범주형자료분석방법론 HW3

이름: 김연주

학과: 통계학과

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1) 표 5.6

```
> t5.6=expand.grid(AZT=factor(c("Yes","No"),levels=c("No","Yes")),Race=factor(c("Wh
ite","Black"), levels=c("Black", "White")))
> t5.6=data.frame(t5.6,Yes=c(14,32,11,12), No=c(93,81,52,43))
> print(t5.6)
  AZT Race Yes No
1 Yes White 14 93
2 No White 32 81
3 Yes Black 11 52
4 No Black 12 43
> fit1=glm(cbind(Yes,No)~AZT+Race, family=binomial, data=t5.6)
> summary(fit1)
call:
glm(formula = cbind(Yes, No) ~ AZT + Race, family = binomial,
    data = t5.6
Coefficients:
            Estimate Std. Error z value Pr(>|z|)
                      0.26294 -4.083 4.45e-05 ***
0.27898 -2.579 0.00991 **
(Intercept) -1.07357
AZTYes
            -0.71946
                                  0.192 0.84755
RaceWhite
             0.05548
                         0.28861
Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
(Dispersion parameter for binomial family taken to be 1)
    Null deviance: 8.3499 on 3 degrees of freedom
Residual deviance: 1.3835 on 1 degrees of freedom
AIC: 24.86
Number of Fisher Scoring iterations: 4
```

 $logit[\pi(x)] = -1.07357 - 0.71946AZT + 0.05548Race$

모델 요약 결과 AZT 변수의 계수는 유의하지만 Race 변수의 계수는 유의하지 않다.

```
> exp(-0.71946)

[1] 0.4870152

> qchisq(df=1,0.95)

[1] 3.841459
```

Residual deviance가 임계값인 3.84보다 작으므로 모델이 saturated model보다 데이터를 더 잘 적합함을 알 수 있다. AZTyes의 계수를 exp처리한 값은 0.487이다. 이는 에이즈 발병의 오즈가 AZT를 사용했을 때 사용하지 않았을 때보다 0.487배라는 것을 의미한다.

```
> fit2=glm(cbind(Yes,No)~AZT*Race,family=binomial, data=t5.6)
> summary(fit2)
call:
glm(formula = cbind(Yes, No) ~ AZT * Race, family = binomial,
    data = t5.6
Coefficients:
                 Estimate Std. Error z value Pr(>|z|)
                              0.3265 -3.909 9.26e-05 ***
(Intercept)
                  -1.2763
AZTYes
                  -0.2771
                              0.4655 -0.595
                                                0.552
RaceWhite
                   0.3476
                              0.3875 0.897
                                                0.370
AZTYes:RaceWhite -0.6878
                              0.5852 - 1.175
                                                0.240
Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
(Dispersion parameter for binomial family taken to be 1)
    Null deviance: 8.3499e+00 on 3 degrees of freedom
Residual deviance: -1.5987e-14 on 0 degrees of freedom
AIC: 25.476
Number of Fisher Scoring iterations: 3
AZT*Race 변수 추가
> fit3=glm(cbind(Yes,No)~AZT, family=binomial, data=t5.6)
> summary(fit3)
glm(formula = cbind(Yes, No) ~ AZT, family = binomial, data = t5.6)
Coefficients:
            Estimate Std. Error z value Pr(>|z|)
 (Intercept) -1.0361
                         0.1755 -5.904 3.54e-09 ***
             -0.7218
                         0.2787 -2.590 0.00961 **
AZTYes
Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
 (Dispersion parameter for binomial family taken to be 1)
    Null deviance: 8.3499 on 3 degrees of freedom
Residual deviance: 1.4206 on 2 degrees of freedom
AIC: 22.897
Number of Fisher Scoring iterations: 4
```

