# **Final Project**

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### **Data Loading**

train\_genetic = read.csv(file="/Users/sakshyamdahal/Desktop/MS\_Data\_Science/Applied P
redictive Modeling/Final project/train\_genetic\_disorders.csv", header= TRUE)

```
df <- train_genetic
#head(df,5)
dim(df)</pre>
```

```
> [1] 22083 45
```

#### Initial feature reduction

Uninformative Feature Reduced

```
#drop unecessary columns
drop_col <- c("Patient.Id", "Patient.First.Name", "Father.s.name", "Institute.Name", "Lo
cation.of.Institute" , "Family.Name", "Parental.consent", "Status", "Place.of.birth",
"Test.1", "Test.2", "Test.3", "Test.4", "Test.5")
df <- df[!(names(df) %in% drop_col)]
#names(df)
dim(df)</pre>
```

```
> [1] 22083 31
```

## Missing value in target vector.

```
#unique(df$Disorder.Subclass)
miss_value_target <- which(is.na(df$Disorder.Subclass) | (df$Disorder.Subclass == "")
)
length(miss_value_target)</pre>
```

```
> [1] 3140
```

#### Excuding data points where our target has missing value.

```
df <- df[-miss_value_target,]
dim(df)</pre>
```

```
> [1] 18943 31
```

#### Check for class imbalance

```
prop.table(table(df$Disorder.Subclass))
```

```
Alzheimer's
>
                                                                        Cancer
                            0.007812912
                                                                  0.004856675
                       Cystic fibrosis
                                                                     Diabetes
>
>
                            0.172992662
                                                                  0.092012881
                       Hemochromatosis Leber's hereditary optic neuropathy
>
                            0.068151824
                                                                  0.032043499
>
>
                        Leigh syndrome
                                                      Mitochondrial myopathy
                            0.258301219
                                                                  0.221823365
>
>
                              Tay-Sachs
                            0.142004962
>
```

### Split the dataframe as per the disorder subclass vector

There is certain class imbalance problem. We will split the data frame using createDataPartition function using target class(Disorder.Subclass) which helps us to get even balance of all classes in both train and test data

```
set.seed(100)
library(caret)
train_row <- createDataPartition(df$Disorder.Subclass, p = .80, list = FALSE)

train_df <- df[train_row,]
test_df <- df[-train_row,]

dim(df)</pre>
```

```
> [1] 18943 31
```

```
dim(train_df)
> [1] 15158
                31
dim(test_df)
> [1] 3785
              31
prop.table(table(train df$Disorder.Subclass))
>
>
                           Alzheimer's
                                                                       Cancer
                           0.007850640
                                                                 0.004881911
                       Cystic fibrosis
>
                                                                     Diabetes
                           0.172977965
                                                                 0.092030611
>
                       Hemochromatosis Leber's hereditary optic neuropathy
>
                           0.068148832
                                                                 0.032062277
>
                        Leigh syndrome
                                                      Mitochondrial myopathy
                           0.258279456
                                                                 0.221797071
>
>
                             Tay-Sachs
                           0.141971236
>
```

```
#lapply(train_df, unique)
```

## Checking and handling missing values for train data

<sup>\*\*</sup>We can see here no difference in the proportion of classes after split

```
#changing ambiguous values from the data to na.
library(caret)
library(dplyr)
library(questionr)
train_df[train_df == ""] <- NA</pre>
test df[test df == ""] <- NA
train_df[train_df == "-"] <- NA</pre>
test_df[test_df == "-"] <- NA
#not chaninging not applicable to na as I think it itself is a class
#df[df == "Not applicable"] <- NA
train_df[ train_df == "Not available"] <- NA</pre>
test_df[ test_df == "Not available"] <- NA</pre>
train_df[train_df == "No record"] <- NA</pre>
test_df[test_df == "No record"] <- NA</pre>
train df[train df == "None"] <- "No"</pre>
test df[test df == "None"] <- "No"</pre>
freq.na(train_df)
```

