

機器學習期末報告

Pima 印地安人糖尿病

HW16_M0928001_黃瑄惠

指導教授: 魏裕中老師

中華民國一百零九年六月十六日

壹、前言

此數據最初來自美國糖尿病、消化與腎臟疾病研究所。數據集目的是基於數據中包含的某些診斷指標，預測患者是否患有糖尿病，數據內的所有患者皆為 Pima 印地安人血統 21 歲以上的女性。將是否患有糖尿病設為 response，此數據共含 768 個實例，1 個目標變數和 8 個反應變數。

貳、變數說明

變數名稱	說明
pregnant	懷孕次數
glucose	葡萄糖，口服葡萄糖耐量測試 2 小時的血漿葡萄糖濃度
pressure	血壓，舒張壓（毫米汞柱）
triceps	皮膚厚度，三頭肌皮膚褶皺厚度（毫米）
insulin	胰島素，2 小時血清胰島素（mu U / ml）
mass	體重指數（體重（kg） / （身高（m）） ^ 2）
pedigree	糖尿病譜系函數
age	年齡（歲）
diabetes	768 個類別變量（0 或 1）中，268 個為 1，其他為 0

參、Diagnostics

<pre>> contrasts(AvaData\$diabetes) pos neg 0 pos 1</pre>	目標變數 Y : neg、pos
--	------------------

```
> table(AvaData$diabetes)
neg pos
500 268
```

neg : 500 筆 pos : 268 筆

```
> table(TrainDataY)
TrainDataY
neg pos
400 214
```

```
> table(ValDataY)
ValDataY
neg pos
100 54
```

為求實驗準確性，資料集依比例將 80% 設為 Training set、20% 設為 Validation set 並套用到所有 model 上

1. KNN

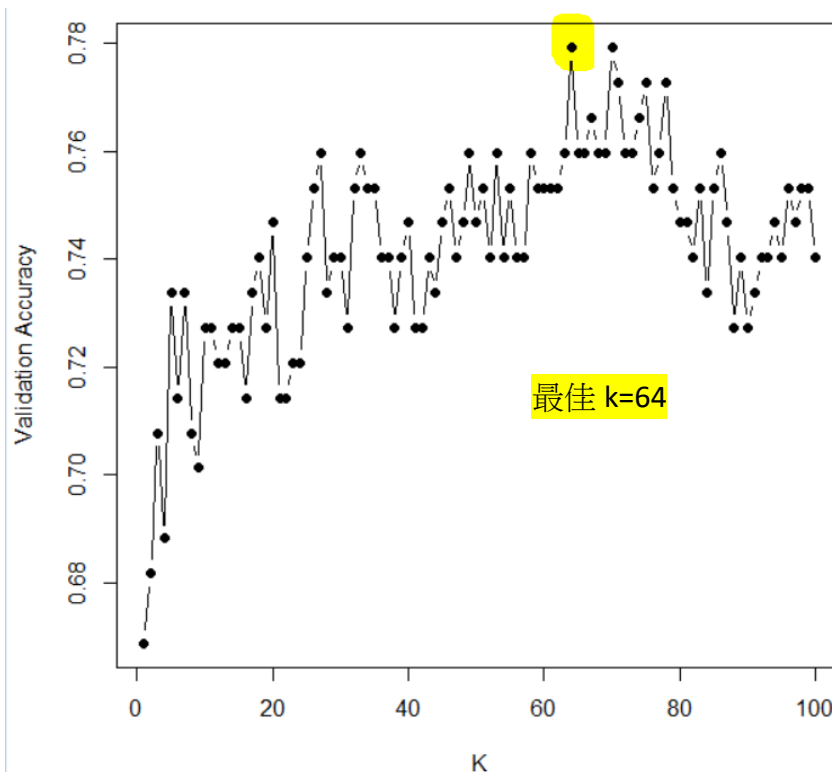
```
> data(PimaIndiansDiabetes)
> AvaData=PimaIndiansDiabetes
> AvaData$diabetes=as.factor(AvaData$diabetes)
> AvaDataX=AvaData[, -9]
> AvaDataY=AvaData[, 9]
> AvaData$diabetes=as.factor(AvaData$diabetes)
> AvaN=nrow(AvaData)
> GN=round(table(AvaData$diabetes)*0.8,0)
> set.seed(3)
> Trainget=strata(AvaData,"diabetes",size=c(GN[[2]],GN[[1]]),method="srswor" )
> TrainData=getdata(AvaData,Trainget)
> TrainInx=TrainData$ID_unit
> ValInx=c(1:AvaN)[-TrainInx]
> TrainDataX=AvaDataX[TrainInx,]
> TrainDataY=AvaDataY[TrainInx]
> ValDataX=AvaDataX[ValInx,]
> ValDataY=AvaDataY[ValInx]
> AccuracyAll=rep(1:100)
> for(i in 1:100){
+   PredY=knn(train=TrainDataX,test=ValDataX, cl=TrainDataY, k=i, prob=F)
+   #AccuracyAll[i]=confusionMatrix(PredY, ValDataY) #前面放正確後面放錯誤
+   #AccuracyAll[i]=confusionMatrix(PredY, ValDataY)$overall
+   AccuracyAll[i]=confusionMatrix(PredY, ValDataY)$overall["Accuracy"]
+ }
> OptimalK=which.max(AccuracyAll)
> OptimalK
[1] 64
> win.graph()
> plot(c(1:100), AccuracyAll, pch=19, xlab="K", ylab="Validation Accuracy", type="b")
> i=OptimalK
> PredY=knn(train=TrainDataX,test=ValDataX, cl=TrainDataY, k=i, prob=F)
> confusionMatrix(PredY, ValDataY)
```

