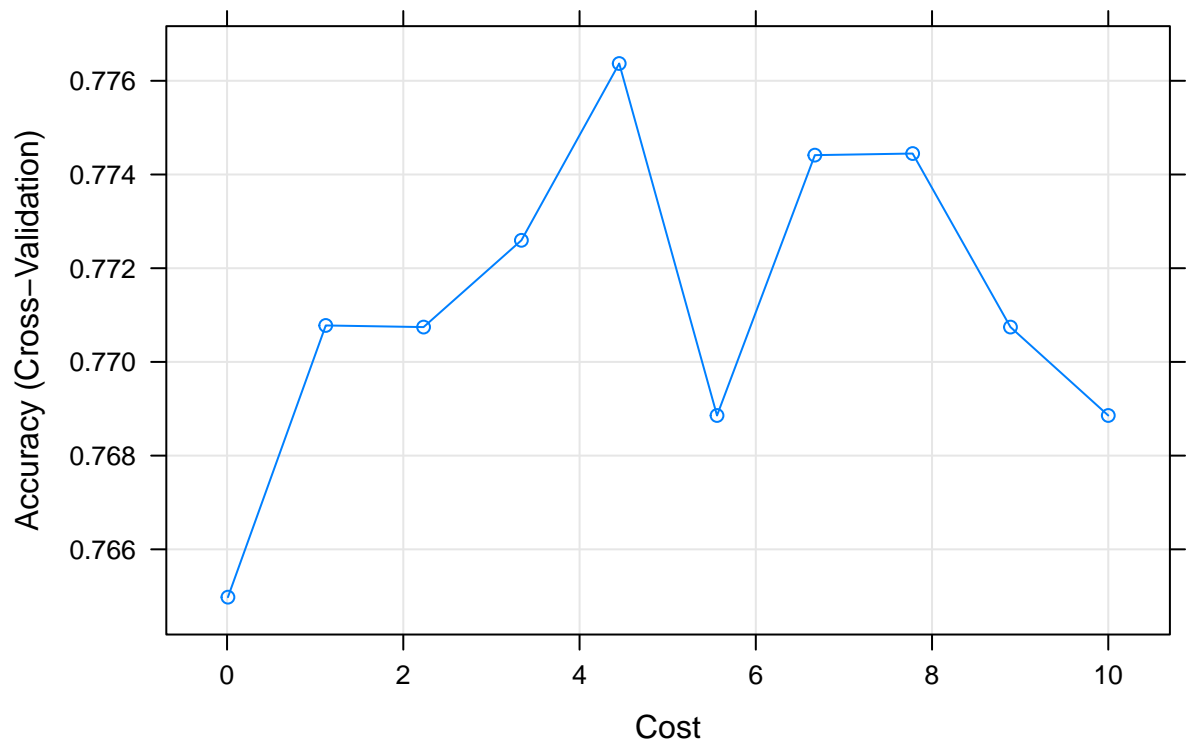
trControl = train.control,
preProcess = c("center", "scale"),
tuneGrid = expand.grid(C = seq(0.01, 10, length = 10))
)
print(modeloSVM1)

## Support Vector Machines with Linear Kernel
##
## 537 samples
## 8 predictor
## 2 classes: 'No', 'Yes'
##
## Pre-processing: centered (8), scaled (8)
## Resampling: Cross-Validated (10 fold)
## Summary of sample sizes: 483, 483, 484, 483, 482, 484, ...
## Resampling results across tuning parameters:
##
##  C      Accuracy  Kappa
##  0.01  0.7649794  0.4734524
##  1.12  0.7707782  0.4824355
##  2.23  0.7707433  0.4808278
##  3.34  0.7725951  0.4843473
##  4.45  0.7763687  0.4917503
##  5.56  0.7688565  0.4769505
##  6.67  0.7744133  0.4896157
##  7.78  0.7744470  0.4862920
##  8.89  0.7707433  0.4778763
## 10.00  0.7688565  0.4754528
##
## Accuracy was used to select the optimal model using the largest value.
## The final value used for the model was C = 4.45.

plot(modeloSVM1, main="SVM Kernel Linear - Acurácia vs Valores de Cost (Imputados)")

```

SVM Kernel Linear – Acurácia vs Valores de Cost (Imputados)

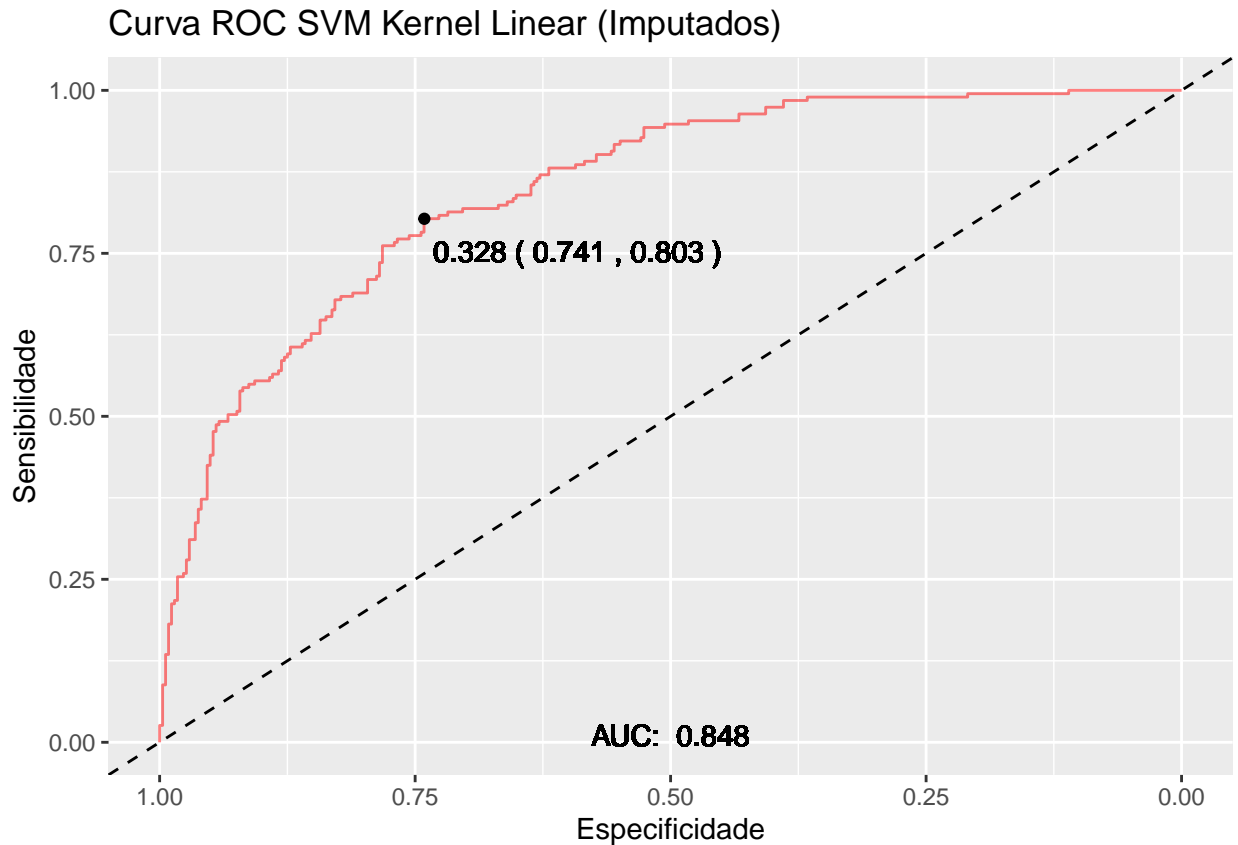


```
rocSVM1 <- roc(response = train$diabetes, predictor = predict(modeloSVM1, train, type = "prob")[,2])
```

