

# MATH 4322 Final Project Group 9

## Introduction

### Logistic Regression (Ryan Nguyen, Alan Johnson)

Paragraph explaining why we are using logistic regression models and the advantages and disadvantages of the model.

## Model Formula

$$P(\text{cardio} = 1|X) = \frac{e^{(\beta_0 + \beta_1 X_1 + \beta_2 X_2 + \dots + \beta_n X_n)}}{1 + e^{(\beta_0 + \beta_1 X_1 + \beta_2 X_2 + \dots + \beta_n X_n)}}$$

### Model 1 - Include all predictors

```
library(readr)
cardio_train <- read_delim("cardio_train.csv",
                           delim = ";", escape_double = FALSE, trim_ws = TRUE)
```

Rows: 70000 Columns: 13

-- Column specification -----

Delimiter: ";"

dbl (13): id, age, gender, height, weight, ap\_hi, ap\_lo, cholesterol, gluc, ...

i Use `spec()` to retrieve the full column specification for this data.

i Specify the column types or set `show\_col\_types = FALSE` to quiet this message.

```
cardio_train$gender = as.factor(cardio_train$gender)
cardio_train$cholesterol = as.factor(cardio_train$cholesterol)
cardio_train$gluc = as.factor(cardio_train$gluc)
cardio_train$smoke = as.factor(cardio_train$smoke)
```

```

cardio_train$alco = as.factor(cardio_train$alco)
cardio_train$active = as.factor(cardio_train$active)
cardio_train$cardio = as.factor(cardio_train$cardio)

heart.logistic1 = glm(cardio ~ . - id, family = "binomial",
                      data = cardio_train)

```

Warning: glm.fit: algorithm did not converge

Warning: glm.fit: fitted probabilities numerically 0 or 1 occurred

```
summary(heart.logistic1)
```

Call:

```
glm(formula = cardio ~ . - id, family = "binomial", data = cardio_train)
```

Deviance Residuals:

Min	1Q	Median	3Q	Max
-8.4904	-0.9635	-0.0980	0.9907	4.6621

Coefficients:

	Estimate	Std. Error	z value	Pr(> z )	
(Intercept)	-8.084e+00	2.213e-01	-36.535	< 2e-16	***
age	1.485e-04	3.557e-06	41.735	< 2e-16	***
gender2	1.430e-02	2.107e-02	0.679	0.497	
height	-5.626e-03	1.232e-03	-4.567	4.95e-06	***
weight	1.521e-02	6.607e-04	23.023	< 2e-16	***
ap_hi	3.951e-02	6.057e-04	65.235	< 2e-16	***
ap_lo	3.004e-04	6.735e-05	4.460	8.18e-06	***
cholesterol2	4.222e-01	2.593e-02	16.285	< 2e-16	***
cholesterol3	1.134e+00	3.444e-02	32.929	< 2e-16	***
gluc2	3.011e-02	3.438e-02	0.876	0.381	
gluc3	-3.387e-01	3.809e-02	-8.894	< 2e-16	***
smoke1	-1.314e-01	3.320e-02	-3.958	7.57e-05	***
alco1	-1.695e-01	4.026e-02	-4.211	2.54e-05	***
active1	-2.101e-01	2.105e-02	-9.981	< 2e-16	***

---

Signif. codes: 0 '\*\*\*' 0.001 '\*\*' 0.01 '\*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for binomial family taken to be 1)

Null deviance: 97041 on 69999 degrees of freedom  
Residual deviance: 80883 on 69986 degrees of freedom  
AIC: 80911

Number of Fisher Scoring iterations: 25

Paragraph explaining which predictors are significant (look at significance table output)

**Model 2 - Only include statistically significant predictors**

```
heart.logistic2 = glm(cardio ~ age+height+weight+ap_hi+ap_lo+cholesterol+smoke+alco+act
                    , family = "binomial",
                    data = cardio_train)

summary(heart.logistic2)
```