設定 tuning parameter 為 $k=1,2,\dots,100$

以 training set 建立 knn 分類模型

計算出所有準確率，挑出準確率最大時的 k

最後列出 confusion matrix



Confusion Matrix and Statistics

```

Reference
Prediction neg pos
neg 92 29
pos 8 25

Accuracy : 0.7597
95% CI : (0.6844, 0.8248)
No Information Rate : 0.6494
P-Value [Acc > NIR] : 0.002122

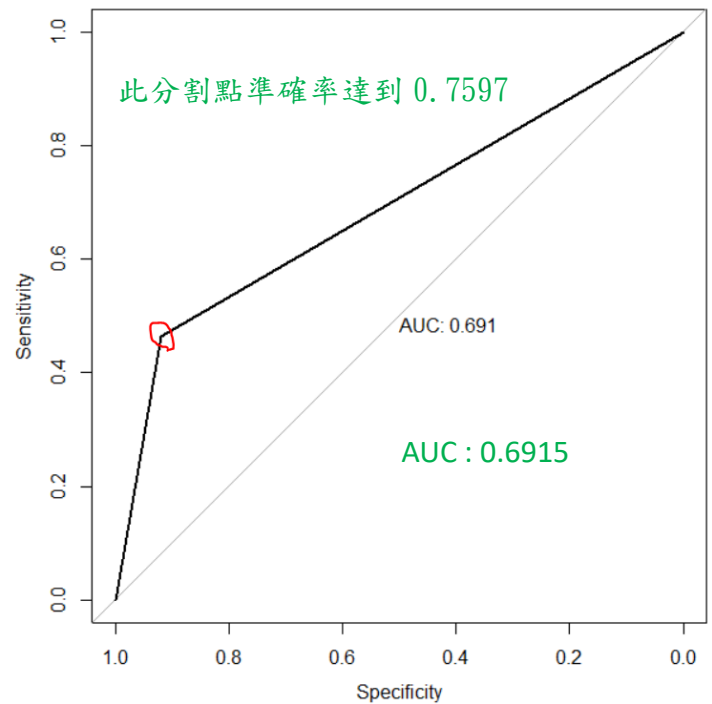
Kappa : 0.4206

McNemar's Test P-Value : 0.001009

Sensitivity : 0.9200
Specificity : 0.4630
Pos Pred Value : 0.7603
Neg Pred Value : 0.7576
Prevalence : 0.6494
Detection Rate : 0.5974
Detection Prevalence : 0.7857
Balanced Accuracy : 0.6915

'Positive' Class : neg

```



```

> win.graph()
> plot.roc(as.numeric(ValDataY),as.numeric(PredY), print.auc=TRUE)

```

	實際 neg (+)	實際 pos (-)	總計
預測 neg	92	29	121
預測 pos	8	25	33
總計	100	54	154
	Se(敏感度) =92/100 =0.92	Sp(特異度) =25/54 =0.463	準確率:117/154=0.7597

2. Logistic regression

```

> data(PimaIndiansDiabetes)
> AvaData=PimaIndiansDiabetes
> AvaData$diabetes=as.factor(AvaData$diabetes)
> AvaN=nrow(AvaData)
> GN=round(table(AvaData$diabetes)*0.8,0)
> set.seed(3)
> Trainget=strata(AvaData,"diabetes",size=c(GN[[2]],GN[[1]]),method="srswor" )
> TrainData=getdata(AvaData,Trainget)
> TrainInx=TrainData$ID_unit
> ValInx=c(1:AvaN)[-TrainInx]
> TrainData=select(TrainData,-c("ID_unit","Prob","Stratum"))
> ValData=AvaData[ValInx,]
> ModelLogl=glm(formula=diabetes~.,family=binomial,data=TrainData)
> PreProbl=predict(ModelLogl, newdata=ValData[,-9],type="response")
> PredYl=as.factor(ifelse(PreProbl>0.5, "pos", "neg"))
> confusionMatrix(PredYl, ValData$diabetes)

```

Training set 帶入建構 Logistic regression，求出預測值並計算準確率

```
> confusionMatrix(PredY1, ValData$diabetes)
Confusion Matrix and Statistics

          Reference
Prediction neg pos
neg      89  23
pos      11  31

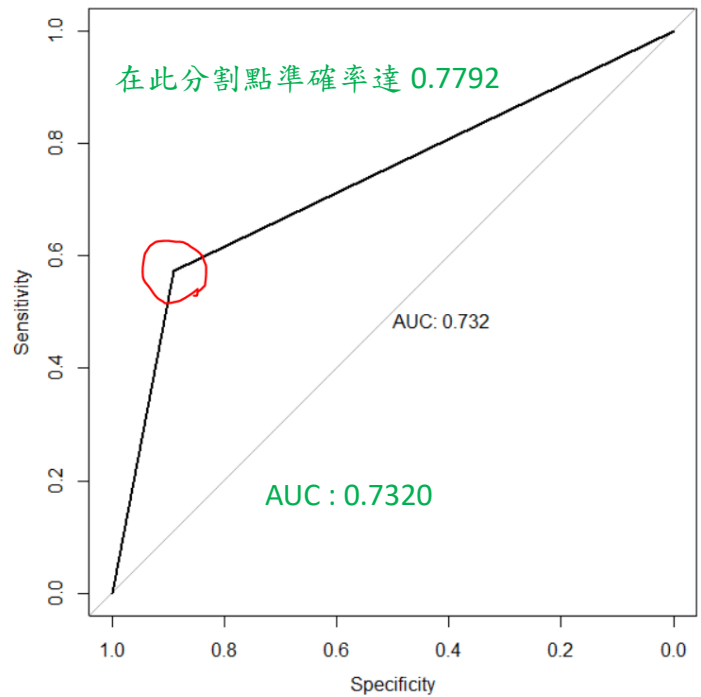
      Accuracy : 0.7792
      95% CI   : (0.7054, 0.842)
No Information Rate : 0.6494
P-Value [Acc > NIR] : 0.0003315

      Kappa : 0.4891

McNemar's Test P-Value : 0.0592297

      Sensitivity : 0.8900
      Specificity : 0.5741
      Pos Pred Value : 0.7946
      Neg Pred Value : 0.7381
      Prevalence : 0.6494
      Detection Rate : 0.5779
      Detection Prevalence : 0.7273
      Balanced Accuracy : 0.7320

      'Positive' Class : neg
```



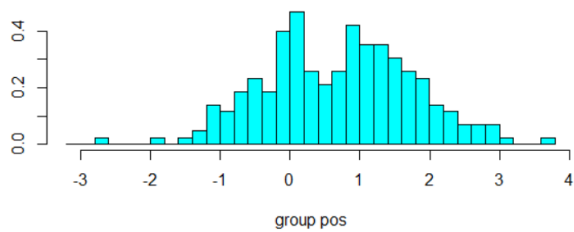
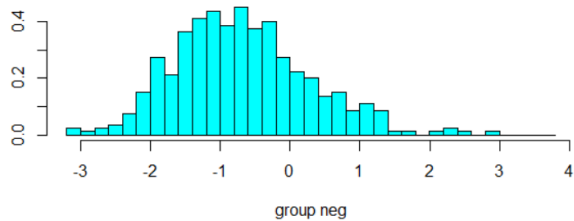
```
> win.graph()
> plot.roc(as.numeric(ValData$diabetes),as.numeric(PredY1), print.auc=TRUE)
```

	實際 neg (+)	實際 pos (-)	總計
預測 neg	89	23	112
預測 pos	11	31	42
總計	100	54	154
	Se(敏感度) =89/100 =0.89	Sp(特異度) =31/54 =0.6494	準確率:120/154=0.7792