>	missing	8
> Birth.asphyxia	8297	55
> H.O.substance.abuse	4925	32
> H.O.radiation.exposurex.ray.	4775	32
> Mother.s.age	3962	26
> Father.s.age	3898	26
> Maternal.gene	1824	12
> Symptom.2	1458	10
> Respiratory.Ratebreaths.min.	1450	10
> Genetic.Disorder	1437	9
> Symptom.5	1434	9
> Symptom.1	1429	9
> Follow.up	1428	9
> Birth.defects	1427	9
> White.Blood.cell.countthousand.per.microlit	er. 1422	9
> Gender	1407	9
> History.of.anomalies.in.previous.pregnancies	1389	9
> H.O.serious.maternal.illness	1388	9
> Blood.test.result	1388	9
> Symptom.4	1382	9
> Heart.Raterates.min	1380	9
> Noof.previous.abortion	1378	9
> Folic.acid.detailsperi.conceptional.	1376	9
> Symptom.3	1376	9
> Assisted.conception.IVF.ART	1373	9
> Patient.Age	971	6
> Autopsy.shows.birth.defectif.applicable.	674	4
> Inherited.from.father	185	1
> Genes.in.mother.s.side	0	0
> Paternal.gene	0	0
> Blood.cell.countmcL.	0	0
> Disorder.Subclass	0	0

```
sum(is.na(train_df))
```

```
> [1] 54833
```

## Droping the columns with high missing value

The imputaion in more than 50% of missing data might hamper our model.

```
#drop Birth.asphyxia column as 57% of data missing and imputation might create inac
curate model prediction
#dim(train_df)
train_df <- subset(train_df, select = -c(Birth.asphyxia))
dim(train_df)</pre>
```

```
> [1] 15158 30
```

```
test_df <- subset(test_df, select = -c(Birth.asphyxia))
dim(test_df)</pre>
```

```
> [1] 3785 30
```

```
str(train df)
```

```
> 'data.frame': 15158 obs. of 30 variables:
 $ Patient.Age
                                                    : int 2 4 6 12 11 14 3 11 4 6 .
  $ Genes.in.mother.s.side
                                                    : chr "Yes" "Yes" "Yes" "Yes" .
  $ Inherited.from.father
                                                           "No" "Yes" "No" "No" ...
                                                    : chr
                                                    : chr "Yes" "No" "No" "Yes" ...
 $ Maternal.gene
 $ Paternal.gene
                                                    : chr
                                                           "No" "No" "No" "No" ...
> $ Blood.cell.count..mcL.
                                                    : num 4.76 4.91 4.89 4.71 4.72
> $ Mother.s.age
                                                    : int NA NA 41 21 32 NA NA 45 4
4 NA ...
> $ Father.s.age
                                                    : int NA 23 22 NA NA NA 63 44 4
2 NA ...
> $ Respiratory.Rate..breaths.min.
                                                    : chr "Normal (30-60)" "Tachypn
ea" "Normal (30-60)" "Tachypnea" ...
> $ Heart.Rate..rates.min
                                                    : chr "Normal" "Tachyc
ardia" "Normal" ...
                                                          "High" "High" "Low" "High
> $ Follow.up
                                                    : chr
 . . .
  $ Gender
                                                    : chr NA NA NA "Male" ...
  $ Autopsy.shows.birth.defect..if.applicable.
                                                           "Not applicable" "No" "No
                                                    : chr
t applicable" "No" ...
                                                    : chr "No" "Yes" "Yes" "No" ...
  $ Folic.acid.details..peri.conceptional.
                                                    : chr NA "Yes" "No" "Yes" ...
  $ H.O.serious.maternal.illness
  $ H.O.radiation.exposure..x.ray.
                                                           "No" "Not applicable" "Ye
                                                    : chr
s" NA ...
  $ H.O.substance.abuse
                                                    : chr
                                                           "No" "Not applicable" NA
```

```
"Not applicable" ...
> $ Assisted.conception.IVF.ART
                                                    : chr "No" "No" "Yes" NA ...
  $ History.of.anomalies.in.previous.pregnancies
                                                            "Yes" "Yes" "Yes" "Yes" .
                                                    : chr
> $ No..of.previous.abortion
                                                    : int NA NA 4 1 4 0 3 0 1 1 ...
  $ Birth.defects
                                                     : chr NA "Multiple" "Singular"
"Singular" ...
 $ White.Blood.cell.count..thousand.per.microliter.: num 9.86 5.52 NA 7.92 4.1 ...
  $ Blood.test.result
                                                     : chr NA "normal" "normal" "inc
onclusive" ...
                                                     : int 1 1 0 0 0 1 0 1 0 1 ...
> $ Symptom.1
  $ Symptom.2
                                                     : int 1 NA 1 0 0 0 0 1 0 NA ...
                                                     : int 1 1 1 1 0 0 0 1 1 0 ...
 $ Symptom.3
                                                     : int 1 1 1 0 0 1 0 0 1 0 ...
  $ Symptom.4
> $ Symptom.5
                                                     : int 1 0 1 0 NA 0 0 1 1 NA ...
> $ Genetic.Disorder
                                                     : chr "Mitochondrial genetic in
heritance disorders" NA "Multifactorial genetic inheritance disorders" "Mitochondrial
genetic inheritance disorders" ...
> $ Disorder.Subclass
                                                     : chr "Leber's hereditary optic
neuropathy" "Cystic fibrosis" "Diabetes" "Leigh syndrome" ...
```