```
## Setting levels: control = No, case = Yes
```

```
## Setting direction: controls < cases
```

```
plotaroc(rocSVM1, titulo = "Curva ROC SVM Kernel Linear (Imputados)")
```



SVM Kernel Linear sem missing

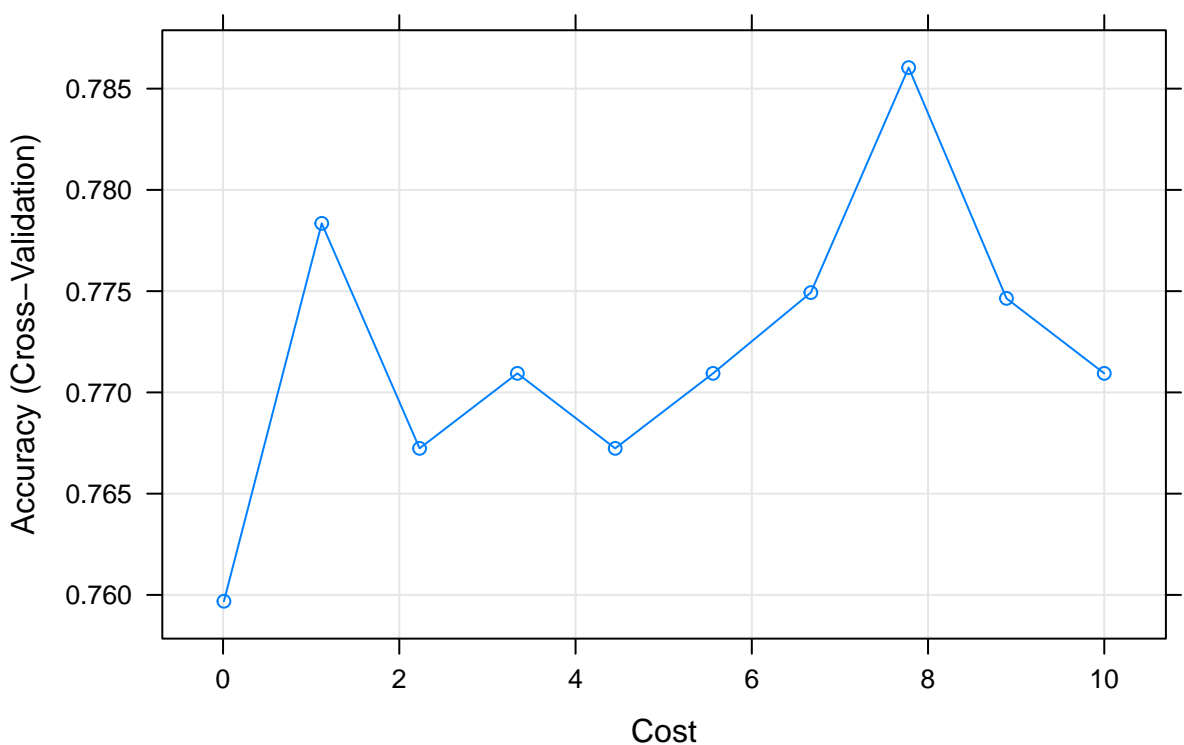
```
## SVM Kernel Linear sem missing
set.seed(23)
modeloSVM2 <- caret::train(
  diabetes ~., data = train_without_NAs, method = "svmLinear",
  trControl = train.control,
  preProcess = c("center", "scale"),
  tuneGrid = expand.grid(C = seq(0.01, 10, length = 10))
)
print(modeloSVM2)
```

```
## Support Vector Machines with Linear Kernel
##
## 266 samples
## 8 predictor
## 2 classes: 'No', 'Yes'
##
## Pre-processing: centered (8), scaled (8)
## Resampling: Cross-Validated (10 fold)
## Summary of sample sizes: 240, 240, 239, 239, 239, 239, ...
## Resampling results across tuning parameters:
##
```

```
##      C      Accuracy      Kappa
##    0.01 0.7596866 0.4629441
##    1.12 0.7783476 0.4980691
##    2.23 0.7672365 0.4744505
##    3.34 0.7709402 0.4839230
##    4.45 0.7672365 0.4774236
##    5.56 0.7709402 0.4781789
##    6.67 0.7749288 0.4893745
##    7.78 0.7860399 0.5103509
##    8.89 0.7746439 0.4906956
##   10.00 0.7709402 0.4754996
##
## Accuracy was used to select the optimal model using the largest value.
## The final value used for the model was C = 7.78.
```

```
plot(modeloSVM2, main="SVM Kernel Linear - Acurácia vs Valores de Cost (Removidos)")
```

SVM Kernel Linear – Acurácia vs Valores de Cost (Removidos)

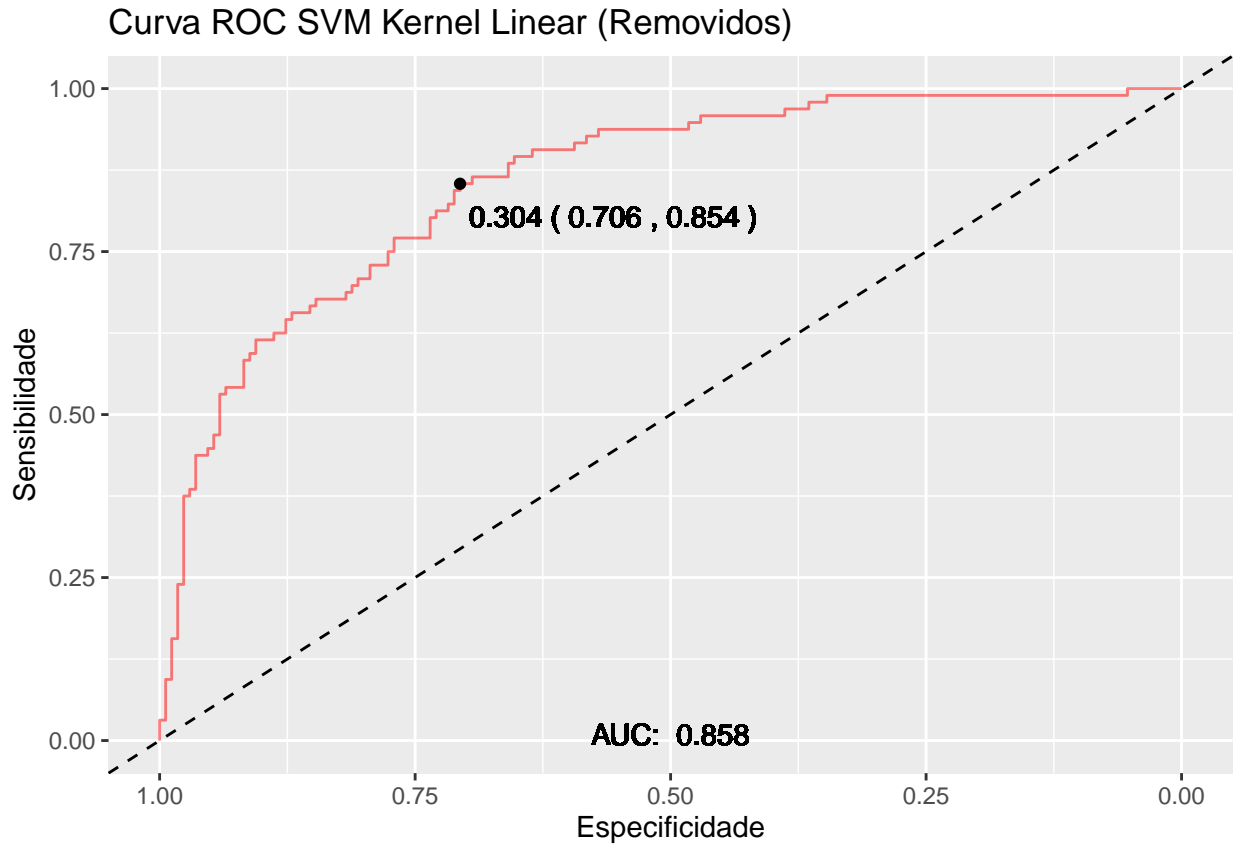


```
rocSVM2 <- roc(response = train_without_NAs$diabetes, predictor = predict(modeloSVM2, train_without_NAs
```

```
## Setting levels: control = No, case = Yes
```

```
## Setting direction: controls < cases
```

```
plotaroc(rocSVM2, titulo = "Curva ROC SVM Kernel Linear (Removidos)")
```



SVM Kernel Não Linear com dados imputados

```
train.control <- caret::trainControl(method = "cv", number = 10, savePred=T, classProb=T)

## SVM Kernel Não Linear com dados imputados
set.seed(23)
modeloSVM3 <- caret::train(
  diabetes ~., data = train, method = "svmRadial",
  trControl = train.control,
  preProcess = c("center", "scale"),
  tuneGrid = expand.grid(sigma = seq(0.01, 1, length = 10) , C = seq(1, 10, length = 10))
)
print(modeloSVM3)
```

```
## Support Vector Machines with Radial Basis Function Kernel
##
## 537 samples
## 8 predictor
## 2 classes: 'No', 'Yes'
```



