Solutions for Examination Categorical Data Analysis, February 14, 2020

Problem 1

a. The linear logistic regression model has

$$\pi(x) = \frac{\exp(\alpha + \beta x)}{1 + \exp(\alpha + \beta x)}.$$
(1)

b. We need to find a confidence interval for $\pi(8)$. We first look at logit $[\pi(8)] = \alpha + 8\beta$, whose point estimate is

$$logit [\hat{\pi}(8)] = \hat{\alpha} + 8\hat{\beta} = -6.0 + 8 \cdot 0.5 = -2.0.$$
(2)

Since

$$\operatorname{Var}\left[\operatorname{logit}(\hat{\pi}(8))\right] = \operatorname{Var}(\hat{\alpha}) + 2 \cdot 8 \cdot \operatorname{Cov}(\hat{\alpha}, \hat{\beta}) + 8^2 \cdot \operatorname{Var}(\hat{\beta}) \\ = \operatorname{Var}(\hat{\alpha}) + 16 \cdot \operatorname{Cov}(\hat{\alpha}, \hat{\beta}) + 64 \cdot \operatorname{Var}(\hat{\beta}),$$

this gives a standard error for the estimate in (2) that equals

$$SE = \sqrt{\widehat{Var}(\hat{\alpha}) + 16 \cdot \widehat{Cov}(\hat{\alpha}, \hat{\beta}) + 64 \cdot \widehat{Var}(\hat{\beta})} \\ = \sqrt{0.1 + 16 \cdot (-0.01) + 64 \cdot 0.005} \\ = \sqrt{0.260} \\ = 0.510$$

and a Wald type confidence interval

$$(-2.0 - 1.96 \cdot \text{SE}, -2.0 + 1.96 \cdot \text{SE}) = (-3.000, -1.001)$$

for logit[$\pi(8)$] with approximate coverage probability 95%, since $z_{0.025} = \sqrt{\chi_1^2(0.05)} =$ 1.96 is the 97.5% quantile of a standard normal distribution. The corresponding confidence interval for $\pi(8)$, with approximate coverage probability 95%, is

$$\left(\frac{\exp(-3.000)}{1+\exp(-3.000)}, \frac{\exp(-1.001)}{1+\exp(-1.001)}\right) = (0.048, 0.269).$$

c. Suppose Scott's PSA level is x, so that James' PSA level is x + 3. The requested odds ratio is

$$OR = \frac{\pi(x+3)/(1-\pi(x+3))}{\pi(x)/(1-\pi(x))} = \frac{\exp(\alpha+\beta(x+3))}{\exp(\alpha+\beta x)} = \exp(3\beta)$$

By a similar argument as in a), we first compute a confidence interval

$$\left(\hat{\beta} - 1.96\sqrt{\widehat{\operatorname{Var}}(\hat{\beta})}, \hat{\beta} + 1.96\sqrt{\widehat{\operatorname{Var}}(\hat{\beta})}\right) = (0.5 - 1.96 \cdot \sqrt{0.005}, 0.5 + 1.96 \cdot \sqrt{0.005})$$

= (0.3614, 0.6386)

for β with approximate coverage probability 95%. The corresponding confidence interval for the odds ratio is

 $(\exp(3 \cdot 0.3614), \exp(3 \cdot 0.6386)) = (2.96, 6.79).$

Problem 2

a. Let $X, Y \in \{1, 2\}$ refer to type of surgery and outcome of surgery respectively, for a randomly chosen individual, and $\pi_{ij} = P(X = i, Y = j)$ the probability that an observation belongs to cell (i, j). Since these probabilities are proportional to the expected cell counts μ_{ij} and sum to 1, we have that $\pi_{ij} = \mu_{ij}/\mu_{++}$. Therefore,

$$\pi_1 = P(Y = 1 | X = 1) = \pi_{11}/(\pi_{11} + \pi_{12}) = \mu_{11}/(\mu_{11} + \mu_{12}),$$

$$\pi_2 = P(Y = 1 | X = 2) = \pi_{21}/(\pi_{21} + \pi_{22}) = \mu_{21}/(\mu_{21} + \mu_{22}).$$

b. If one conditions on the two row sums $N_{1+} = n_{1+} = n_1$ and $N_{2+} = n_{2+} = n_2$, we get independent binomial rows sampling. This corresponds to N_{11} and N_{21} being independent binomially distributed random variables

$$N_{11} \sim \text{Bin}(n_1, \pi_1),$$

 $N_{21} \sim \text{Bin}(n_2, \pi_2).$

c. The maximum likelihood estimator of r is

$$\hat{r} = \frac{\hat{\pi}_1}{\hat{\pi}_2} = \frac{N_{11}/n_1}{N_{21}/n_2} = \frac{30/200}{20/100} = 0.75.$$