### Identifying the dtypes and changing into suitable dtypes.

- 1. Numeric vectors are mostly acceptable by all models. There are 11 numeric vectors in the data.
- 2. Vectors in string/character needs to be converted into factor for running smooth model. 19 character vectors are converted into factors.

```
#subsetting numeric columns and character for handling missing values and
library(dplyr)
# Subset numeric columns with dplyr
train_numeric <- select_if(train_df, is.numeric)
test_numeric <- select_if(test_df, is.numeric)

# Subset categorical columns with dplyr
train_categorical <- select_if(train_df,is.character)
test_categorical <- select_if(test_df,is.character)
dim(train_numeric)</pre>
```

```
> [1] 15158 11
```

```
dim(train_categorical)
```

```
> [1] 15158
               19
dim(test_numeric)
> [1] 3785
             11
dim(test_categorical)
> [1] 3785
             19
lapply(train_categorical, unique)
> $Genes.in.mother.s.side
> [1] "Yes" "No"
> $Inherited.from.father
> [1] "No" "Yes" NA
>
> $Maternal.gene
> [1] "Yes" "No"
> $Paternal.gene
> [1] "No" "Yes"
>
> $Respiratory.Rate..breaths.min.
> [1] "Normal (30-60)" "Tachypnea"
                                        NA
> $Heart.Rate..rates.min
> [1] "Normal"
                    "Tachycardia" NA
> $Follow.up
> [1] "High" "Low" NA
> $Gender
> [1] NA
                  "Male"
                               "Female"
                                           "Ambiguous"
> $Autopsy.shows.birth.defect..if.applicable.
> [1] "Not applicable" "No"
                                         "Yes"
                                                          NA
> $Folic.acid.details..peri.conceptional.
```

> [1] "No" "Yes" NA

>

```
> $H.O.serious.maternal.illness
> [1] NA
           "Yes" "No"
> $H.O.radiation.exposure..x.ray.
> [1] "No"
                       "Not applicable" "Yes"
                                                          NA
> $H.O.substance.abuse
> [1] "No"
                       "Not applicable" NA
                                                          "Yes"
> $Assisted.conception.IVF.ART
> [1] "No" "Yes" NA
> $History.of.anomalies.in.previous.pregnancies
> [1] "Yes" "No" NA
> $Birth.defects
                 "Multiple" "Singular"
> [1] NA
> $Blood.test.result
                          "normal"
                                              "inconclusive"
> [1] NA
> [4] "slightly abnormal" "abnormal"
> $Genetic.Disorder
> [1] "Mitochondrial genetic inheritance disorders"
> [2] NA
> [3] "Multifactorial genetic inheritance disorders"
> [4] "Single-gene inheritance diseases"
> $Disorder.Subclass
> [1] "Leber's hereditary optic neuropathy" "Cystic fibrosis"
> [3] "Diabetes"
                                             "Leigh syndrome"
                                             "Tay-Sachs"
> [5] "Cancer"
> [7] "Hemochromatosis"
                                             "Mitochondrial myopathy"
> [9] "Alzheimer's"
```

```
#convert all character columns to factor

df_factor_train <- as.data.frame(unclass(train_categorical), stringsAsFactors = TRUE)

df_factor_test <- as.data.frame(unclass(test_categorical), stringsAsFactors = TRUE)

str(df_factor_train)</pre>
```

```
> 'data.frame': 15158 obs. of 19 variables:
> $ Genes.in.mother.s.side
                                                 : Factor w/ 2 levels "No", "Yes": 2 2
2 2 2 2 2 1 1 2 ...
> $ Inherited.from.father
                                                 : Factor w/ 2 levels "No", "Yes": 1 2
1 1 1 1 1 1 2 1 ...
> $ Maternal.gene
                                                 : Factor w/ 2 levels "No", "Yes": 2 1
1 2 NA 2 2 2 2 NA ...
                                                 : Factor w/ 2 levels "No", "Yes": 1 1
> $ Paternal.gene
1 1 2 1 2 1 2 1 ...
> $ Respiratory.Rate..breaths.min.
                                                 : Factor w/ 2 levels "Normal (30-60)
",..: 1 2 1 2 2 NA 1 2 2 2 ...
> $ Heart.Rate..rates.min
                                                 : Factor w/ 2 levels "Normal", "Tachy
cardia": 1 1 2 1 2 1 NA 2 2 NA ...
> $ Follow.up
                                                 : Factor w/ 2 levels "High", "Low": 1
1 2 1 2 2 2 2 2 2 ...
                                                 : Factor w/ 3 levels "Ambiguous", "Fe
> $ Gender
male",..: NA NA NA 3 3 2 3 3 3 3 ...
> $ Autopsy.shows.birth.defect..if.applicable. : Factor w/ 3 levels "No", "Not appli
cable",..: 2 1 2 1 2 1 2 2 2 1 ...
> $ Folic.acid.details..peri.conceptional. : Factor w/ 2 levels "No", "Yes": 1 2
2 1 1 1 NA 2 2 1 ...
> $ H.O.serious.maternal.illness
                                                 : Factor w/ 2 levels "No", "Yes": NA
2 1 2 2 1 2 2 1 2 ...
> $ H.O.radiation.exposure..x.ray.
                                                 : Factor w/ 3 levels "No", "Not appli
cable",..: 1 2 3 NA NA 1 1 1 1 3 ...
> $ H.O.substance.abuse
                                                 : Factor w/ 3 levels "No", "Not appli
cable",..: 1 2 NA 2 2 1 2 1 1 2 ...
> $ Assisted.conception.IVF.ART
                                                 : Factor w/ 2 levels "No", "Yes": 1 1
2 NA 2 NA 2 1 2 2 ...
> $ History.of.anomalies.in.previous.pregnancies: Factor w/ 2 levels "No", "Yes": 2 2
2 2 1 1 1 2 2 NA ...
> $ Birth.defects
                                                 : Factor w/ 2 levels "Multiple", "Sin
gular": NA 1 2 2 1 1 1 1 1 2 ...
> $ Blood.test.result
                                                 : Factor w/ 4 levels "abnormal", "inc
onclusive",..: NA 3 3 2 NA 3 3 4 1 1 ...
> $ Genetic.Disorder
                                                 : Factor w/ 3 levels "Mitochondrial
genetic inheritance disorders",..: 1 NA 2 1 2 3 3 1 2 3 ...
> $ Disorder.Subclass
                                                 : Factor w/ 9 levels "Alzheimer's",.
.: 6 3 4 7 2 3 9 7 4 5 ...
```

```
#merge the tranformed dataframe with numeric vectors data
train_df_transformed= cbind(train_numeric,df_factor_train)
test_df_transformed= cbind(test_numeric,df_factor_test)
```