```

##
## Pre-processing: centered (8), scaled (8)
## Resampling: Cross-Validated (10 fold)
## Summary of sample sizes: 483, 483, 484, 483, 482, 484, ...
## Resampling results across tuning parameters:
##
##   sigma C   Accuracy   Kappa
##   0.01  1  0.7632323  0.4654255
##   0.01  2  0.7651541  0.4686906
##   0.01  3  0.7633371  0.4652478
##   0.01  4  0.7578502  0.4499811
##   0.01  5  0.7596334  0.4565153
##   0.01  6  0.7578153  0.4521902
##   0.01  7  0.7597021  0.4544051
##   0.01  8  0.7652576  0.4649773
##   0.01  9  0.7652576  0.4667128
##   0.01 10  0.7652227  0.4665617
##   0.12  1  0.7410412  0.4172729
##   0.12  2  0.7409688  0.4130742
##   0.12  3  0.7428905  0.4189473
##   0.12  4  0.7429928  0.4173560
##   0.12  5  0.7410724  0.4173790
##   0.12  6  0.7298901  0.3891193
##   0.12  7  0.7280382  0.3836402
##   0.12  8  0.7280382  0.3865933
##   0.12  9  0.7206296  0.3666429
##   0.12 10  0.7186754  0.3627544
##   0.23  1  0.7316746  0.3933253
##   0.23  2  0.7225513  0.3738773
##   0.23  3  0.7113354  0.3457537
##   0.23  4  0.7057449  0.3323967
##   0.23  5  0.7056750  0.3313558
##   0.23  6  0.6982314  0.3163946
##   0.23  7  0.6983362  0.3096625
##   0.23  8  0.6870853  0.2825074
##   0.23  9  0.6944578  0.3009454
##   0.23 10  0.6870167  0.2782018
##   0.34  1  0.7224827  0.3714102
##   0.34  2  0.7019376  0.3273452
##   0.34  3  0.7036872  0.3239199
##   0.34  4  0.6926421  0.2970354
##   0.34  5  0.6963808  0.3111219
##   0.34  6  0.6889721  0.2906708
##   0.34  7  0.6850975  0.2810966
##   0.34  8  0.6814624  0.2675126
##   0.34  9  0.6890071  0.2828918
##   0.34 10  0.6759742  0.2551517
##   0.45  1  0.7150753  0.3509448
##   0.45  2  0.6963833  0.3062685
##   0.45  3  0.6908589  0.2926231
##   0.45  4  0.6945988  0.3106269
##   0.45  5  0.6928156  0.3041198
##   0.45  6  0.6889734  0.2984869
##   0.45  7  0.6816009  0.2875827

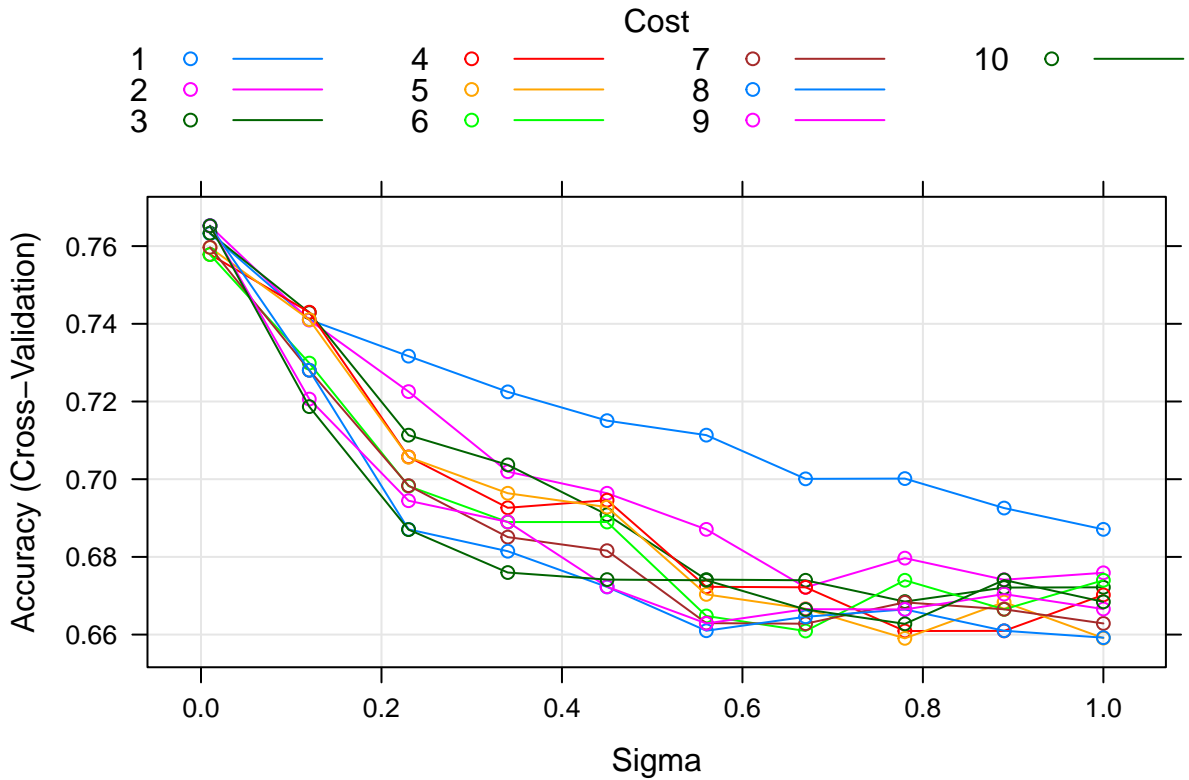
```

##	0.45	8	0.6722718	0.2590777
##	0.45	9	0.6723067	0.2552091
##	0.45	10	0.6741586	0.2616750
##	0.56	1	0.7113379	0.3428026
##	0.56	2	0.6870866	0.2825257
##	0.56	3	0.6741573	0.2575748
##	0.56	4	0.6723054	0.2584329
##	0.56	5	0.6703500	0.2495136
##	0.56	6	0.6647958	0.2365721
##	0.56	7	0.6629439	0.2385834
##	0.56	8	0.6609872	0.2203032
##	0.56	9	0.6628740	0.2280941
##	0.56	10	0.6739864	0.2501550
##	0.67	1	0.7000870	0.3084599
##	0.67	2	0.6721670	0.2416235
##	0.67	3	0.6739864	0.2500257
##	0.67	4	0.6721695	0.2490706
##	0.67	5	0.6666463	0.2338014
##	0.67	6	0.6609186	0.2177422
##	0.67	7	0.6628054	0.2275974
##	0.67	8	0.6645887	0.2323034
##	0.67	9	0.6665428	0.2321039
##	0.67	10	0.6664754	0.2375644
##	0.78	1	0.7001569	0.3088335
##	0.78	2	0.6796805	0.2585207
##	0.78	3	0.6684982	0.2323193
##	0.78	4	0.6609186	0.2142967
##	0.78	5	0.6590318	0.2111359
##	0.78	6	0.6739877	0.2470423
##	0.78	7	0.6682924	0.2377308
##	0.78	8	0.6664754	0.2266465
##	0.78	9	0.6664754	0.2258914
##	0.78	10	0.6627705	0.2125283
##	0.89	1	0.6925761	0.2908961
##	0.89	2	0.6741236	0.2431205
##	0.89	3	0.6721009	0.2334575
##	0.89	4	0.6609536	0.2093185
##	0.89	5	0.6683610	0.2275698
##	0.89	6	0.6665091	0.2235560
##	0.89	7	0.6665091	0.2188595
##	0.89	8	0.6609872	0.2072523
##	0.89	9	0.6704212	0.2236802
##	0.89	10	0.6740550	0.2476682
##	1.00	1	0.6870891	0.2753881
##	1.00	2	0.6759081	0.2379519
##	1.00	3	0.6721695	0.2306436
##	1.00	4	0.6702128	0.2300421
##	1.00	5	0.6591017	0.2023119
##	1.00	6	0.6739864	0.2383797
##	1.00	7	0.6628728	0.2097380
##	1.00	8	0.6592040	0.2043506
##	1.00	9	0.6665790	0.2159346
##	1.00	10	0.6684308	0.2258624
##				

```
## Accuracy was used to select the optimal model using the largest value.
## The final values used for the model were sigma = 0.01 and C = 9.
```