AZT 변수만 사용

```
> fit4=glm(cbind(Yes,No)~Race, family=binomial, data=t5.6)
> summary(fit4)
call:
glm(formula = cbind(Yes, No) ~ Race, family = binomial, data = t5.6)
Coefficients:
            Estimate Std. Error z value Pr(>|z|)
                        0.23239 -6.103 1.04e-09 ***
 (Intercept) -1.41838
            0.08797
                        0.28547
                                  0.308
RaceWhite
                                           0.758
Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
 (Dispersion parameter for binomial family taken to be 1)
    Null deviance: 8.3499 on 3 degrees of freedom
Residual deviance: 8.2544 on 2 degrees of freedom
AIC: 29.731
Number of Fisher Scoring iterations: 4
Race 변수만 사용
> fit5=glm(cbind(Yes,No)~1, family=binomial, data=t5.6)
> summary(fit5)
glm(formula = cbind(Yes, No) ~ 1, family = binomial, data = t5.6)
Coefficients:
            Estimate Std. Error z value Pr(>|z|)
                         0.1349 -10.08 <2e-16 ***
(Intercept) -1.3606
Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' '1
(Dispersion parameter for binomial family taken to be 1)
    Null deviance: 8.3499 on 3 degrees of freedom
Residual deviance: 8.3499 on 3 degrees of freedom
AIC: 27.826
Number of Fisher Scoring iterations: 4
```

Null model

```
> model=c("AZT*Race","AZT+Race","AZT","Race","null")
> df=c(0,1,2,2,3)
> deviance=c(fit1$deviance,fit2$deviance,fit3$deviance,fit4$deviance,fit5$deviance)
> AIC=c(fit1$aic,fit2$aic,fit3$aic,fit4$aic,fit5$aic)
> compare_model=c("_","2-1","3-2","4-3","5-3")
> compare_dev=c("_",c(fit2$deviance-fit1$deviance, fit3$deviance-fit2$deviance, fit4$deviance-fit3
$deviance, fit5$deviance-fit3$deviance))
> compare_df=c(1,1,1,1,1)
> model_diag=data.frame(model,df,deviance,AIC,compare_model,compare_dev,compare_df)
> print(model_diag)
     model df
                     deviance
                                      AIC compare_model
                                                                 compare_dev compare_df
1 AZT*Race 0 1.383530e+00 24.85981
2 AZT+Race 1 -1.598721e-14 25.47628
                                                      2-1 -1.38353009248638
                                                                                          1
      AZT 2 1.420614e+00 22.89689
Race 2 8.254436e+00 29.73071
null 3 8.349946e+00 27.82622
3
                                                     3-2 1.42061364205088
                                                                                          1
                                                     4-3 6.83382275208821
                                                     5-3 6.92933240995818
                                                                                          1
> #fit3 선택
> residuals(fit3,type='pearson')
> residuals(fit3, type = "pearson")/sqrt(1 - lm.influence(fit3)$hat)
         1
                                   3
-0.7780908 0.8992455 0.7780908 -0.8992455
```

다섯개의 모델을 표로 비교한 결과는 위와 같다. 1,2번 모델을 비교한 결과 compared_dev값이 임계값인 3.84보다 작기에 더 작은 모델인 두번째 모델이 채택된다. 2,3번 모델을 비교하면 같은 이유로 인해 더 작은 모델인 3번 모델이 적합하다. 3,4번 모델을 비교하면 deviance값의 차이가 3.84보다 크기 때문에 3번 모델이 채택된다. 마지막으로 3,5번 모델을 비교하면 같은 이유로 3번 모델이 채택된다. AIC를 비교해 보아도 3번 모델의 AIC가 가장 낮기에 이를 택하는 것이 적합하다.

채택된 모델 (AZT변수만 사용)의 잔차와 표준화잔차를 도출하였다. 그 결과 모델이 잘 적합함을 확인할 수 있었다.

2) 표 6.5, 6.7

```
> #6.5, 6.7
> sp=factor(c("<117","117-126","127-136","137-146","147-156","157-166","167-186",">
186"))
> n=c(156,252,284,271,139,85,99,43)
> obs=c(3,17,12,16,12,8,16,8)
> bp=c(111.5,121.5,131.5,141.5,151.5,161.6,176.5,191.5)
> t6.5=data.frame(sp,bp,n,obs)
> t6.5
            bp n obs
       sp
     <117 111.5 156
1
                     3
2 117-126 121.5 252
                     17
3 127-136 131.5 284
                     12
4 137-146 141.5 271
5 147-156 151.5 139
                     12
6 157-166 161.6 85
                     8
7 167-186 176.5 99 16
     >186 191.5 43
> fit6=glm(obs/n~bp,family=binomial,weights=n,data=t6.5)
> summary(fit6)
call:
glm(formula = obs/n \sim bp, family = binomial, data = t6.5, weights = n)
Coefficients:
             Estimate Std. Error z value Pr(>|z|)
(Intercept) -6.081046
                        0.724230 -8.397 < 2e-16 ***
                                 5.024 5.05e-07 ***
             0.024330
                        0.004842
Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' '1
(Dispersion parameter for binomial family taken to be 1)
    Null deviance: 30.0226 on 7
                                 degrees of freedom
Residual deviance: 5.9134 on 6 degrees of freedom
AIC: 42.615
Number of Fisher Scoring iterations: 4
> qchisq(0.95,df=6)
[1] 12.59159
```

