Mini Project Report On

"Diabetes Prediction Using Data Mining"

BACHELOR OF COMPUTER ENGINEERING

SUBMITTED BY

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UNDER THE GUIDANCE OF

Prof. S.B.Shirke



DEPARTMENT OF COMPUTER ENGINEERING

SAVITRIBAI PHULE PUNE UNIVERSITY

Batch of 2021-22



CERTIFICATE

This is to certify that the seminar report entitled

"Diabetes Prediction Using Data Mining"

Submitted by

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Are bonafide student of this institute and the work has been carried out by her under the supervision of **Prof.S.B.Shirke** and it is approved for the partial fulfilment of the requirement of Savitribai Phule Pune University Computer Engineering.

Prof.S.B.Shirke Prof.B.D.Thorat **Guide** HOD

Dr.S.B.Patil **Principal**

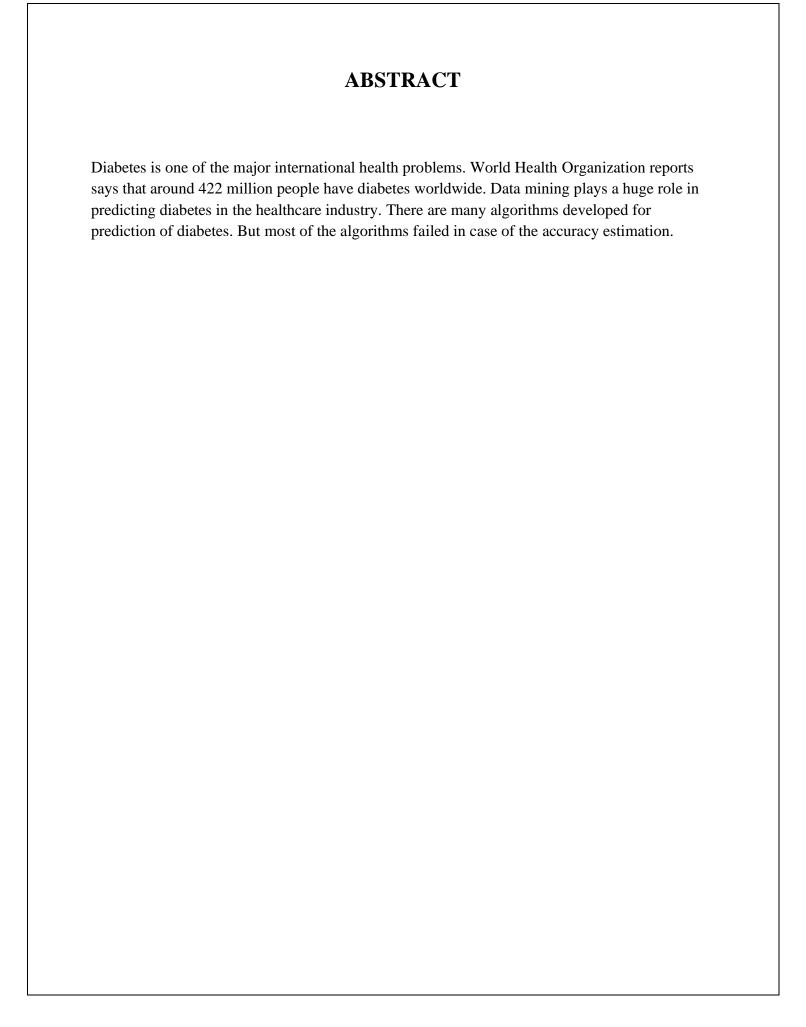
Shri Chhatrapati Shivajiraje college of Engineering Pune

Place: Dhangwadi Date:

ACKNOWLEDGEMENT

We extend our sincere and heartfelt thanks to our esteemed guide, **Prof.S.B.Shirke** for this exemplary guidance, monitoring and constant encouragement throughout the course at crucial junctures and for showing us the write way.

We would like to extends thanks to our respected HEAD of the division **Prof. B.D.Thorat** for following us to use to the facilities available. We would like to thanks faculty members also Last but not the least, We would like to thanks our friends and family for the support and encouragement they have give us during the course of our work



Introduction

Objective:

•To predict diabetes in healthcare industry using data mining.