str(train\_df\_transformed)

```
> 'data.frame': 15158 obs. of 30 variables:
 $ Patient.Age
                                                     : int 2 4 6 12 11 14 3 11 4 6 .
                                                     : num 4.76 4.91 4.89 4.71 4.72
 $ Blood.cell.count..mcL.
                                                     : int NA NA 41 21 32 NA NA 45 4
> $ Mother.s.age
4 NA ...
                                                     : int NA 23 22 NA NA NA 63 44 4
  $ Father.s.age
2 NA ...
  $ No..of.previous.abortion
                                                     : int NA NA 4 1 4 0 3 0 1 1 ...
  $ White.Blood.cell.count..thousand.per.microliter.: num 9.86 5.52 NA 7.92 4.1 ...
                                                     : int 1 1 0 0 0 1 0 1 0 1 ...
  $ Symptom.1
                                                     : int 1 NA 1 0 0 0 0 1 0 NA ...
  $ Symptom.2
  $ Symptom.3
                                                     : int 1 1 1 1 0 0 0 1 1 0 ...
> $ Symptom.4
                                                     : int 1 1 1 0 0 1 0 0 1 0 ...
> $ Symptom.5
                                                     : int 1 0 1 0 NA 0 0 1 1 NA ...
                                                     : Factor w/ 2 levels "No", "Yes":
> $ Genes.in.mother.s.side
2 2 2 2 2 2 2 1 1 2 ...
> $ Inherited.from.father
                                                     : Factor w/ 2 levels "No", "Yes":
1 2 1 1 1 1 1 1 2 1 ...
> $ Maternal.gene
                                                     : Factor w/ 2 levels "No", "Yes":
2 1 1 2 NA 2 2 2 2 NA ...
> $ Paternal.gene
                                                     : Factor w/ 2 levels "No", "Yes":
1 1 1 1 2 1 2 1 2 1 ...
> $ Respiratory.Rate..breaths.min.
                                                     : Factor w/ 2 levels "Normal (30
-60)",..: 1 2 1 2 2 NA 1 2 2 2 ...
> $ Heart.Rate..rates.min
                                                     : Factor w/ 2 levels "Normal", "T
achycardia": 1 1 2 1 2 1 NA 2 2 NA ...
> $ Follow.up
                                                     : Factor w/ 2 levels "High", "Low
": 1 1 2 1 2 2 2 2 2 2 ...
> $ Gender
                                                     : Factor w/ 3 levels "Ambiguous"
,"Female",..: NA NA NA 3 3 2 3 3 3 3 ...
 $ Autopsy.shows.birth.defect..if.applicable.
                                                    : Factor w/ 3 levels "No", "Not a
pplicable",..: 2 1 2 1 2 1 2 2 2 1 ...
> $ Folic.acid.details..peri.conceptional.
                                                    : Factor w/ 2 levels "No", "Yes":
1 2 2 1 1 1 NA 2 2 1 ...
> $ H.O.serious.maternal.illness
                                                     : Factor w/ 2 levels "No", "Yes":
NA 2 1 2 2 1 2 2 1 2 ...
> $ H.O.radiation.exposure..x.ray.
                                                     : Factor w/ 3 levels "No", "Not a
pplicable",..: 1 2 3 NA NA 1 1 1 1 3 ...
> $ H.O.substance.abuse
                                                     : Factor w/ 3 levels "No", "Not a
pplicable",..: 1 2 NA 2 2 1 2 1 1 2 ...
                                                     : Factor w/ 2 levels "No", "Yes":
> $ Assisted.conception.IVF.ART
1 1 2 NA 2 NA 2 1 2 2 ...
> $ History.of.anomalies.in.previous.pregnancies : Factor w/ 2 levels "No", "Yes":
2 2 2 2 1 1 1 2 2 NA ...
 $ Birth.defects
                                                     : Factor w/ 2 levels "Multiple",
```