3. LDA

```
> data(PimaIndiansDiabetes)
> AvaData=PimaIndiansDiabetes
> AvaData$diabetes=as.factor(AvaData$diabetes)
> AvaN=nrow(AvaData)
> GN=round(table(AvaData$diabetes)*0.8,0)
> set.seed(3)
> Trainget=strata(AvaData,"diabetes",size=c(GN[[2]],GN[[1]]),method="srswor" )
> TrainData=getdata(AvaData,Trainget)
> TrainInx=TrainData$ID_unit
> ValInx=c(1:AvaN)[-TrainInx]
> TrainData=TrainData[,-c(10,11,12)]
> ValData=AvaData[ValInx,]
> ModelLDA=lda(formula=diabetes~.,data=AvaData, subset=TrainInx) #此處放全資料
> plot(ModelLDA)
> PredY=predict(ModelLDA, newdata=ValData[,-9],type="response")$class
> confusionMatrix(PredY, ValData$diabetes)
```

以 training set 建立 LDA 模型並以 validation set 的代入建構的 LDA model，得到 Y 的預測情況並計算準確率



Y 變數 neg、pos 接近呈常態
兩者分散程度差異不大
適合用 Logistic regression

```
Confusion Matrix and Statistics

      Reference
Prediction neg pos
neg      90  23
pos      10  31

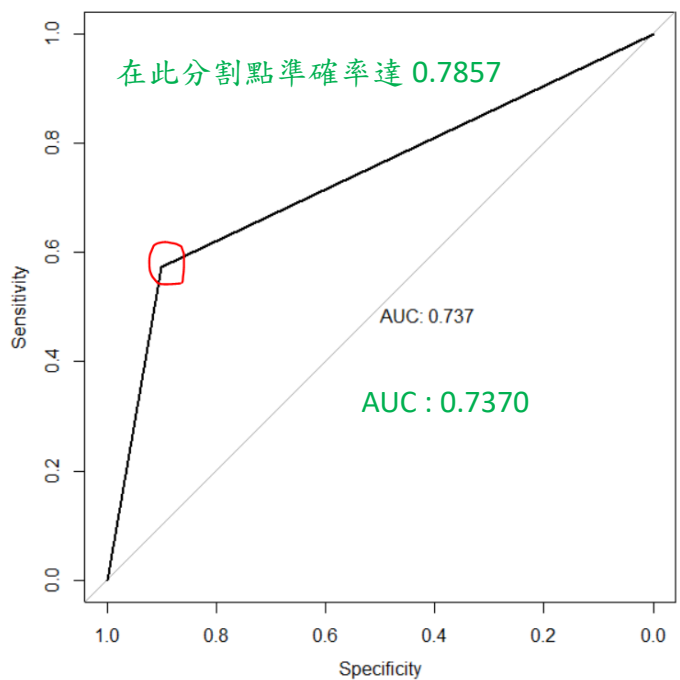
      Accuracy : 0.7857
      95% CI   : (0.7124, 0.8477)
No Information Rate : 0.6494
P-Value [Acc > NIR] : 0.0001665

      Kappa : 0.5019

McNemar's Test P-Value : 0.0367139

      Sensitivity : 0.9000
      Specificity : 0.5741
      Pos Pred Value : 0.7965
      Neg Pred Value : 0.7561
      Prevalence : 0.6494
      Detection Rate : 0.5844
      Detection Prevalence : 0.7338
      Balanced Accuracy : 0.7370

      'Positive' Class : neg
```



```
> win.graph()
> plot.roc(as.numeric(ValData$diabetes), as.numeric(PredY), print.auc=TRUE)
```

	實際 neg (+)	實際 pos (-)	總計
預測 neg	90	23	113
預測 pos	10	31	41
總計	100	54	154
	Se(敏感度) =90/100 =0.9	Sp(特異度) =31/54 =0.5741	正確率:121/154=0.7857

4. QDA

```
> data(PimaIndiansDiabetes)
> AvaData=PimaIndiansDiabetes
> AvaData$diabetes=as.factor(AvaData$diabetes)
> AvaN=nrow(AvaData)
> GN=round(table(AvaData$diabetes)*0.8,0)
> set.seed(3)
> Trainget=strata(AvaData,"diabetes",size=c(GN[[2]],GN[[1]]), method="srswor" )
> TrainData=getdata(AvaData,Trainget)
> TrainInx=TrainData$ID_unit
> ValInx=c(1:AvaN)[-TrainInx]
> TrainData=TrainData[,-c(10,11,12)]
> ValData=AvaData[ValInx,]
> ModelQDA=qda(formula=diabetes~.,data=AvaData, subset=TrainInx)
> PredY=predict(ModelQDA, newdata=ValData[,-9],type="response")$class
> confusionMatrix(PredY, ValData$diabetes)
```

以 training set 建構 QDA model，以 validation set 代入建構的 QDA model 得到 Y 的預測情況並計算出準確率

```
Confusion Matrix and Statistics

      Reference
Prediction neg pos
neg      88  24
pos      12  30

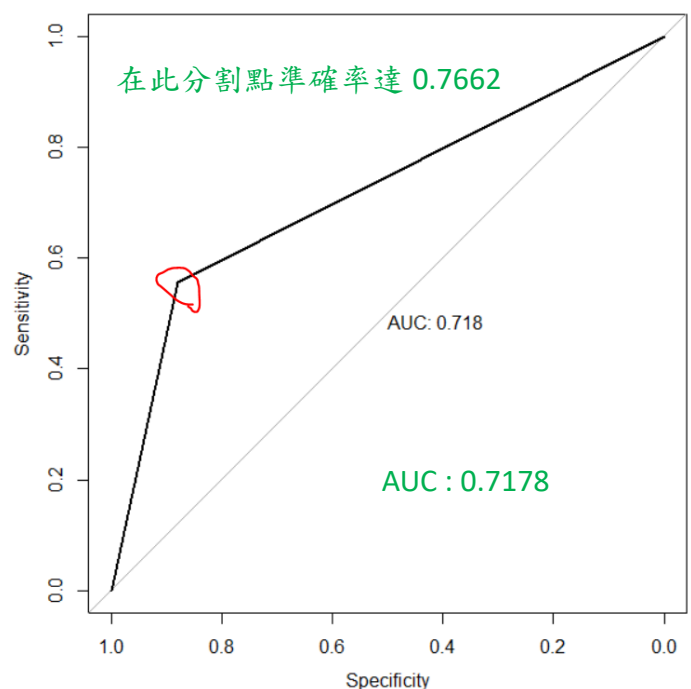
      Accuracy : 0.7662
      95% CI   : (0.6914, 0.8306)
No Information Rate : 0.6494
P-Value [Acc > NIR] : 0.001182

      Kappa : 0.459

McNemar's Test P-Value : 0.066753

      Sensitivity : 0.8800
      Specificity : 0.5556
      Pos Pred Value : 0.7857
      Neg Pred Value : 0.7143
      Prevalence : 0.6494
      Detection Rate : 0.5714
      Detection Prevalence : 0.7273
      Balanced Accuracy : 0.7178

      'Positive' Class : neg
```