```
plot(modeloSVM3, main="SVM Kernel não Linear - Acurácia vs Valores de Sigma (Imputados)")
```

SVM Kernel não Linear – Acurácia vs Valores de Sigma (Imputados)

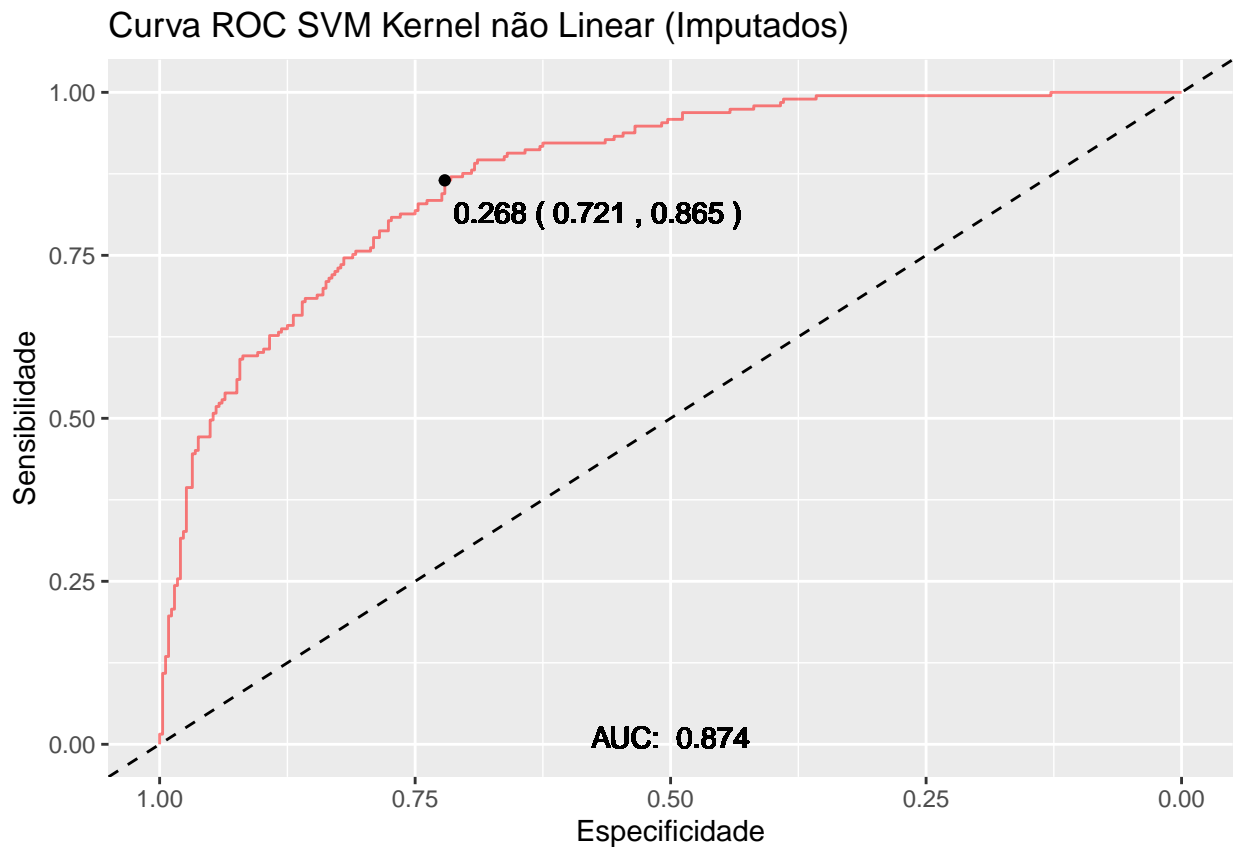


```
rocSVM3 <- roc(response = train$diabetes, predictor = predict(modeloSVM3, train, type = "prob")[,2])
```

```
## Setting levels: control = No, case = Yes
```

```
## Setting direction: controls < cases
```

```
plotaroc(rocSVM3, titulo = "Curva ROC SVM Kernel não Linear (Imputados)")
```



SVM Kernel Não Linear sem missing

```
## SVM Kernel Não Linear sem missing
set.seed(23)
modeloSVM4 <- caret::train(
  diabetes ~., data = train_without_NAs, method = "svmRadial",
  trControl = train.control,
  preProcess = c("center", "scale"),
  tuneGrid = expand.grid(sigma = seq(0.01, 1, length = 10) , C = seq(1, 10, length = 10))
)
print(modeloSVM4)
```

```
## Support Vector Machines with Radial Basis Function Kernel
##
## 266 samples
## 8 predictor
## 2 classes: 'No', 'Yes'
##
## Pre-processing: centered (8), scaled (8)
## Resampling: Cross-Validated (10 fold)
## Summary of sample sizes: 240, 240, 239, 239, 239, 239, ...
## Resampling results across tuning parameters:
##
```

##	sigma	C	Accuracy	Kappa
##	0.01	1	0.7596866	0.4650398
##	0.01	2	0.7670940	0.4812458
##	0.01	3	0.7750712	0.4961683
##	0.01	4	0.7635328	0.4698621
##	0.01	5	0.7786325	0.4972411
##	0.01	6	0.7784900	0.4946139
##	0.01	7	0.7746439	0.4843767
##	0.01	8	0.7709402	0.4757028
##	0.01	9	0.7710826	0.4782618
##	0.01	10	0.7710826	0.4782618
##	0.12	1	0.7787749	0.4951781
##	0.12	2	0.7712251	0.4802989
##	0.12	3	0.7635328	0.4566007
##	0.12	4	0.7598291	0.4445875
##	0.12	5	0.7524217	0.4302310
##	0.12	6	0.7562678	0.4371691
##	0.12	7	0.7522792	0.4200541
##	0.12	8	0.7562678	0.4370246
##	0.12	9	0.7373219	0.3911532
##	0.12	10	0.7376068	0.3892707
##	0.23	1	0.7598291	0.4523565
##	0.23	2	0.7561254	0.4451085
##	0.23	3	0.7337607	0.4021808
##	0.23	4	0.7413105	0.4199586
##	0.23	5	0.7225071	0.3736789
##	0.23	6	0.7223647	0.3670588
##	0.23	7	0.7111111	0.3417905
##	0.23	8	0.7111111	0.3408347
##	0.23	9	0.7109687	0.3405406
##	0.23	10	0.7148148	0.3367158
##	0.34	1	0.7522792	0.4378279
##	0.34	2	0.7336182	0.4009842
##	0.34	3	0.7339031	0.3950800
##	0.34	4	0.7225071	0.3770402
##	0.34	5	0.7111111	0.3487114
##	0.34	6	0.7183761	0.3638857
##	0.34	7	0.7109687	0.3366912
##	0.34	8	0.7109687	0.3441970
##	0.34	9	0.7185185	0.3508626
##	0.34	10	0.7186610	0.3590489
##	0.45	1	0.7450142	0.4217840
##	0.45	2	0.7450142	0.4322630
##	0.45	3	0.7222222	0.3712480
##	0.45	4	0.7072650	0.3469918
##	0.45	5	0.7188034	0.3656221
##	0.45	6	0.7149573	0.3580174
##	0.45	7	0.7186610	0.3583701
##	0.45	8	0.7186610	0.3619806
##	0.45	9	0.7112536	0.3476623
##	0.45	10	0.7075499	0.3454457
##	0.56	1	0.7448718	0.4243368
##	0.56	2	0.7262108	0.3809057
##	0.56	3	0.7109687	0.3386173

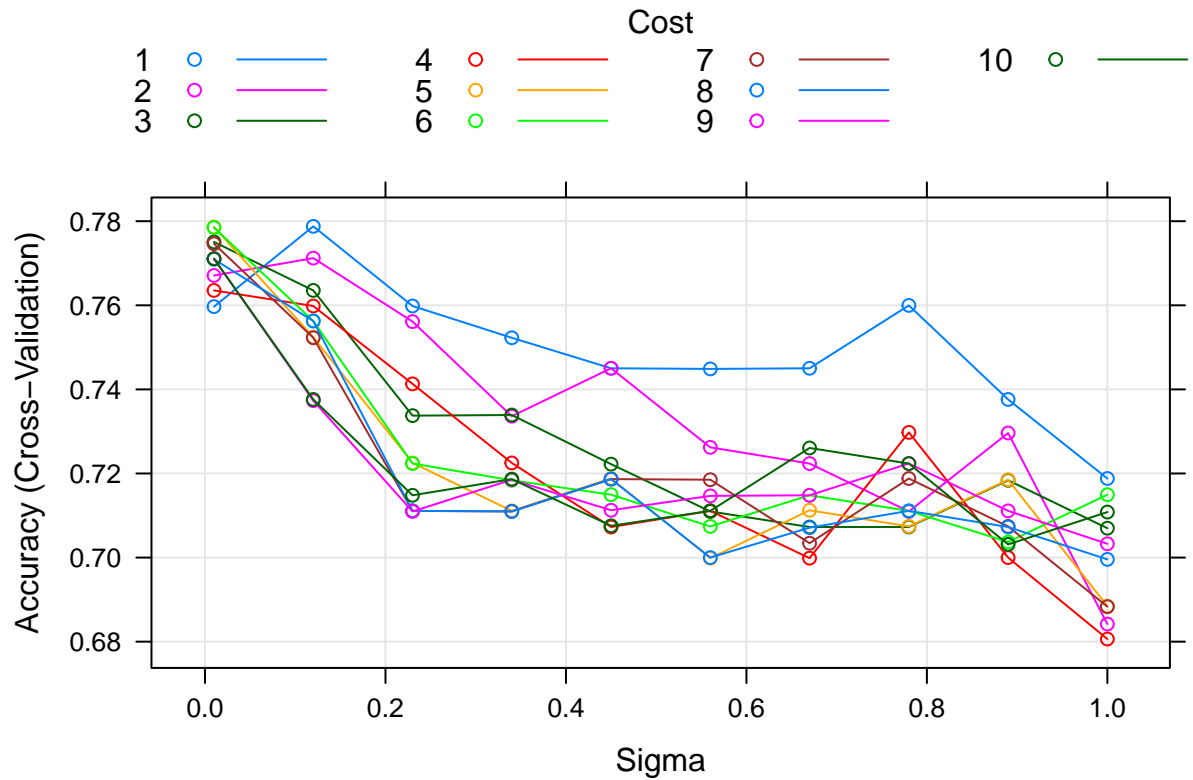
```