Project Overview:

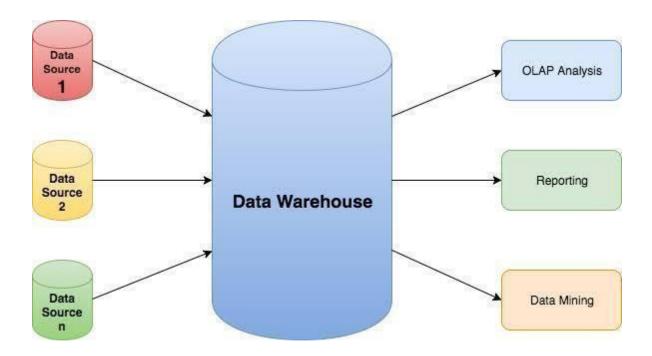
Diabetes is one of the major international health problems. World Health Organization reports says that around 422 million people have diabetes worldwide. Data mining plays a huge role in predicting diabetes in the healthcare industry. There are many algorithms developed for prediction of diabetes. But most of the algorithms failed in case of the accuracy estimation. Also, there is a need to automate the overall process of diabetes prediction. This automation of diabetic database helps in identification of impact of diabetes on various human organs. More the accuracy of prediction, more the chances of accurate severity estimation. Therefore this project concentrated on providing different prediction methods of diabetes.

Propose system

Dataset:

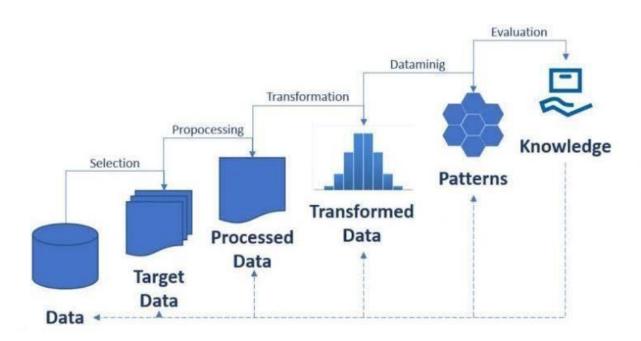
Here PIMA Indian diabetes data set is considered. The data set is taken from UCI machine learning repository. The data set consists of 9 attributes: number of times pregnant, plasma glucose concentration, diastolic blood pressure, triceps skin folds thickness, serum insulin, body mass index, pedigree type, age, and class. Here, the class label is binary classification. It has two values

- Tested positive (1) which means diabetic
- Tested negative (0) which says nondiabetic



Methodology

Data preprocessing and data mining algorithms are used for the further process in the project. Data preprocessing technique data transformation is applied to the data set before applying data mining algorithms. The decision tree and regression models are built. Decision trees and Regression models are used to predict the final binary target variable. After running different types of models, model comparison needed to select the best algorithm. The best algorithm and best model is selected based on the high accuracy rate.



HARDWARE AND SOFTWARE REQUIREMENTS

Software Requirements:

Windows OS

Weka

Hardware Requirements:

Hard Disk − 1 TB or Above

RAM required – 8 GB or Above

Processor – Core i3 or Above

Technology Used:

Data Mining

Data Visualization

Coding:

```
opetion(repos = c(CRAN = "http://cran.rstudio.com"))
library(RSQLite)
library(DBI)
library(datasets)
library(caTools)
library(e1071)
f<-file.choose("diab1.csv")
mydata<-read.csv(f)
datasetss<-read.csv(f)
#mydata<-read.csv(file = "/home/spllab01/Diabetes/diab1.csv",header = TRUE,sep = ",")</pre>
#datasetss<-read.csv(file = "/home/spllab01/Diabetes/diab1.csv",header = TRUE,sep = ",")
View(mydata)
View(datasetss)
#dataset<-read.table('diab.csv',header = T)</pre>
datasetss$SkinThickness = ifelse(is.na(datasetss$SkinThickness),
ave(datasetss$SkinThickness, FUN = function(x) mean(x, na.rm = 'TRUE')),
datasetss$SkinThickness)
```

#Fills the NUll values with the average of that column values.

View(datasetss)

 $datasetss\$Glucose = ifelse(is.na(datasetss\$Glucose), ave(datasetss\$Glucose, FUN = function(x) \\ mean(x, na.rm = 'TRUE')), datasetss\$Glucose)$

View(datasetss)

h<-hist(datasetssSkinThickness,main="SkinThickness frquencies - histogram", xlab = "SkinThickness", xlim = c(5,50),col = "blue")

h<-hist(datasetss\$Glucose,main="Glucose frquencies - histogram", xlab = "Glucose Value",col = "blue",labels = TRUE, breaks = 8, border = "green",las=3)

train<-as.data.frame(datasetss[1:200,]) View(train)

test<-as.data.frame(datasetss[201:299,]) View(test)

head(train)

#Pregnancies Glucose BloodPressure SkinThickness Insulin BMI DiabetesPedigreeFunction Age

#201	0	113	80	16	0 31.0	0.874
#202	1	138	82	0	0 40.1	21 0.236
#203	0	108	68	20	0 27.3	28 0.787
11203	O	100	00	20	0 21.3	32