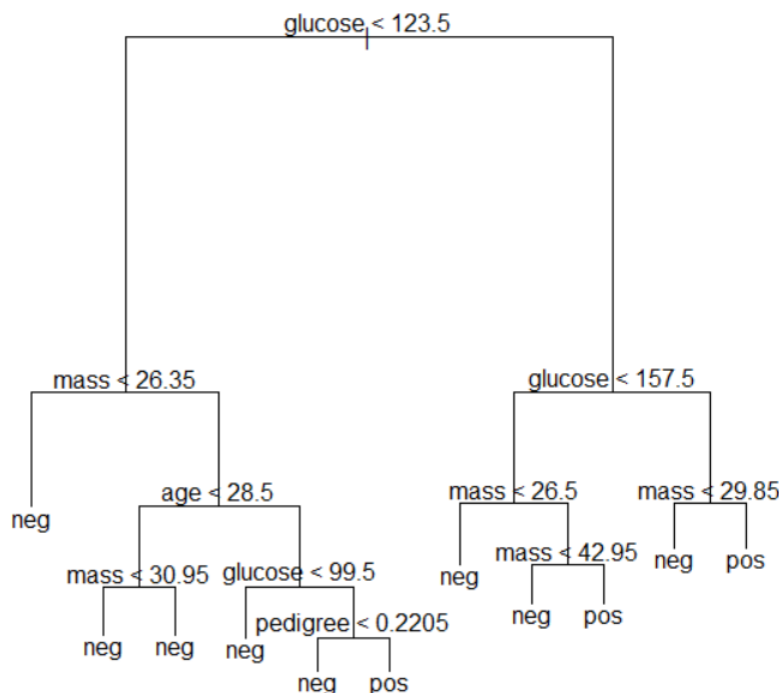
```
> win.graph()
> plot.roc(as.numeric(ValData$diabetes),as.numeric(PredY), print.auc=TRUE)
```

	實際 neg (+)	實際 pos (-)	總計
預測 neg	88	24	112
預測 pos	12	30	42
總計	100	54	154
	Se(敏感度) =88/100 =0.88	Sp(特異度) =30/54 =0.5556	正確率:118/154=0.7662

5. Classification Tree

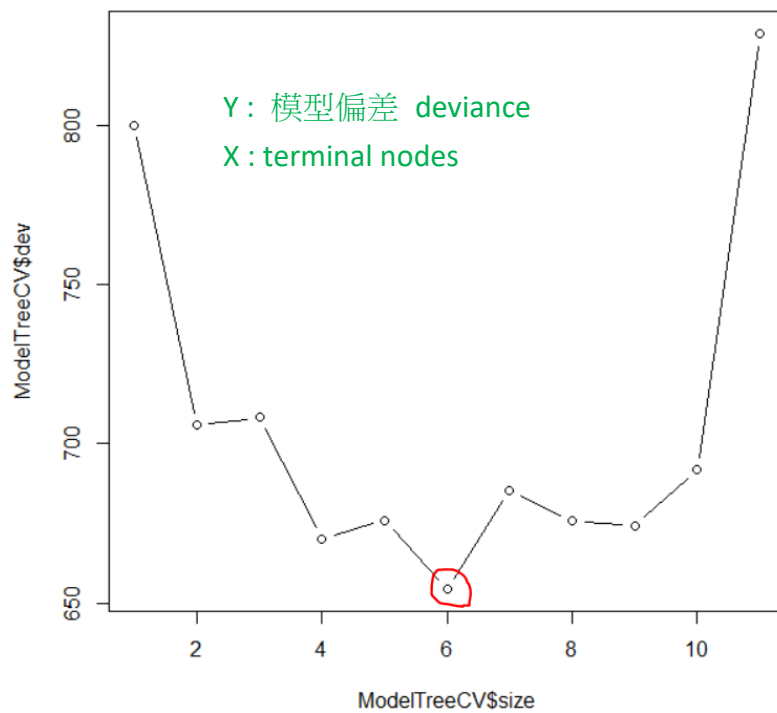
```
> data(PimaIndiansDiabetes)
> AvaData=PimaIndiansDiabetes
> AvaData$diabetes=as.factor(AvaData$diabetes)
> AvaN=nrow(AvaData)
> GN=round(table(AvaData$diabetes)*0.8,0) #先做一個table分成R、S0,再各取80%,才
> set.seed(3)
> Trainget=strata(AvaData,"diabetes",size=c(GN[[2]],GN[[1]]),method="srswor" )
> TrainData=getdata(AvaData,Trainget)
> TrainInx=TrainData$ID_unit
> ValInx=c(1:AvaN)[-TrainInx]
> TrainData=TrainData[,-c(10,11,12)]
> ValData=AvaData[ValInx,]
>
> ModelTree=tree(diabetes~., data=AvaData, subset=TrainInx) #data放全資料
> win.graph()
> plot(ModelTree)
> text(ModelTree) #文字附上
> ModelTreeCV=cv.tree(ModelTree) #用cross validation方法看不同尺寸的樹
> win.graph()
> plot(ModelTreeCV$size, ModelTreeCV$dev, type="b")
>
> ModelPruneTree=prune.tree(ModelTree,best=6)
> win.graph()
> plot(ModelPruneTree)
> text(ModelPruneTree)
>
> PredProbY=predict(ModelPruneTree,ValData[, -9])
> PredY=as.factor(ifelse(as.data.frame(PredProbY)$pos>0.5,"pos","neg"))
> confusionMatrix(PredY, ValData$diabetes)
```