## 0.56 4 0.7111111 0.3454171
## 0.56 5 0.6998575 0.3195248
## 0.56 6 0.7074074 0.3369069
## 0.56 7 0.7185185 0.3779832
## 0.56 8 0.7000000 0.3153407
## 0.56 9 0.7146724 0.3515126
## 0.56 10 0.7111111 0.3485484
## 0.67 1 0.7450142 0.4361749
## 0.67 2 0.7223647 0.3844141
## 0.67 3 0.7072650 0.3418878
## 0.67 4 0.6998575 0.3169579
## 0.67 5 0.7112536 0.3447735
## 0.67 6 0.7148148 0.3577632
## 0.67 7 0.7034188 0.3278498
## 0.67 8 0.7071225 0.3375983
## 0.67 9 0.7148148 0.3570661
## 0.67 10 0.7260684 0.3767459
## 0.78 1 0.7599715 0.4719043
## 0.78 2 0.7109687 0.3479731
## 0.78 3 0.7072650 0.3401029
## 0.78 4 0.7297721 0.3925998
## 0.78 5 0.7074074 0.3478802
## 0.78 6 0.7111111 0.3430724
## 0.78 7 0.7188034 0.3574965
## 0.78 8 0.7111111 0.3475439
## 0.78 9 0.7223647 0.3713868
## 0.78 10 0.7223647 0.3729232
## 0.89 1 0.7376068 0.4239749
## 0.89 2 0.7296296 0.3890878
## 0.89 3 0.7183761 0.3677189
## 0.89 4 0.7000000 0.3174928
## 0.89 5 0.7185185 0.3622729
## 0.89 6 0.7037037 0.3369286
## 0.89 7 0.7074074 0.3383008
## 0.89 8 0.7072650 0.3383578
## 0.89 9 0.7111111 0.3472359
## 0.89 10 0.7031339 0.3065973
## 1.00 1 0.7188034 0.3915682
## 1.00 2 0.6841880 0.2662962
## 1.00 3 0.7069801 0.3218391
## 1.00 4 0.6806268 0.2520008
## 1.00 5 0.6884615 0.2735769
## 1.00 6 0.7149573 0.3506383
## 1.00 7 0.6883191 0.2773509
## 1.00 8 0.6995726 0.2933212
## 1.00 9 0.7032764 0.3099552
## 1.00 10 0.7108262 0.3449889
##
## Accuracy was used to select the optimal model using the largest value.
## The final values used for the model were sigma = 0.12 and C = 1.

```

```
plot(modeloSVM4, main="SVM Kernel não Linear - Acurácia vs Valores de Sigma (Removidos)")
```

SVM Kernel não Linear – Acurácia vs Valores de Sigma (Removidos)

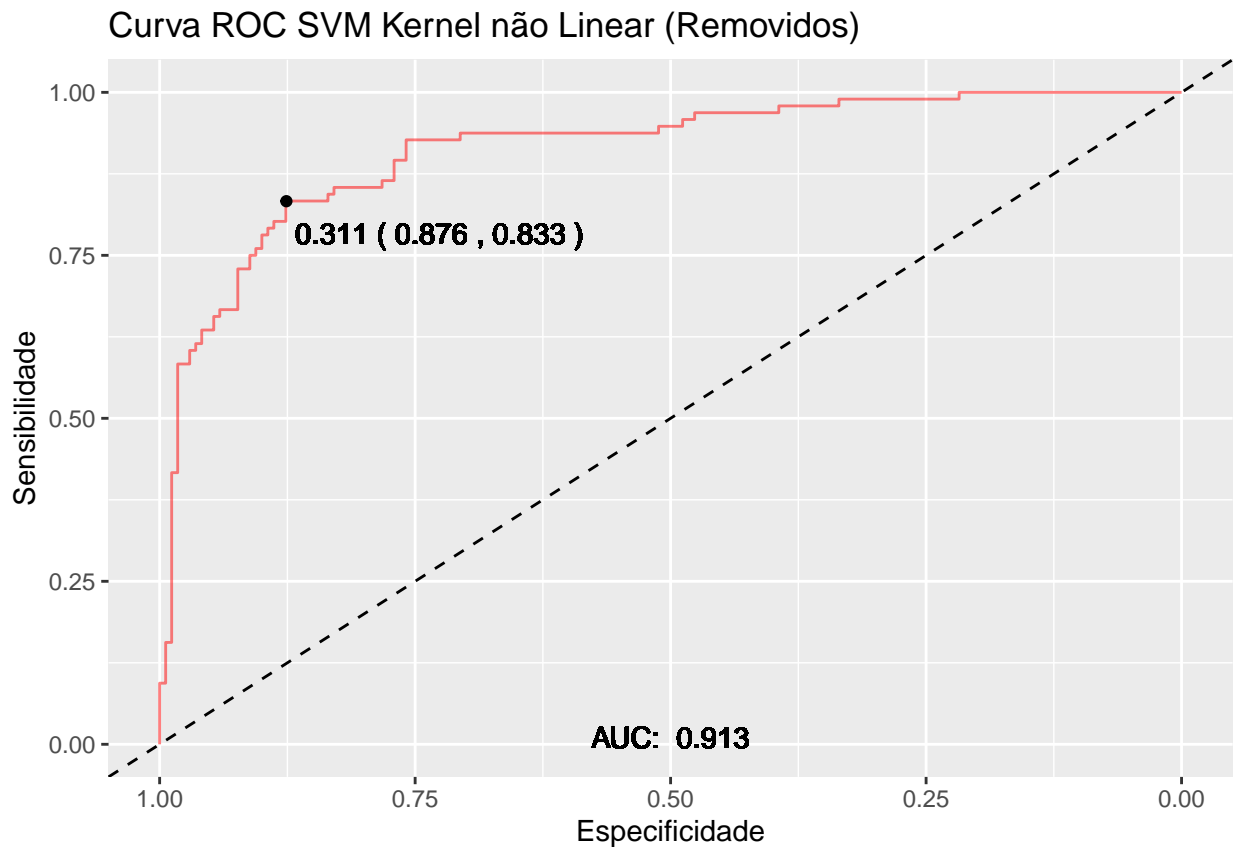


```
rocSVM4 <- roc(response = train_without_NAs$diabetes, predictor = predict(modeloSVM4, train_without_NAs
```

```
## Setting levels: control = No, case = Yes
```

```
## Setting direction: controls < cases
```

```
plotaroc(rocSVM4, titulo = "Curva ROC SVM Kernel não Linear (Removidos)")
```



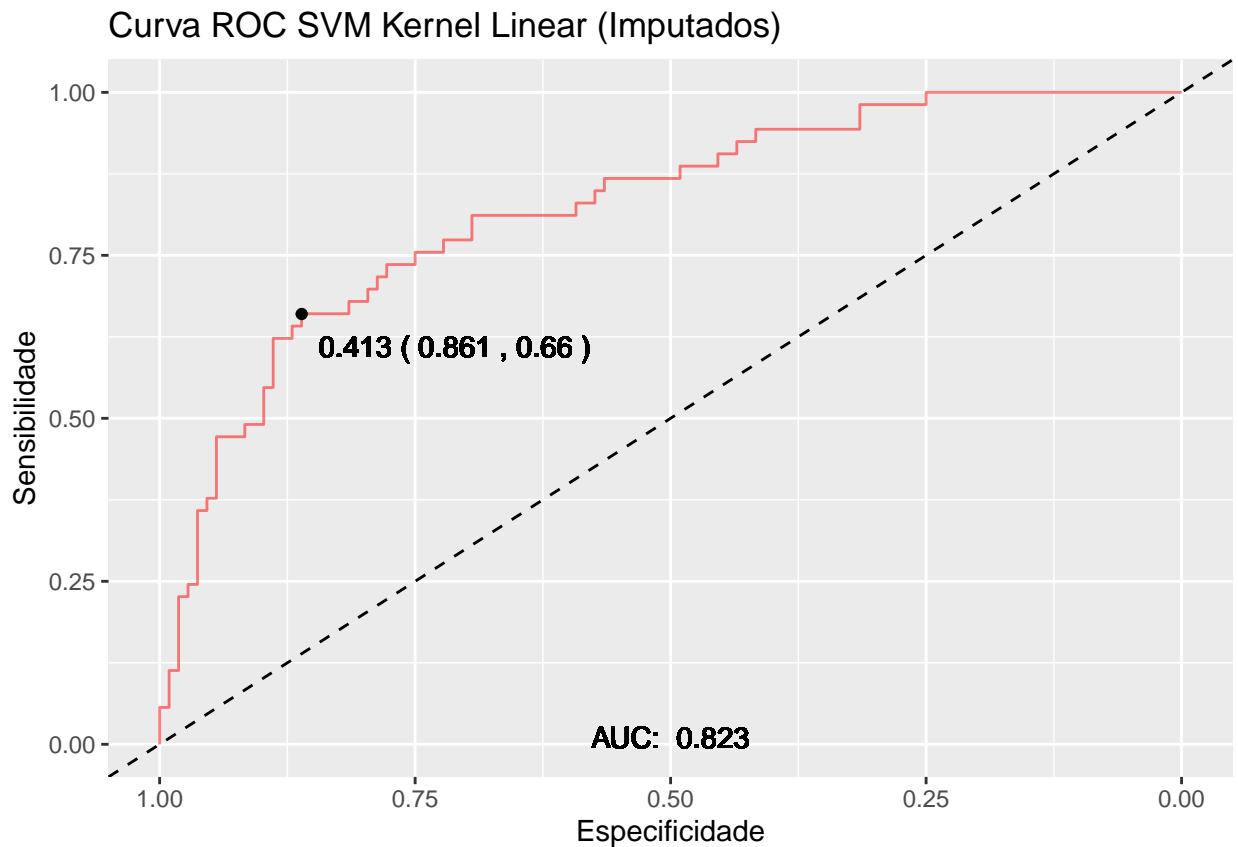
Testes