모델 적합 결과 residual deviance가 자유도 6인 카이제곱 분포의 임계값 12.592보다 작으므로 해당 모델이 saturated model보다 데이터를 잘 적합함을 알 수 있다.

```
> #standard residual
> s_res=resid(fit6, type = "pearson")/sqrt(1 - lm.influence(fit6)$hat)
> r1=glm(obs/n~bp,family=binomial,weights=n,data=t6.5[-1,])
> r2=glm(obs/n~bp,family=binomial,weights=n,data=t6.5[-2,])
> r3=glm(obs/n~bp,family=binomial,weights=n,data=t6.5[-3,])
> r4=glm(obs/n~bp,family=binomial,weights=n,data=t6.5[-4,])
> r5=glm(obs/n~bp,family=binomial,weights=n,data=t6.5[-5,])
> r6=qlm(obs/n~bp,family=binomial,weights=n,data=t6.5[-6,])
> r7=glm(obs/n~bp,family=binomial,weights=n,data=t6.5[-7,])
> r8=glm(obs/n~bp,family=binomial,weights=n,data=t6.5[-8,])
> G=c(fit6$deviance-r1$deviance,fit6$deviance-r2$deviance,fit6$deviance-r3$devianc
e,fit6$deviance-r4$deviance,fit6$deviance-r5$deviance,fit6$deviance-r6$deviance,fit
6$deviance-r7$deviance, fit6$deviance-r8$deviance)
[1] 1.38682177 5.13817728 0.93478517 0.33606512 0.01599573 0.11325019 0.42199844
[8] 0.03092674
> #pearson residual
> p_res=resid(fit6, type = "pearson")
> p_diff=c(p_res[1]^2,p_res[2]^2,p_res[3]^2,p_res[4]^2,p_res[5]^2,p_res[6]^2,p_res
[7]^2, p_res[8]^2
> p_diff
                               3
                                                     5
                    2
                                         4
                                                               6
0.95952629 4.02309509 0.66098125 0.25607027 0.01402959 0.09602632 0.26541626
0.01903237
> #standardized residual (table 6.7)
> s_diff=c(s_res[1]^2,s_res[2]^2,s_res[3]^2,s_res[4]^2,s_res[5]^2,s_res[6]^2,s_res
[7]^2, s_res[8]^2
> s_diff
1.22311954 5.63944496 0.89286868 0.32714094 0.01613177 0.11038294 0.42763364
0.03075717
> #df
> df = c((fit6$coefficients[2] - r1$coefficients[2]) / sqrt(vcov(fit6)[2, 2]),(fit6
$coefficients[2] - r2$coefficients[2]) / sqrt(vcov(fit6)[2, 2]),(fit6$coefficients
[2] - r3$coefficients[2]) / sqrt(vcov(fit6)[2, 2]),(fit6$coefficients[2] - r4$coeff
icients[2]) / sqrt(vcov(fit6)[2, 2]),(fit6$coefficients[2] - r5$coefficients[2]) /
sqrt(vcov(fit6)[2, 2]),(fit6$coefficients[2] - r6$coefficients[2]) / sqrt(vcov(fit
6)[2, 2]),(fit6$coefficients[2] - r7$coefficients[2]) / sqrt(vcov(fit6)[2, 2]),(fit
6$coefficients[2] - r8$coefficients[2]) / sqrt(vcov(fit6)[2, 2]))
> df
          bp
                       bp
                                   bp
                                                bp
                                                             bp
 0.484546941 -1.233261336
                          bp
                       bp
 0.412778680 -0.121788108
> #표로 만들어서 비교
> diag2=data.frame(bp,df,p_diff,s_diff,G)
> diag2
                  df
                         p_diff
                                    s_diff
1 111.5 0.484546941 0.95952629 1.22311954 1.38682177
2 121.5 -1.233261336 4.02309509 5.63944496 5.13817728
3 131.5 0.315906335 0.66098125 0.89286868 0.93478517
4 141.5 0.079300520 0.25607027 0.32714094 0.33606512
5 151.5 0.007879018 0.01402959 0.01613177 0.01599573
6 161.6 -0.066291266 0.09602632 0.11038294 0.11325019
7 176.5 0.412778680 0.26541626 0.42763364 0.42199844
8 191.5 -0.121788108 0.01903237 0.03075717 0.03092674
```