由 training set 建構一棵 classification tree



11 個 terminal nodes

接著用 10—fold cross validation 方法來看不同尺寸的樹，並判斷是否修剪樹

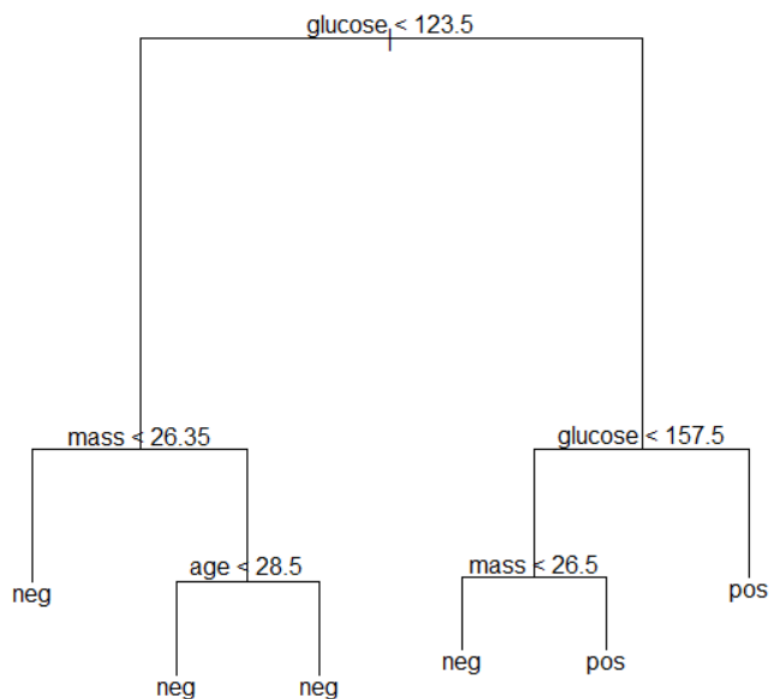


Deviance :模型偏差

模型偏差大小可以反映一個模型你和數據的程度，偏差高代表該模型對數據的擬合越差。

因而選擇 deviance(dev)最低的，或者 deviance 下降程度趨於平緩時的 terminal nodes 數(size)來修剪樹。

將 validation data 的 X 帶入修剪過後的樹，得到 Y 預測值並計算 accuracy



由挑選出的最佳 tree size:6 來建構樹

Confusion Matrix and Statistics

```

Reference
Prediction neg pos
neg 83 19
pos 17 35

Accuracy : 0.7662
95% CI : (0.6914, 0.8306)
No Information Rate : 0.6494
P-Value [Acc > NIR] : 0.001182

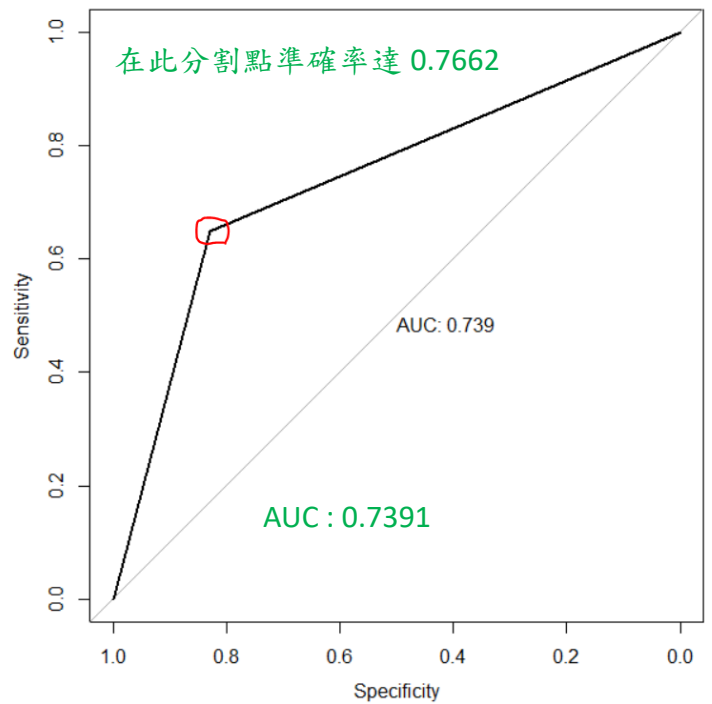
Kappa : 0.4823

McNemar's Test P-Value : 0.867632

Sensitivity : 0.8300
Specificity : 0.6481
Pos Pred Value : 0.8137
Neg Pred Value : 0.6731
Prevalence : 0.6494
Detection Rate : 0.5390
Detection Prevalence : 0.6623
Balanced Accuracy : 0.7391

'Positive' Class : neg

```



```

> win.graph()
> plot.roc(as.numeric(ValData$diabetes),as.numeric(PredY), print.auc=TRUE)

```

	實際 neg (+)	實際 pos (-)	總計
預測 neg	83	19	102
預測 pos	17	35	52
總計	100	54	154
	Se(敏感度) =83/100 =0.83	Sp(特異度) =35/54 =0.6481	正確率:118/154=0.7662

6. Bagging

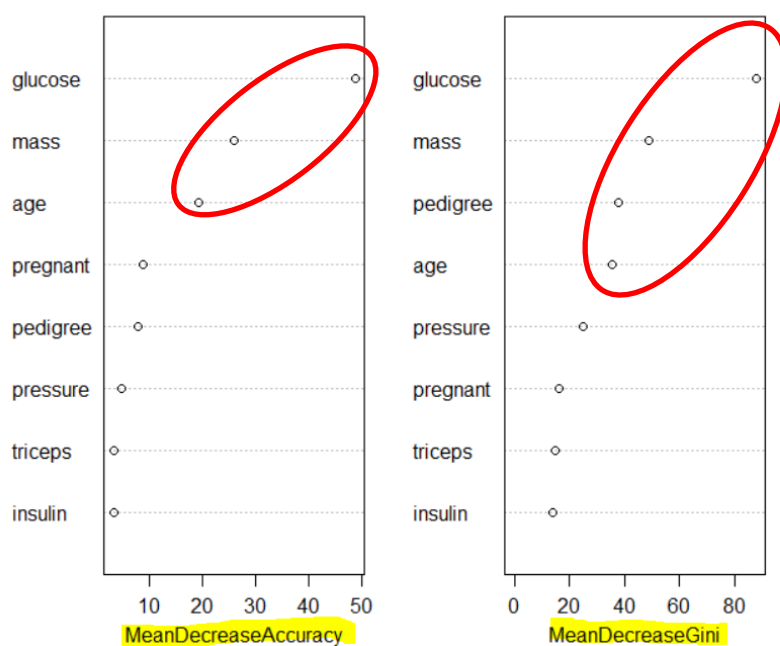
```

> data(PimaIndiansDiabetes)
> AvaData=PimaIndiansDiabetes
> AvaData$diabetes=as.factor(AvaData$diabetes)
> AvaN=nrow(AvaData)
> GN=round(table(AvaData$diabetes)*0.8,0)
> set.seed(3)
> Trainget=strata(AvaData,"diabetes",size=c(GN[[2]],GN[[1]]),method="srswor" )
> TrainData=getdata(AvaData,Trainget)
> TrainInx=TrainData$ID_unit
> ValInx=c(1:AvaN)[-TrainInx]
> TrainData=TrainData[,-c(10,11,12)]
> ValData=AvaData[ValInx,]
> ModelBag=randomForest(diabetes~., data=AvaData,subset=TrainInx, mtry=8, importance=T)
> PredY=predict(ModelBag,newdata=ValData[,-9],type="response")
> confusionMatrix(PredY, ValData$diabetes)