```
## SVM Kernel Linear com dados imputados
rocSVM1 <- roc(response = test$diabetes, predictor = predict(modeloSVM1, test, type = "prob")[,2])

## Setting levels: control = No, case = Yes

## Setting direction: controls < cases

plotaroc(rocSVM1, titulo = "Curva ROC SVM Kernel Linear (Imputados)")
```

```
predictSVM1 <- predict(modeloSVM1, newdata = test)
confusionMatrix(predictSVM1, test$diabetes)
```

```
## Confusion Matrix and Statistics
##
##           Reference
## Prediction No Yes
##      No  98  27
##      Yes  10  26
##
##           Accuracy : 0.7702
##           95% CI : (0.6974, 0.8327)
##      No Information Rate : 0.6708
##      P-Value [Acc > NIR] : 0.003815
##
##           Kappa : 0.4334
##
##  Mcnemar's Test P-Value : 0.008529
##
##           Sensitivity : 0.9074
##           Specificity : 0.4906
##      Pos Pred Value : 0.7840
##      Neg Pred Value : 0.7222
##           Prevalence : 0.6708
##      Detection Rate : 0.6087
```

```
## Detection Prevalence : 0.7764
## Balanced Accuracy : 0.6990
##
## 'Positive' Class : No
##
```

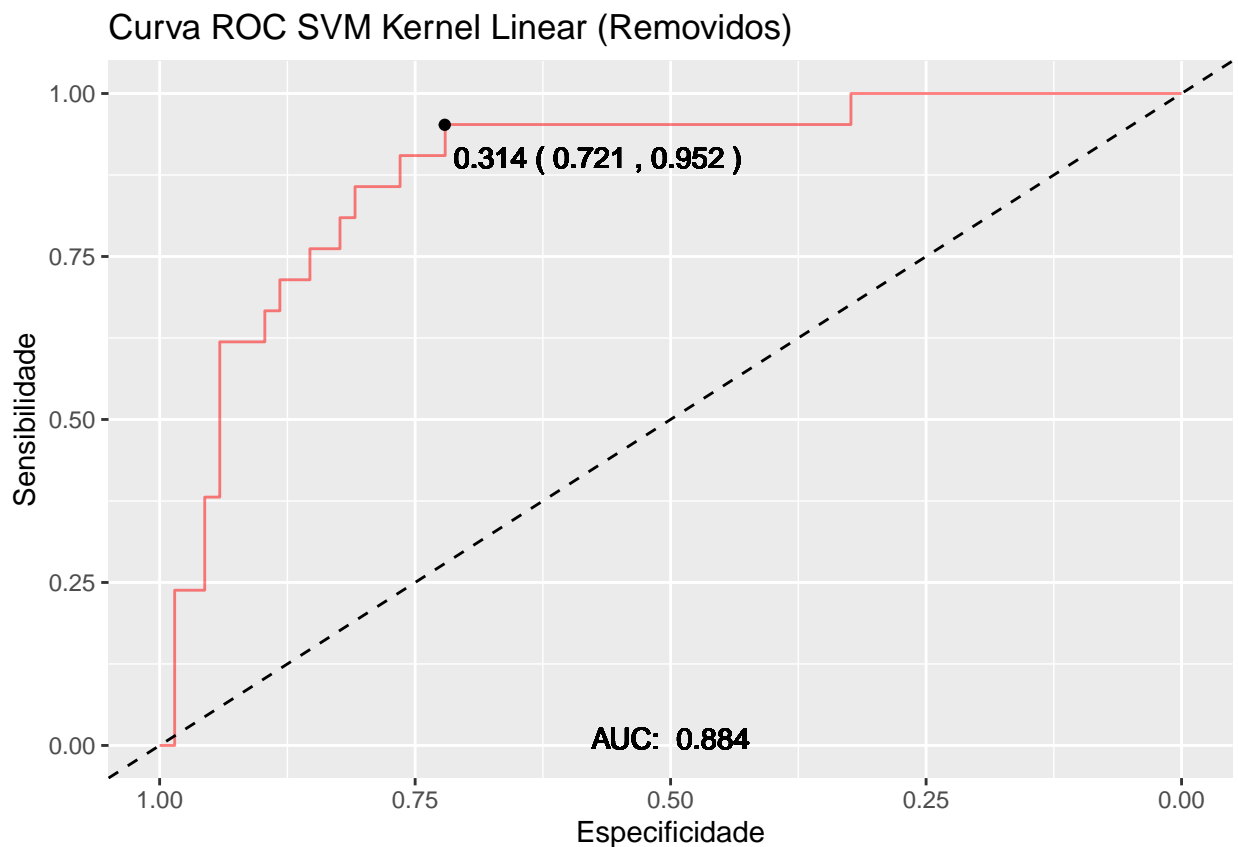
```
## SVM Kernel Linear sem missing
```

```
rocSVM2 <- roc(response = test_without_NAs$diabetes, predictor = predict(modeloSVM2, test_without_NAs, "
```

```
## Setting levels: control = No, case = Yes
```

```
## Setting direction: controls < cases
```

```
plotaroc(rocSVM2, titulo = "Curva ROC SVM Kernel Linear (Removidos)")
```



```
predictSVM2 <- predict(modeloSVM2, newdata = test_without_NAs)
```

```
confusionMatrix(predictSVM2, test_without_NAs$diabetes)
```

```
## Confusion Matrix and Statistics
```

```
##
```

```
## Reference
```

```
## Prediction No Yes
```

```
## No 64 12
```

```
## Yes 4 9
```

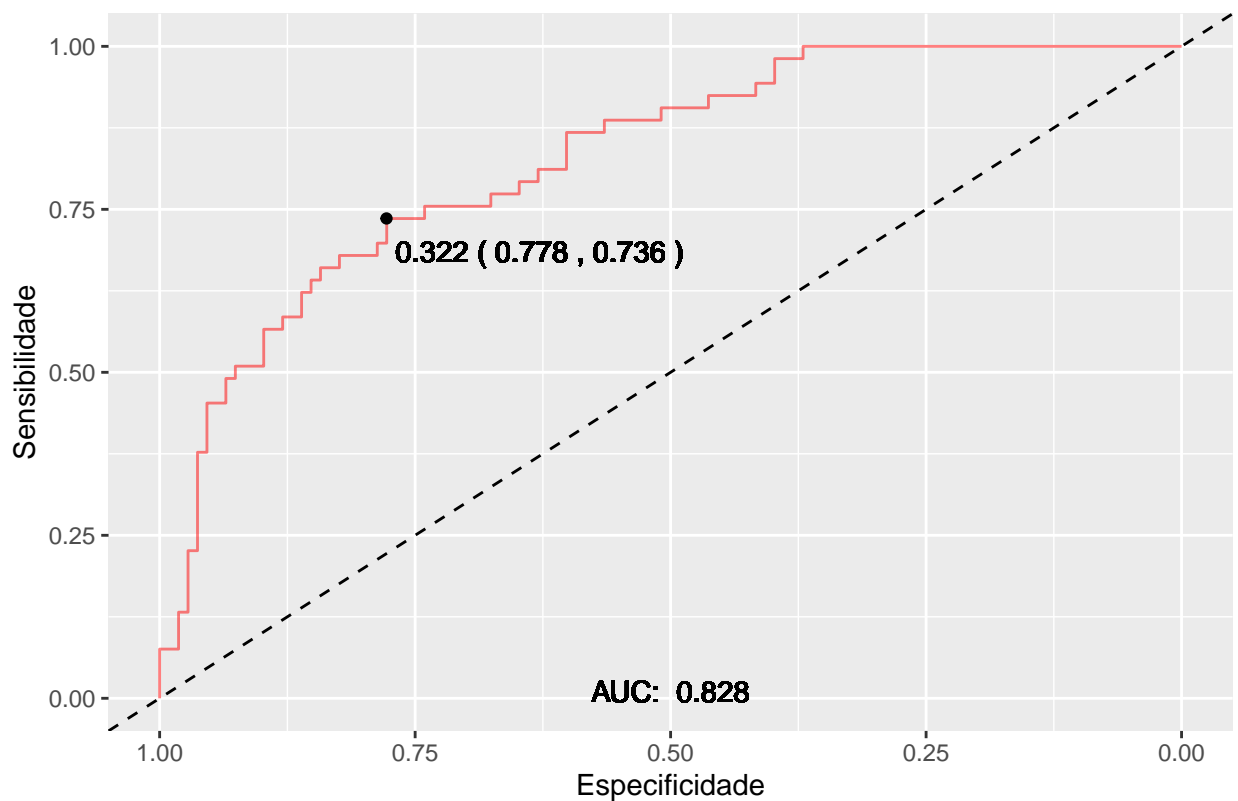
```
##
##           Accuracy : 0.8202
##           95% CI : (0.7245, 0.8936)
##      No Information Rate : 0.764
##      P-Value [Acc > NIR] : 0.12910
##
##           Kappa : 0.4258
##
##      McNemar's Test P-Value : 0.08012
##
##           Sensitivity : 0.9412
##           Specificity : 0.4286
##      Pos Pred Value : 0.8421
##      Neg Pred Value : 0.6923
##           Prevalence : 0.7640
##      Detection Rate : 0.7191
##      Detection Prevalence : 0.8539
##      Balanced Accuracy : 0.6849
##
##      'Positive' Class : No
##
```