표 6.7의 결과를 diag2로 표현하였다. 표에서 두번째 행의 값들이 다른 값들보다 훨씬 큰 것을 알 수 있다. 이를 통해 두번째 관측값이 가장 큰 영향력을 가짐을 확인할 수 있다. 이는 두번째 관찰값이 이상치로 작용할 수 있음을 시사한다.

```
> fit7=glm(obs/n~1, family=binomial, weights=n, data=t6.5)
> summarv(fit7)
glm(formula = obs/n \sim 1, family = binomial, data = t6.5, weights = n)
Coefficients:
            Estimate Std. Error z value Pr(>|z|)
                                          <2e-16 ***
            -2.5987
                         0.1081 - 24.05
Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
(Dispersion parameter for binomial family taken to be 1)
    Null deviance: 30.023 on 7 degrees of freedom
Residual deviance: 30.023 on 7 degrees of freedom
AIC: 64.724
Number of Fisher Scoring iterations: 4
> s_res.ind=resid(fit7, type = "pearson")/sqrt(1 - lm.influence(fit7)$hat)
> r1.ind=glm(obs/n~1,family=binomial,weights=n,data=t6.5[-1,])
> r2.ind=glm(obs/n~1,family=binomial,weights=n,data=t6.5[-2,])
> r3.ind=glm(obs/n~1,family=binomial,weights=n,data=t6.5[-3,])
> r4.ind=glm(obs/n~1,family=binomial,weights=n,data=t6.5[-4,])
> r5.ind=glm(obs/n~1,family=binomial,weights=n,data=t6.5[-5,])
> r6.ind=glm(obs/n~1,family=binomial,weights=n,data=t6.5[-6,])
> r7.ind=glm(obs/n~1,family=binomial,weights=n,data=t6.5[-7,])
> r8.ind=glm(obs/n~1,family=binomial,weights=n,data=t6.5[-8,])
> G.ind=c(fit7$deviance-r1.ind$deviance.fit7$deviance-r2.ind$deviance.fit7$deviance
-r3.ind$deviance,fit7$deviance-r4.ind$deviance,fit7$deviance-r5.ind$deviance,fit7$d
eviance-r6.ind$deviance,fit7$deviance-r7.ind$deviance,fit7$deviance-r8.ind$devianc
e)
> #pearson residual
> p_res.ind=resid(fit7, type = "pearson")
> p_diff.ind=c(p_res.ind[1]^2,p_res.ind[2]^2,p_res.ind[3]^2,p_res.ind[4]^2,p_res.in
d[5]^2,p_res.ind[6]^2,p_res.ind[7]^2,p_res.ind[8]^2)
> #standardized residual (table 6.7)
> s_diff.ind=c(s_res.ind[1]^2,s_res.ind[2]^2,s_res.ind[3]^2,s_res.ind[4]^2,s_res.in
d[5]^2,s_res.ind[6]^2,s_res.ind[7]^2,s_res.ind[8]^2)
> diag2.ind=data.frame(p_diff.ind,s_diff.ind,G.ind)
> diag2.ind
   p_diff.ind s_diff.ind
                                G. ind
1 6.05140873 6.85619967 9.10672745
2 0.01217915 0.01502886 0.01512068
  3.20641469 4.07782308 4.53582651
   0.43624638 0.54798813 0.56812426
5 0.63125065 0.70498497 0.66274271
6 0.81743679 0.87329059 0.79656796
7 13.11561504 14.17126210 10.87035484
8 9.10765714 9.41219000 6.74014336
```