```

由 training set 建構 bagging model

ModelBag



MeanDecreaseAccuracy:

準確率越高越好，平均下降的準確率越大代表變數 X 越重要。

MeanDecreaseGini:

Gini 越大純度越高，平均下降的 Gini 值越大代表變數 X 越重要。

```
> importance(ModelBag)
```

	neg	pos	MeanDecreaseAccuracy	MeanDecreaseGini
pregnant	10.170509	-1.2240719	8.967861	16.22616
glucose	36.625175	35.9267150	48.686084	87.66958
pressure	7.292382	-0.9882496	4.824362	24.63866
triceps	5.353317	-1.5340826	3.524558	14.59854
insulin	4.275927	-0.2977598	3.432922	13.59205
mass	17.571579	20.8004269	26.078701	48.67089
pedigree	8.431507	2.9673263	7.875888	37.92952
age	16.098479	8.8769861	19.238684	35.42319

由上圖也可看出 glucose、mass、age 三變數較為重要

左邊兩行數據顯示，少了此變數後影響 Y(neg、pos)準確率，使之下降程度

接著由 validation set 的 X 帶入 bagging model 預測 Y，並計算準確率

Confusion Matrix and Statistics

Reference

Prediction neg pos

neg 82 21

pos 18 33

Accuracy : 0.7468

95% CI : (0.6705, 0.8133)

No Information Rate : 0.6494

P-Value [Acc > NIR] : 0.006192

Kappa : 0.4367

Mcnemar's Test P-Value : 0.748774

Sensitivity : 0.8200

Specificity : 0.6111

Pos Pred Value : 0.7961

Neg Pred Value : 0.6471

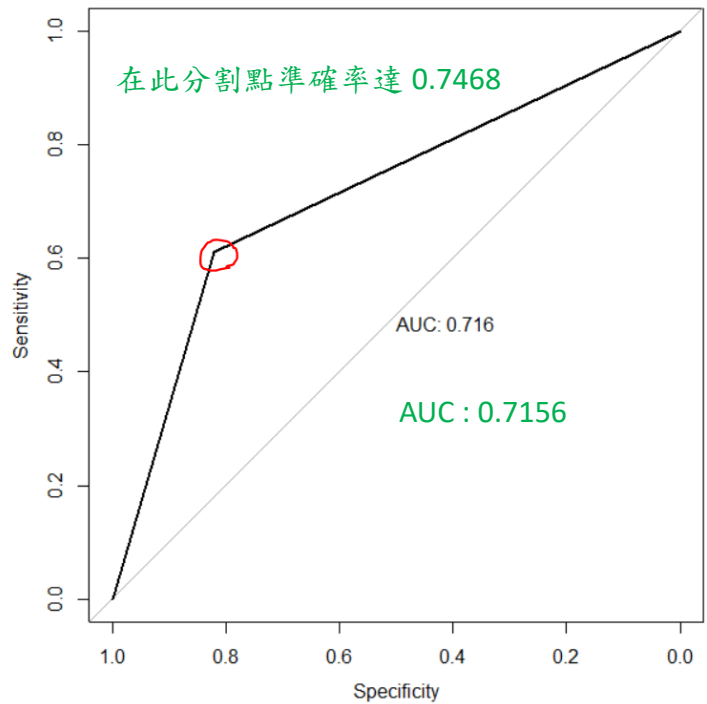
Prevalence : 0.6494

Detection Rate : 0.5325

Detection Prevalence : 0.6688

Balanced Accuracy : 0.7156

'Positive' Class : neg



```
> win.graph()
```

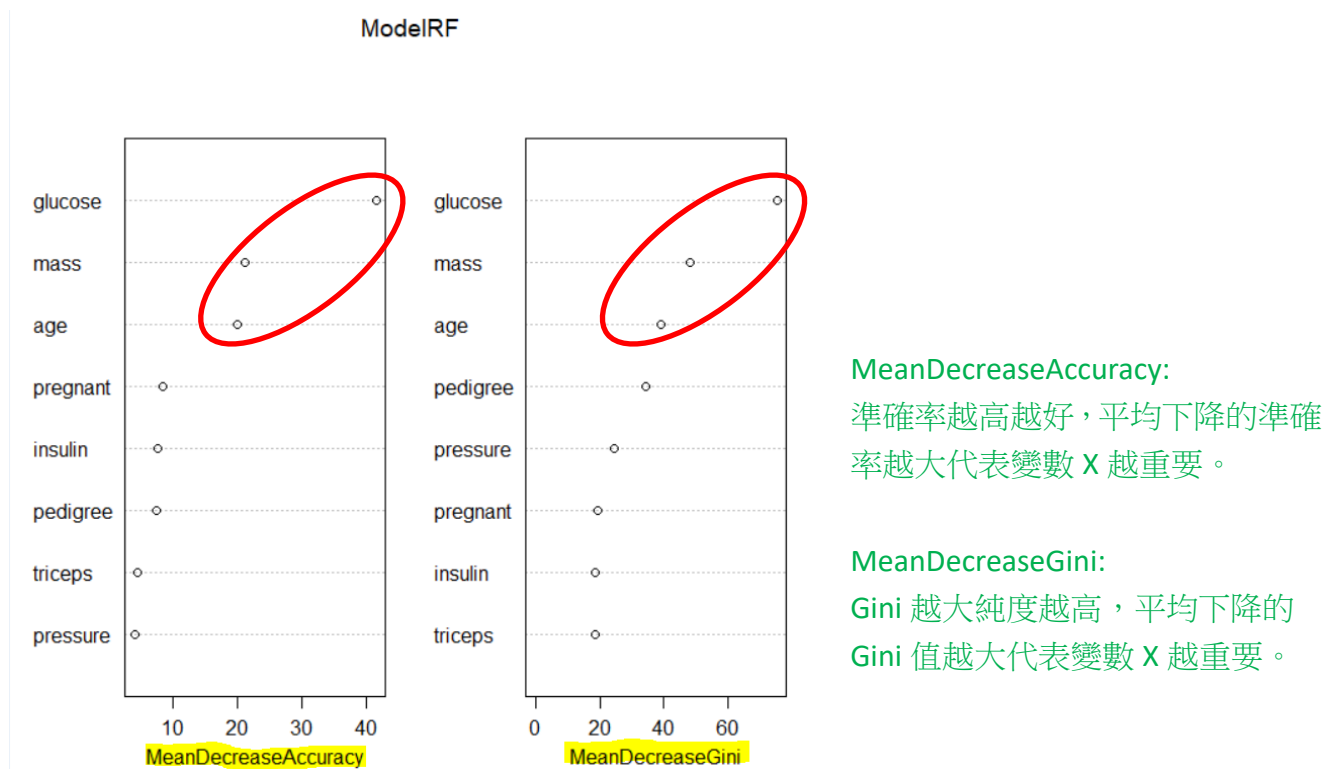
```
> plot.roc(as.numeric(ValData$diabetes),as.numeric(PredY), print.auc=TRUE)
```