Call:

```
glm(formula = cardio ~ age + height + weight + ap_hi + ap_lo +
     cholesterol + smoke + alco + active, family = "binomial",
     data = cardio_train)
```

Deviance Residuals:

Min	1Q	Median	3Q	Max
-8.4904	-0.9639	-0.0992	0.9900	4.6678

Coefficients:

	Estimate	Std. Error	z value	Pr(> z )
(Intercept)	-8.124e+00	2.028e-01	-40.062	< 2e-16 ***
age	1.476e-04	3.551e-06	41.550	< 2e-16 ***
height	-5.356e-03	1.103e-03	-4.857	1.19e-06 ***
weight	1.516e-02	6.586e-04	23.023	< 2e-16 ***
ap_hi	3.960e-02	6.047e-04	65.485	< 2e-16 ***
ap_lo	3.028e-04	6.765e-05	4.475	7.63e-06 ***
cholesterol2	4.234e-01	2.497e-02	16.959	< 2e-16 ***
cholesterol3	9.855e-01	2.962e-02	33.275	< 2e-16 ***
smoke1	-1.222e-01	3.205e-02	-3.812	0.000138 ***
alco1	-1.641e-01	4.013e-02	-4.090	4.31e-05 ***
active1	-2.085e-01	2.104e-02	-9.909	< 2e-16 ***

---

Signif. codes: 0 '\*\*\*' 0.001 '\*\*' 0.01 '\*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for binomial family taken to be 1)

Null deviance: 97041 on 69999 degrees of freedom  
Residual deviance: 80984 on 69989 degrees of freedom  
AIC: 81006

Number of Fisher Scoring iterations: 8

```
step(heart.logistic2)
```

Start: AIC=81005.71

cardio ~ age + height + weight + ap\_hi + ap\_lo + cholesterol +  
smoke + alco + active

	Df	Deviance	AIC
- smoke	1	80978	80998
- alco	1	80978	80998
<none>		80984	81006
- ap_lo	1	80991	81011
- height	1	80999	81019
- active	1	81062	81082
- weight	1	81324	81344
- cholesterol	2	82180	82198
- age	1	82526	82546
- ap_hi	1	88091	88111

Step: AIC=80998.02

cardio ~ age + height + weight + ap\_hi + ap\_lo + cholesterol +  
alco + active

	Df	Deviance	AIC
<none>		80978	80998
- ap_lo	1	81009	81027
- alco	1	81015	81033
- height	1	81024	81042
- active	1	81083	81101
- weight	1	81342	81360
- cholesterol	2	82196	82212
- age	1	82558	82576
- ap_hi	1	88096	88114

```
Call: glm(formula = cardio ~ age + height + weight + ap_hi + ap_lo +
          cholesterol + alco + active, family = "binomial", data = cardio_train)
```

Coefficients:

(Intercept)	age	height	weight	ap_hi
-8.0256093	0.0001480	-0.0060018	0.0151616	0.0395437
ap_lo	cholesterol2	cholesterol3	alco1	active1
0.0003027	0.4216361	0.9849712	-0.2143313	-0.2101295

Degrees of Freedom: 69999 Total (i.e. Null); 69990 Residual

Null Deviance: 97040

Residual Deviance: 80980 AIC: 81000

### Model 3 - Using predictors from stepwise regression

Paragraph explaining the results of stepwise regression

```
heart.logistic3 = glm(formula = cardio ~ age + height + weight + ap_hi + ap_lo + cholesterol + alco + active, family = "binomial", data = cardio_train)
summary(heart.logistic3)
```

Call:

```
glm(formula = cardio ~ age + height + weight + ap_hi + ap_lo +
    cholesterol + alco + active, family = "binomial", data = cardio_train)
```

Deviance Residuals:

Min	1Q	Median	3Q	Max
-8.4904	-0.9635	-0.1015	0.9910	4.6663

Coefficients:

	Estimate	Std. Error	z value	Pr(> z )
(Intercept)	-8.026e+00	2.011e-01	-39.908	< 2e-16 ***
age	1.480e-04	3.550e-06	41.684	< 2e-16 ***
height	-6.002e-03	1.090e-03	-5.507	3.66e-08 ***
weight	1.516e-02	6.586e-04	23.022	< 2e-16 ***
ap_hi	3.954e-02	6.043e-04	65.434	< 2e-16 ***
ap_lo	3.027e-04	6.753e-05	4.482	7.38e-06 ***
cholesterol2	4.216e-01	2.496e-02	16.894	< 2e-16 ***
cholesterol3	9.850e-01	2.961e-02	33.260	< 2e-16 ***
alco1	-2.143e-01	3.787e-02	-5.660	1.51e-08 ***
active1	-2.101e-01	2.104e-02	-9.989	< 2e-16 ***

---

Signif. codes: 0 '\*\*\*' 0.001 '\*\*' 0.01 '\*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for binomial family taken to be 1)

Null deviance: 97041 on 69999 degrees of freedom  
Residual deviance: 80978 on 69990 degrees of freedom  
AIC: 80998

Number of Fisher Scoring iterations: 25

## Determining Best Model

```
extract_info = function(model) {  
  deviance <- summary(model)$null.deviance  
  residual_deviance <- summary(model)$deviance  
  r_squared <- 1 - (residual_deviance / deviance)  
  AIC <- AIC(model)  
  BIC <- BIC(model)  
  
  return(c(Null_Deviance = deviance,  
           Residual_Deviance = residual_deviance,  
           R_Squared = r_squared,  
           AIC = AIC,  
           BIC = BIC))  
}  
  
# Extract information from each model  
info1 <- extract_info(heart.logistic1)  
info2 <- extract_info(heart.logistic2)  
info3 <- extract_info(heart.logistic3)  
  
# Create a data frame to store the information  
model_info <- data.frame(  
  Model = c("heart.logistic1", "heart.logistic2", "heart.logistic3"),  
  Null_Deviance = c(info1["Null_Deviance"], info2["Null_Deviance"], info3["Null_Deviance"]),  
  Residual_Deviance = c(info1["Residual_Deviance"], info2["Residual_Deviance"], info3["Residual_Deviance"]),  
  R_Squared = c(info1["R_Squared"], info2["R_Squared"], info3["R_Squared"]),  
  AIC = c(info1["AIC"], info2["AIC"], info3["AIC"]),  
  BIC = c(info1["BIC"], info2["BIC"], info3["BIC"])  
)  
(model_info)
```

	Model	Null_Deviance	Residual_Deviance	R_Squared	AIC	BIC
1	heart.logistic1	97040.58	80882.62	0.1665072	80910.62	81038.81
2	heart.logistic2	97040.58	80983.71	0.1654656	81005.71	81106.43
3	heart.logistic3	97040.58	80978.02	0.1655242	80998.02	81089.58

## Final Equation for Logistic Regression Model

[Insert latex equation here](#)

## Training/ Validation

```
set.seed(100)
for(i in 1:10){
  # initialize vector to store prediction errors
  test_errors = numeric(10)
  sample= sample.int(n = nrow(cardio_train), size = floor(0.80*nrow(cardio_train)))
  train.heart.logistic = cardio_train[sample,]
  test.heart.logistic = cardio_train[-sample,]

  train.logistic = glm(cardio ~ age + height + weight + ap_hi + ap_lo +cholesterol + alco +

  glm.pred = predict.glm(train.logistic, newdata = test.heart.logistic, type = "response")

  # Convert probability to binary
  test_predictions_binary = ifelse(glm.pred > 0.5, 1, 0)

  # Calculate test prediction error
  test_error= mean(test_predictions_binary != test.heart.logistic$cardio)

  test_errors[i] = test_error
}

(mean_test_error = mean(test_errors))
```

```
[1] 0.02805
```

Paragraph explaining the the procedure above and the mean error rate

## Results

Insert graphics

Two paragraphs to provide the interpretation of results and your conclusions as it pertains to the original overall question.

## Neural Network