```
## SVM Kernel não Linear com dados imputados
rocSVM3 <- roc(response = test$diabetes, predictor = predict(modeloSVM3, test, type = "prob")[,2])
```

```
## Setting levels: control = No, case = Yes
## Setting direction: controls < cases
```

```
plotaroc(rocSVM3, titulo = "Curva ROC SVM Kernel não Linear (Imputados)")
```

Curva ROC SVM Kernel não Linear (Imputados)



```
predictSVM3 <- predict(modeloSVM3, newdata = test)
confusionMatrix(predictSVM3, test$diabetes)
```

```
## Confusion Matrix and Statistics
```

```
##
```

```
##           Reference
```

```
## Prediction No Yes
```

```
##           No  99  26
```

```
##           Yes   9  27
```

```
##
```

```
##           Accuracy : 0.7826
```

```
##           95% CI : (0.7109, 0.8437)
```

```
## No Information Rate : 0.6708
```

```
## P-Value [Acc > NIR] : 0.001230
```

```
##
```

```
##           Kappa : 0.464
```

```
##
```

```
## McNemar's Test P-Value : 0.006841
```

```
##
```

```
##           Sensitivity : 0.9167
```

```
##           Specificity : 0.5094
```

```
## Pos Pred Value : 0.7920
```

```
## Neg Pred Value : 0.7500
```

```
## Prevalence : 0.6708
```

```
## Detection Rate : 0.6149
```

```
## Detection Prevalence : 0.7764
## Balanced Accuracy : 0.7131
##
## 'Positive' Class : No
##
```

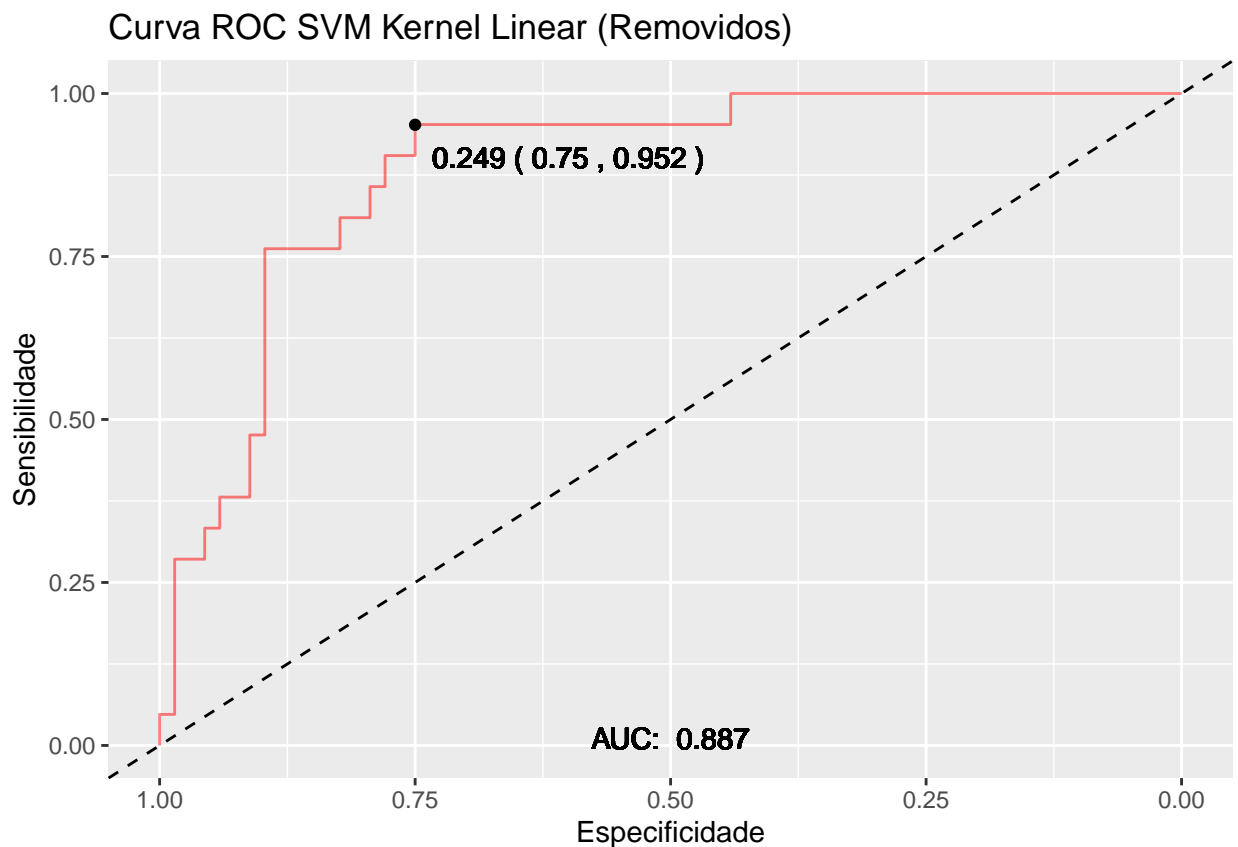
```
## SVM Kernel Linear sem missing
```

```
rocSVM4 <- roc(response = test_without_NAs$diabetes, predictor = predict(modeloSVM4, test_without_NAs, "
```

```
## Setting levels: control = No, case = Yes
```

```
## Setting direction: controls < cases
```

```
plotaroc(rocSVM4, titulo = "Curva ROC SVM Kernel Linear (Removidos)")
```



```
predictSVM4 <- predict(modeloSVM4, newdata = test_without_NAs)
confusionMatrix(predictSVM4, test_without_NAs$diabetes)
```