#204	2	99	70	16	44 20.4	0.235					
#205	6	103	72	32	190 37.7	27 0.324 55					
#206	5	111	72	28	0 23.9	0.407 27					
						_,					
#Outcome											
#201 0											
#202 0 #203 0)										
#204 0 #205 0											
#206 0	1										
my_model<-naiveBayes(as.factor(train\$Outcome)~.,train) pred1<-											
predict(my_model,test[,-9])											
pred1											
#[1] 0 0	001011	01011	011000	001101	0000100	0.1.0.0.0					
#[1]00001011010110110000110100001001000 11110000111											
$\#[47]\ 1\ 1\ 1\ 0\ 0\ 0\ 0\ 1\ 0\ 0\ 1\ 1\ 1\ 0\ 0\ 1\ 0\ 0\ 1\ 0\ 0\ 0\ 1\ 0\ 0\ 0\ 0\ 0$											
#[93] 1 1 1 1 1 0 1											
#Levels: 0 1											
#generate the confusion matrix											

table(pred1,test\$Outcome,dnn=c("predicted","actual"))

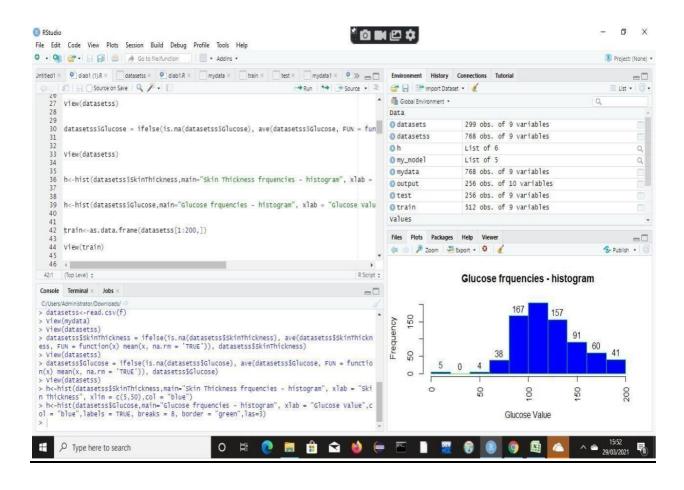
```
#actual
 #predicted 0 1
 #0 41 16
 #1 19 23
 #Build Classifier Models using Different Techniques.
 #Cross Validation.
 #Cross Validation K fold cross validation
 library(caret) library(lattice)
 library(ggplot2)
 # Define train control for k fold cross validation train_control
 <- trainControl(method="cv", number=10)
# Fit Naive Bayes Model model <- train(SkinThickness~., data=datasetss,
trControl=train control, method="knn")
# Summarise Results print(model)
#k-Nearest Neighbors
#299 samples
#8 predictor
#No pre-processing
#Resampling: Cross-Validated (10 fold)
#Summary of sample sizes: 269, 269, 269, 268, 270, 270, ...
```

```
#Resampling results across tuning parameters:
# k RMSE Rsquared MAE
#5 11.87092 0.3645257 9.515780
#7 12.01004 0.3420931 9.699612
#9 11.83102 0.3558637 9.702612
#RMSE was used to select the optimal model using the smallest value.
#The final value used for the model was k = 9.
#scatterPolt matrix
head(datasetss)
pairs(datasetss[,1:4], pch = 19)
pairs(datasetss[,5:9], pch = 19)
pairs(datasetss[,1:4], pch = 19, lower.panel = NULL)
pairs(datasetss[,5:9], pch = 19, lower.panel = NULL)
View(datasetss)
```

```
#One more classifier model
library(mlbench)
library(caret)
# prepare training scheme control <-
trainControl(method="repeatedcv", number=10, repeats=3)
# CART
set.seed(7) fit.cart <- train(SkinThickness~., data=datasetss,
method="rpart", trControl=control)
# SVM
set.seed(7) fit.svm <- train(SkinThickness~., data=datasetss,
method="svmRadial", trControl=control)
# kNN
set.seed(7)
fit.knn <- train(SkinThickness~., data=datasetss, method="knn", trControl=control)
# Random Forest set.seed(7) fit.rf <- train(SkinThickness~., data=datasetss, method="rf",
trControl=control)
# collect resamples
results <- resamples(list(CART=fit.cart, SVM=fit.svm, KNN=fit.knn, RF=fit.rf))
summary(results)
```