	實際 neg (+)	實際 pos (-)	總計
預測 neg	82	21	103
預測 pos	18	33	51
總計	100	54	154
	Se(敏感度) =82/100 =0.82	Sp(特異度) =33/54 0.6111	正確率:115/154=0.7468

7. Random Forest

```
> data(PimaIndiansDiabetes)
> AvaData=PimaIndiansDiabetes
> AvaData$diabetes=as.factor(AvaData$diabetes)
> AvaN=nrow(AvaData)
> GN=round(table(AvaData$diabetes)*0.8,0)
> set.seed(3)
> Trainget=strata(AvaData,"diabetes",size=c(GN[[2]],GN[[1]]),method="srswor" )
> TrainData=getdata(AvaData,Trainget)
> TrainInx=TrainData$ID_unit
> ValInx=c(1:AvaN)[-TrainInx]
> TrainData=TrainData[-c(10,11,12)]
> ValData=AvaData[ValInx,]
> ModelRF=randomForest(diabetes~., data=AvaData, subset=TrainInx, mtry=3, importance=T)
> PredY=predict(ModelRF,newdata=ValData[, -9], type="response")
> confusionMatrix(PredY, ValData$diabetes)
```

由 training set 建構 random forest model



```
> importance (ModelRF)
```

	neg	pos	MeanDecreaseAccuracy	MeanDecreaseGini
pregnant	9.218628	0.9054926	8.318209	19.29858
glucose	32.264027	31.1673503	41.588111	75.25744
pressure	7.311201	-3.0296233	4.046469	24.47332
triceps	3.112462	2.7401431	4.413186	18.56175
insulin	4.760389	5.7707602	7.691685	18.67357
mass	14.149911	17.7109559	21.136527	48.26376
pedigree	7.700656	2.1184884	7.369882	34.50302
age	14.194014	12.0399585	20.018753	39.20679

由上圖也可看出 glucose、mass、age 三變數較為重要
左邊兩行數據顯示，少了此變數後影響 Y(neg、pos)準確率，使之下降程度

接著由 validation set 的 X 帶入 bagging model 預測 Y，並計算準確率

Confusion Matrix and Statistics

```

Reference
Prediction neg pos
neg 82 21
pos 18 33

Accuracy : 0.7468
95% CI : (0.6705, 0.8133)
No Information Rate : 0.6494
P-Value [Acc > NIR] : 0.006192

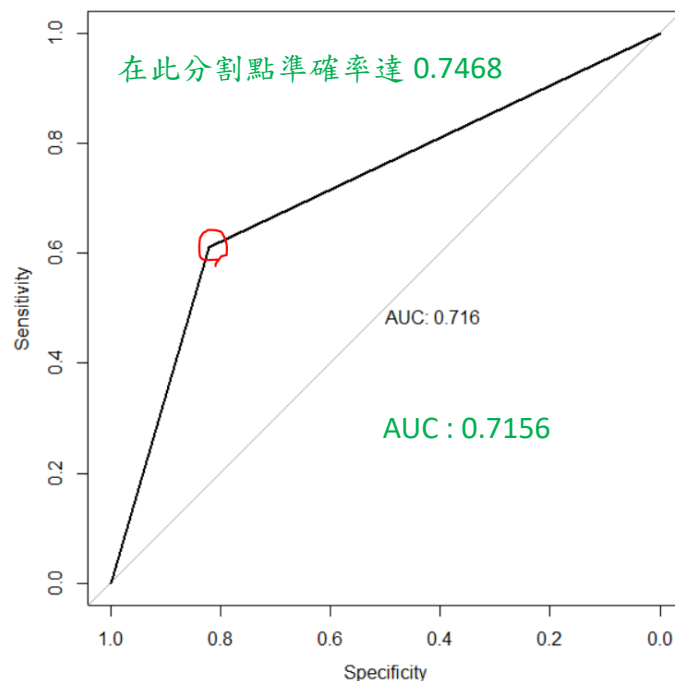
Kappa : 0.4367

McNemar's Test P-Value : 0.748774

Sensitivity : 0.8200
Specificity : 0.6111
Pos Pred Value : 0.7961
Neg Pred Value : 0.6471
Prevalence : 0.6494
Detection Rate : 0.5325
Detection Prevalence : 0.6688
Balanced Accuracy : 0.7156

'Positive' Class : neg

```



```

> win.graph()
> plot.roc(as.numeric(ValData$diabetes), as.numeric(PredY), print.auc=TRUE)

```

	實際 neg (+)	實際 pos (-)	總計
預測 neg	82	21	103
預測 pos	18	33	51
總計	100	54	154
	Se(敏感度) =82/100 =0.82	Sp(特異度) =33/54 =0.6111	正確率:115/154=0.7468

發現：Bagging 和 Random forest 的預測結果差不多。

8. Support Vector

```

> data(PimaIndiansDiabetes)
> AvaData=PimaIndiansDiabetes
> AvaData$diabetes=as.factor(AvaData$diabetes)
> AvaN=nrow(AvaData)
> GN=round(table(AvaData$diabetes)*0.8,0)
> set.seed(3)
> Trainget=strata(AvaData,"diabetes",size=c(GN[[2]],GN[[1]]),method="srswor" )
> TrainData=getdata(AvaData,Trainget)
> TrainInx=TrainData$ID_unit
> ValInx=c(1:AvaN)[-TrainInx]
> TrainData=TrainData[-c(10,11,12)]
> ValData=AvaData[ValInx,]
> Costlist=c(0.001,0.01,0.1,1,5,10,100)
> AccuSumm=rep(0,length(Costlist))
> for(i in 1:length(Costlist)){
+ svmfitTemp=svm(diabetes~., data=TrainData,kernel="linear", cost=Costlist[i], scale=F)
+ PredYTemp=predict(svmfitTemp, newdata=ValData[, -9], type="response")
+ AccuSumm[i]=confusionMatrix(PredYTemp, ValData$diabetes)$overall["Accuracy"]
+ }

> BestC=which.max(AccuSumm)
> svmfit=svm(diabetes~., data=AvaData, kernel="linear", cost=Costlist[BestC], scale=F)
> PredY=predict(svmfit, newdata=ValData[, -9], type="response")
> confusionMatrix(PredY, ValData$diabetes)

```

設定各種 cost(tuning parameter)的值(0.001,0.01,0.1,1,5,10,100)