```
## Confusion Matrix and Statistics
```

```
##
```

```
## Reference
```

```
## Prediction No Yes
```

```
## No 63 13
```

```
## Yes 5 8
```

```
##
##           Accuracy : 0.7978
##           95% CI   : (0.6993, 0.8755)
##    No Information Rate : 0.764
##    P-Value [Acc > NIR] : 0.27088
##
##           Kappa : 0.354
##
##    McNemar's Test P-Value : 0.09896
##
##           Sensitivity : 0.9265
##           Specificity : 0.3810
##           Pos Pred Value : 0.8289
##           Neg Pred Value : 0.6154
##           Prevalence : 0.7640
##           Detection Rate : 0.7079
##           Detection Prevalence : 0.8539
##           Balanced Accuracy : 0.6537
##
##           'Positive' Class : No
##
```

Validação

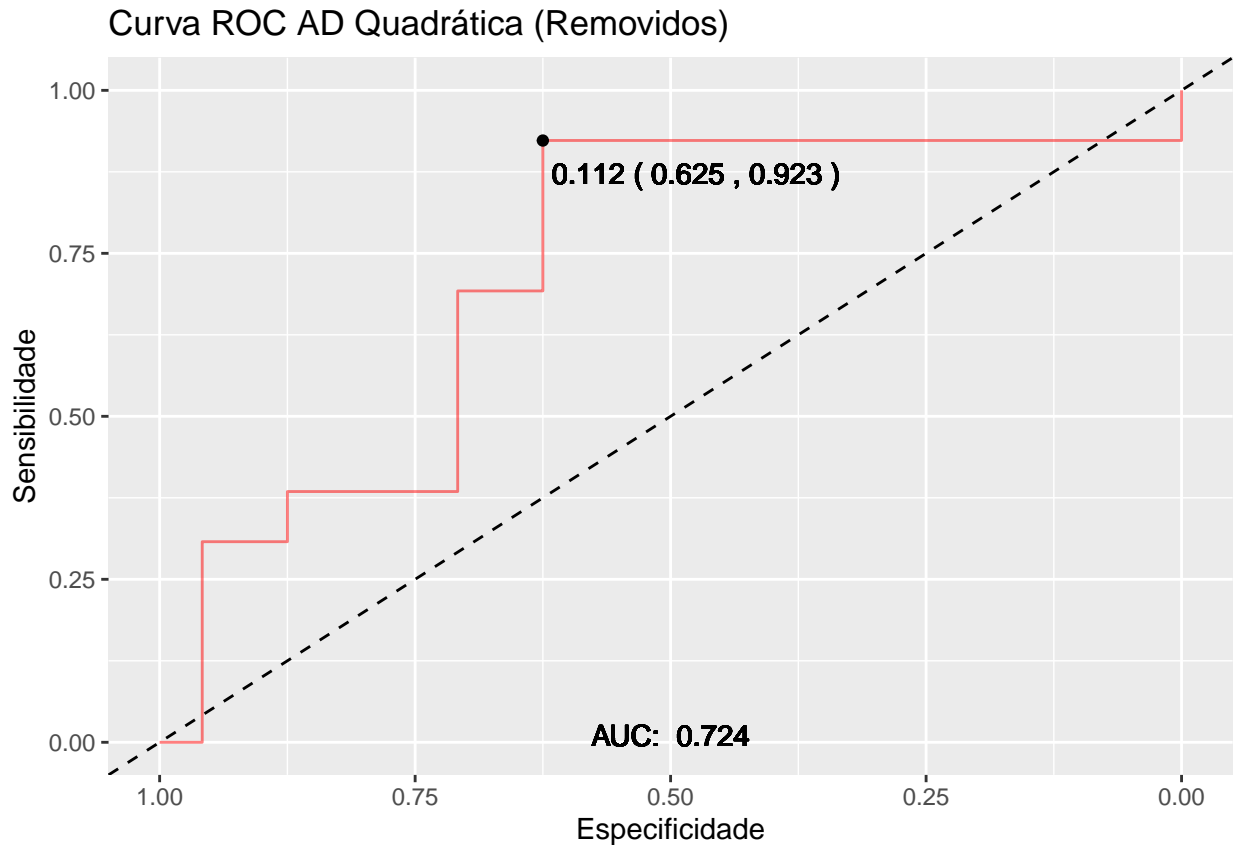
AD Quadrática - Dados Removidos

```
rocAD6out <- roc(response = test_out_sample_whitout_NAs$diabetes, predictor = predict(modeloAD6, test_o

## Setting levels: control = No, case = Yes

## Setting direction: controls < cases

plotaroc(rocAD6out, titulo = "Curva ROC AD Quadrática (Removidos)")
```



```
predictAD6out <- predict(modeloAD6, newdata = test_out_sample_whitout_NAs)
confusionMatrix(predictAD6out, test_out_sample_whitout_NAs$diabetes)
```

```
## Confusion Matrix and Statistics
##
##           Reference
## Prediction No Yes
##      No  17   5
##      Yes   7   8
##
##           Accuracy : 0.6757
##           95% CI : (0.5021, 0.8199)
##      No Information Rate : 0.6486
##      P-Value [Acc > NIR] : 0.4384
##
##           Kappa : 0.3127
##
##  Mcnemar's Test P-Value : 0.7728
##
##           Sensitivity : 0.7083
##           Specificity : 0.6154
##      Pos Pred Value : 0.7727
##      Neg Pred Value : 0.5333
##           Prevalence : 0.6486
##      Detection Rate : 0.4595
```

```
## Detection Prevalence : 0.5946
## Balanced Accuracy : 0.6619
##
## 'Positive' Class : No
##
```

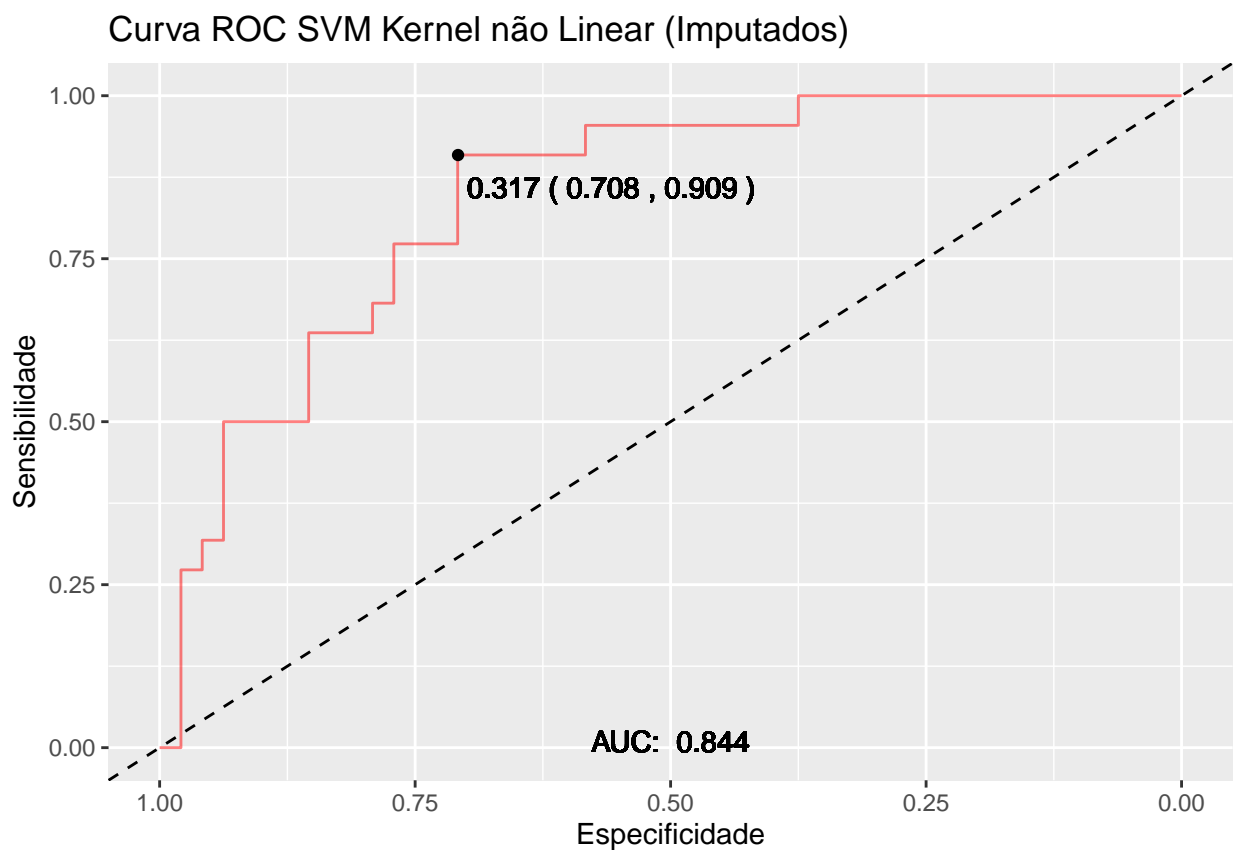
SVM Kernel Não Linear - Dados Imputados

```
## SVM Kernel não Linear com dados imputados
rocSVM3 <- roc(response = test_out_sample$diabetes, predictor = predict(modeloSVM3, test_out_sample, type = "raw"))

## Setting levels: control = No, case = Yes

## Setting direction: controls < cases

plotaroc(rocSVM3, titulo = "Curva ROC SVM Kernel não Linear (Imputados)")
```



```
predictSVM3 <- predict(modeloSVM3, newdata = test_out_sample)
confusionMatrix(predictSVM3, test_out_sample$diabetes)
```

```
## Confusion Matrix and Statistics
```



```

##
##           Reference
## Prediction No Yes
##           No  38   8
##           Yes 10  14
##
##           Accuracy : 0.7429
##           95% CI : (0.6244, 0.8399)
##           No Information Rate : 0.6857
##           P-Value [Acc > NIR] : 0.1846
##
##           Kappa : 0.4177
##
## Mcnemar's Test P-Value : 0.8137
##
##           Sensitivity : 0.7917
##           Specificity : 0.6364
##           Pos Pred Value : 0.8261
##           Neg Pred Value : 0.5833
##           Prevalence : 0.6857
##           Detection Rate : 0.5429
##           Detection Prevalence : 0.6571
##           Balanced Accuracy : 0.7140
##
##           'Positive' Class : No
##

```