Mice imputation: MICE is a multiple imputation method used to replace missing data values in a data set under certain assumptions about the data missingness mechanism (e.g., the data are missing at random, the data are missing completely at random).

```
# mice imputation
library(mice)
set.seed(100)
train_df_imputed <- mice(train_df_transformed, m=5, maxit = 3, method = 'pmm')</pre>
```

```
iter imp variable
       1 Patient.Age Mother.s.age Father.s.age No..of.previous.abortion White.B
lood.cell.count..thousand.per.microliter. Symptom.1 Symptom.2 Symptom.3 Symptom.4
Symptom.5 Inherited.from.father Maternal.gene Respiratory.Rate..breaths.min. Hear
t.Rate..rates.min Follow.up Gender Autopsy.shows.birth.defect..if.applicable.
ic.acid.details..peri.conceptional. H.O.serious.maternal.illness H.O.radiation.expo
sure..x.ray. H.O.substance.abuse Assisted.conception.IVF.ART History.of.anomalies.
in.previous.pregnancies Birth.defects Blood.test.result Genetic.Disorder
       2 Patient.Age Mother.s.age Father.s.age No..of.previous.abortion White.B
lood.cell.count..thousand.per.microliter. Symptom.1 Symptom.2 Symptom.3 Symptom.4
Symptom.5 Inherited.from.father Maternal.gene Respiratory.Rate..breaths.min. Hear
t.Rate..rates.min Follow.up Gender Autopsy.shows.birth.defect..if.applicable. Fol
ic.acid.details..peri.conceptional. H.O.serious.maternal.illness H.O.radiation.expo
sure..x.ray. H.O.substance.abuse Assisted.conception.IVF.ART History.of.anomalies.
in.previous.pregnancies Birth.defects Blood.test.result Genetic.Disorder
       3 Patient.Age Mother.s.age Father.s.age No..of.previous.abortion White.B
lood.cell.count..thousand.per.microliter.
                                         Symptom.1 Symptom.2 Symptom.3 Symptom.4
Symptom.5 Inherited.from.father Maternal.gene Respiratory.Rate..breaths.min. Hear
t.Rate..rates.min Follow.up Gender Autopsy.shows.birth.defect..if.applicable. Fol
ic.acid.details..peri.conceptional. H.O.serious.maternal.illness H.O.radiation.expo
sure..x.ray. H.O.substance.abuse Assisted.conception.IVF.ART History.of.anomalies.
in.previous.pregnancies Birth.defects Blood.test.result Genetic.Disorder
       4 Patient.Age Mother.s.age Father.s.age No..of.previous.abortion White.B
```

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test\_df\_imputed <- mice(test\_df\_transformed, m=5, maxit = 3, method = 'pmm')</pre>

> iter imp variable

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```
set.seed(100)
train_df_imputed <- complete(train_df_imputed,3)
test_df_imputed <- complete(test_df_imputed,3)

sum(is.na(train_df_imputed))</pre>
```

> [1] 0

```
sum(is.na(test_df_imputed))
```

> [1] 0

#### **EDA**