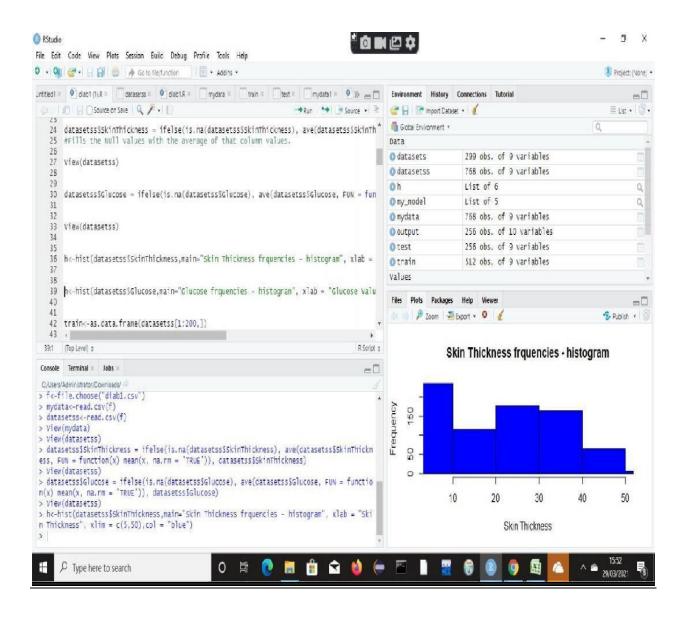
```
#box wisker scales <- list(x=list(relation="free"),
y=list(relation="free")) bwplot(results, scales=scales)
#density plots scales <- list(x=list(relation="free"),
y=list(relation="free")) densityplot(results, scales=scales,
pch = "|")
#dot plots scales <- list(x=list(relation="free"),
y=list(relation="free")) dotplot(results, scales=scales)
#parallel plots parallelplot(results)
#scatter plot splom(results)
#pair wise x and y plots
xyplot(results, models=c("KNN", "SVM"))
#statisticall significance test #
difference in model predictions
diffs <- diff(results)
# summarize p-values for pair-wise comparisons summary(diffs)
```

Glucose Frequencies-histogram:

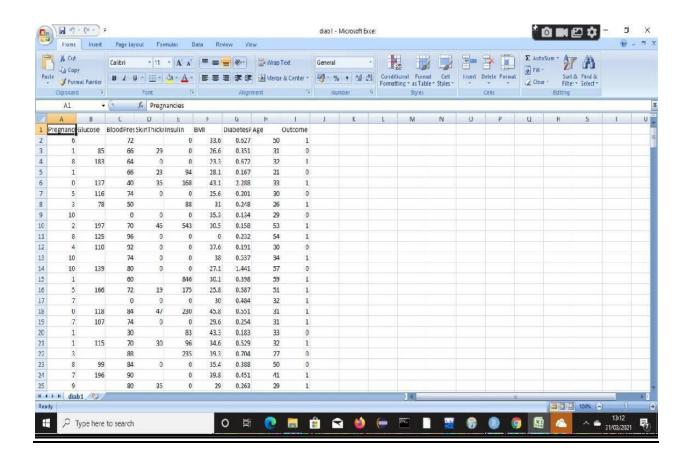


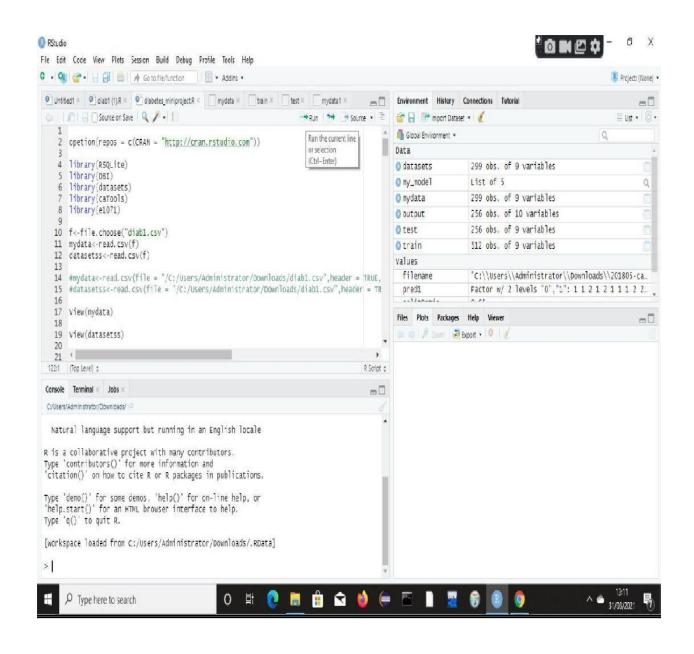
Outputs:

Skin Thickness Frequencies:



Database:





Conclusion: Finally, decision is built using c4.5 decision tree algorithm. All the results are displayed to the end user using weka data visualization, regression provides the predicted outcome to the end user.