對每個 tuning parameter 值，以 training set 建構 model 並以 validation set 的 X 代入建構的 model，得到 Y 的預測情況並計算準確率

```
> svmfit
```

```
Call:
```

```
svm(formula = diabetes ~ ., data = AvaData, kernel = "linear", cost = Costlist[BestC],  
     scale = F)
```

```
Parameters:
```

```
SVM-Type: C-classification
```

```
SVM-Kernel: linear
```

```
cost: 0.01
```

```
Number of Support Vectors: 406
```

找到準確率最高的最佳 tuning parameter value

-->0.01

```
Confusion Matrix and Statistics
```

```
Reference
```

```
Prediction neg pos
```

```
neg 91 22
```

```
pos 9 32
```

```
Accuracy : 0.7987
```

```
95% CI : (0.7266, 0.8589)
```

```
No Information Rate : 0.6494
```

```
P-Value [Acc > NIR] : 3.76e-05
```

```
Kappa : 0.5321
```

```
McNemar's Test P-Value : 0.03114
```

```
Sensitivity : 0.9100
```

```
Specificity : 0.5926
```

```
Pos Pred Value : 0.8053
```

```
Neg Pred Value : 0.7805
```

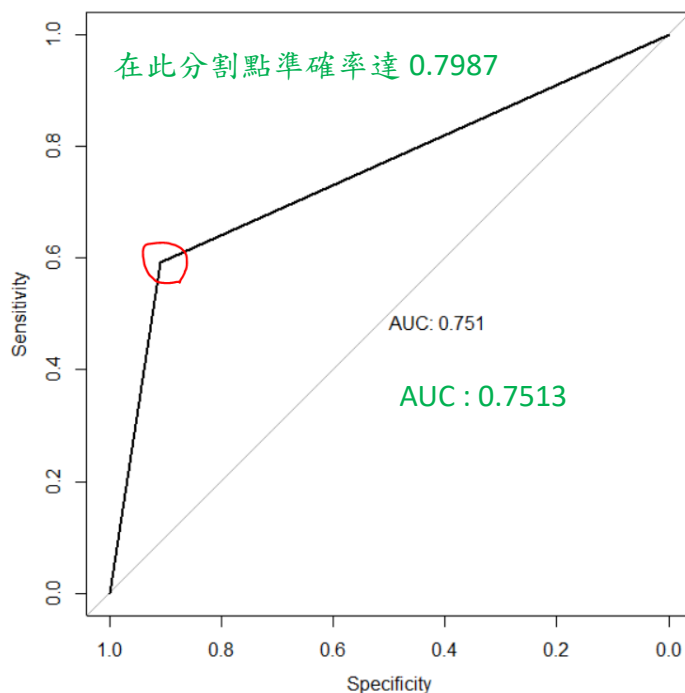
```
Prevalence : 0.6494
```

```
Detection Rate : 0.5909
```

```
Detection Prevalence : 0.7338
```

```
Balanced Accuracy : 0.7513
```

```
'Positive' Class : neg
```



```
> win.graph()
```

```
> plot.roc(as.numeric(ValData$diabetes), as.numeric(PredY), print.auc=TRUE)
```

	實際 neg (+)	實際 pos (-)	總計
預測 neg	91	22	113
預測 pos	9	32	41
總計	100	54	154
	Se(敏感度) =91/100 =0.91	Sp(特異度) =32/54 =0.5926	正確率:123/154=0.7987

9. Neural network

```
> data(PimaIndiansDiabetes)
> AvaData=PimaIndiansDiabetes
> AvaData$diabetes=as.factor(AvaData$diabetes)
> AvaN=nrow(AvaData)
> GN=round(table(AvaData$diabetes)*0.8,0)
> set.seed(3)
> Trainget=strata(AvaData,"diabetes",size=c(GN[[2]],GN[[1]]),method="srswor" )
> TrainData=getdata(AvaData,Trainget)
> TrainInx=TrainData$ID_unit
> ValInx=c(1:AvaN)[-TrainInx]
> TrainData=TrainData[-c(10,11,12)]
> ValData=AvaData[ValInx,]
> NNfit=nnet(diabetes~., data=TrainData, size=25, range=0.7)
# weights: 251
initial value 493.071588
iter 10 value 368.892983
iter 20 value 354.408205
iter 30 value 343.291718
iter 40 value 333.208536
iter 50 value 320.507387
iter 60 value 314.469485
iter 70 value 303.304727
iter 80 value 297.108047
iter 90 value 288.648807
iter 100 value 281.086314
final value 281.086314
stopped after 100 iterations
> PredY=predict(NNfit,newdata=ValData[,-9],type="class")
> confusionMatrix(as.factor(PredY), ValData$diabetes)
```

由 training set 建構 neural network model

設定 number of unit in the hidden layer (size) =25

設定 initial weight 的 range 為 [-0.7,0.7]

Confusion Matrix and Statistics

	Reference	
Prediction	neg	pos
neg	80	28
pos	20	26

AUC : 0.6883

在此分割點準確率達 0.6407

Accuracy : 0.6883
95% CI : (0.6088, 0.7604)
No Information Rate : 0.6494
P-Value [Acc > NIR] : 0.1768

Kappa : 0.2914

McNemar's Test P-Value : 0.3123

Sensitivity : 0.8000
Specificity : 0.4815
Pos Pred Value : 0.7407
Neg Pred Value : 0.5652
Prevalence : 0.6494
Detection Rate : 0.5195
Detection Prevalence : 0.7013
Balanced Accuracy : 0.6407

'Positive' Class : neg

用此方法的準確率卻不高的原因可能是因為本身資料集的 Y 就不均勻

```
> table(AvaData$diabetes)
```

neg	pos
500	268

	實際 neg (+)	實際 pos (-)	總計
預測 neg	80	28	108
預測 pos	20	26	46
總計	100	54	154
	Se(敏感度) =80/100 =0.8	Sp(特異度) =26/54 =0.4815	正確率:106/154=0.6883

肆、總結

	Se(敏感度)	Sp(特異度)	準確率
KNN	0.92	0.4630	0.7597
LOGISTIC	0.89	0.6494	0.7792
LDA	0.90	0.5741	0.7857
QDA	0.88	0.5556	0.7662
CLASSIFICATION TREE	0.83	0.6481	0.7662
BAGGING	0.82	0.6111	0.7468
RANDOM FOREST	0.82	0.6111	0.7468
SUPPORT VECTOR	0.91	0.5926	0.7987
NEURAL NETWORK	0.80	0.4815	0.6883

用 neural network 方法的準確率卻不高的原因可能是因為本身資料集的 Y 就不均勻

```
> table(AvaData$diabetes)

neg pos
500 268
```

伍、資料來源

<https://www.kaggle.com/uciml/pima-indians-diabetes-database>