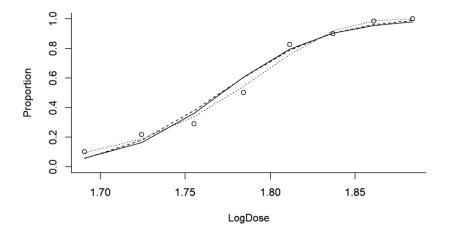
독립성 모델. 관측값을 하나씩 제거했을 때 독립성 모델을 살펴보았다.

```
> #7.1
> t7.1=data.frame(LogDose=c(1.6907,1.7242,1.7552,1.7842,1.8113,1.8369,1.8610,1.883
9), n=c(59,60,62,56,63,59,62,60), y=c(6,13,18,28,52,53,61,60))
> fit8=glm(y/n ~ LogDose, weights=n, family=binomial,data=t7.1)
> summary(fit8)
call:
glm(formula = y/n \sim LogDose, family = binomial, data = t7.1,
    weights = n
Coefficients:
            Estimate Std. Error z value Pr(>|z|)
                                         <2e-16 ***
                          5.181 -11.72
(Intercept) -60.717
LogDose
              34.270
                          2.912
                                  11.77
                                          <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: 284.202 on 7 degrees of freedom
Residual deviance: 11.232 on 6 degrees of freedom
AIC: 41.43
Number of Fisher Scoring iterations: 4
   logit[π(x)] = −60.717 + 34.270LogDose. LogDose의 한 단위 증가는 사건 발생의 오즈를
exp(34.27)배 증가시킨다. 또한 해당 계수는 통계적으로 유의하다.
> fit9=glm(y/n ~ LogDose, weights=n,family=binomial(link=probit),data=t7.1)
> summary(fit9)
call:
glm(formula = y/n \sim LogDose, family = binomial(link = probit),
    data = t7.1, weights = n)
Coefficients:
            Estimate Std. Error z value Pr(>|z|)
(Intercept) -34.935
                          2.648 -13.19 <2e-16 ***
LogDose
              19.728
                          1.487 13.27
                                          <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: 284.20 on 7 degrees of freedom
Residual deviance: 10.12 on 6 degrees of freedom
AIC: 40.318
Number of Fisher Scoring iterations: 4
   Probit link 사용. \varphi^{-1}[\pi(x)] = -34.935 + 19.728 \text{LogDose}.
```

```
> fit10=glm(y/n ~ LogDose, weights=n,family=binomial(link=cloglog),data=t7.1)
> summary(fit10)
call:
glm(formula = y/n \sim LogDose, family = binomial(link = cloglog),
    data = t7.1, weights = n)
Coefficients:
            Estimate Std. Error z value Pr(>|z|)
                                           <2e-16 ***
                           3.240 -12.21
(Intercept)
             -39.572
LogDose
              22.041
                          1.799
                                 12.25
                                           <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: 284.2024 on 7
                                    degrees of freedom
Residual deviance:
                     3.4464 on 6
                                   degrees of freedom
AIC: 33.644
Number of Fisher Scoring iterations: 4
   cloglog link 사용. Log[-log(1-π(x))] =-39.572 + 22.041LogDose.
```

해당 모델은 앞선 fit8, fit9와 달리 $\pi(x)$ 가 0으로는 천천히, 1로는 빠른 속도로 가까워진다. 또한 세모델 중 AIC가 가장 작다.

```
> plot(t7.1$LogDose,t7.1$y/t7.1$n,ylim=c(0,1), xlab="LogDose",ylab="Proportion",bty
="L")
> axis(side=1, at=seq(from=1.5,to=2,by=0.05))
> lines(t7.1$LogDose,predict(fit8, type="response"),lty=1)
> lines(t7.1$LogDose,predict(fit9, type="response"),lty=2)
> lines(t7.1$LogDose,predict(fit10, type="response"),lty=3)
```



세 모델을 그래프를 통해 비교하면 실선은 fit8, 점실선은 fit9, 점선은 fit10이다. 비교 결과 세 모델의 예측값이 비슷함을 알 수 있다.