By means of a first order Taylor expansion, we have that

$$\log(\hat{r}) = \log(\hat{\pi}_1) - \log(\hat{\pi}_2) \approx \log(\pi_1) + \frac{\hat{\pi}_1 - \pi_1}{\pi_1} - \log(\pi_2) - \frac{\hat{\pi}_2 - \pi_2}{\pi_2}.$$

Consequently, since $\hat{\pi}_1$ and $\hat{\pi}_2$ are independent for the sampling scheme in b), the variance of $\log(\hat{r})$ satisfies

$$\operatorname{Var}[\log(\hat{r})] \approx \operatorname{Var}\left(\frac{\hat{\pi}_{1}-\pi_{1}}{\pi_{1}}\right) + \operatorname{Var}\left(\frac{\hat{\pi}_{2}-\pi_{2}}{\pi_{2}}\right) \\ = \frac{\operatorname{Var}(\hat{\pi}_{1})}{\pi_{1}^{2}} + \frac{\operatorname{Var}(\hat{\pi}_{2})}{\pi_{2}^{2}} \\ = \frac{\operatorname{Var}(N_{11})}{n_{1}^{2}\pi_{1}^{2}} + \frac{\operatorname{Var}(N_{21})}{n_{2}^{2}\pi_{2}^{2}} \\ = \frac{n_{1}\pi_{1}(1-\pi_{1})}{n_{1}^{2}\pi_{1}^{2}} + \frac{n_{2}\pi_{2}(1-\pi_{2})}{n_{2}^{2}\pi_{2}^{2}} \\ = \frac{\pi_{1}(1-\pi_{1})}{n_{1}\pi_{1}^{2}} + \frac{\pi_{2}(1-\pi_{2})}{n_{2}\pi_{2}^{2}} \\ = \frac{1-\pi_{1}}{n_{1}\pi_{1}} + \frac{1-\pi_{2}}{n_{2}\pi_{2}}.$$
(3)

d. The standard error of $\log(\hat{r})$ is obtained by plugging in estimates of π_1 and π_2 into the variance formula (3) and then taking the square root. That is,

$$\begin{aligned} \operatorname{SE}[\log(\hat{r})] &= \sqrt{\widehat{\operatorname{Var}}[\log(\hat{r})]} \\ &= \sqrt{\frac{1-\hat{\pi}_1}{n_1\hat{\pi}_1} + \frac{1-\hat{\pi}_2}{n_2\hat{\pi}_2}} \\ &= \sqrt{\frac{n_{12}}{n_1n_{11}} + \frac{n_{22}}{n_2n_{21}}} \\ &= \sqrt{\frac{170}{200\cdot30} + \frac{80}{100\cdot20}} \\ &= \sqrt{0.0683} \\ &= 0.2614. \end{aligned}$$

This gives a Wald-based confidence interval

$$(\log(\hat{r}) - 1.96 \cdot SE[\log(\hat{r})], \log(\hat{r}) + 1.96 \cdot SE[\log(\hat{r})]) = (\log(0.75) - 1.96 \cdot 0.2614, \log(0.75) + 1.96 \cdot 0.2614) = (-0.800, 0.225)$$

for $\log(r)$ with approximate coverage probability 95%. Applying the exponential transformation to both sides of this interval, we finally obtain a confidence interval

$$(\exp(-0.800), \exp(0.225)) = (0.45, 1.25)$$

for r with approximate coverage probability 95%. Since 1 belongs to this interval, the null hypothesis H_0 that type of surgery does not influence the probability of outcome (i.e. $\pi_1 = \pi_2$ or r = 1) is not rejected.

Problem 3

a. For the loglinear model (AC, AM, CM) we have that

$$\mu_{acm} = \exp(\lambda + \lambda_a^A + \lambda_c^C + \lambda_m^M + \lambda_{ac}^{AC} + \lambda_{am}^{AM} + \lambda_{cm}^{CM}),$$

for all cells $a, c, m \in \{0, 1\}$. If a = c = m = 0 are chosen as baseline levels, then all loglinear parameters equal 0 if at least index is 0. This gives a parameter vector with the remaining nonzero loglinear parameters

$$\boldsymbol{\beta} = (\lambda, \lambda_1^A, \lambda_1^C, \lambda_1^M, \lambda_{11}^{AC}, \lambda_{11}^{AM}, \lambda_{11}^{CM}).$$

The number of parameters is thus p(AC, AM, CM) = 7.

b. All of the listed models in the tables are balanced, and all three categorical variables are binary. Therefore, each model has 1 baseline parameter, 3 main effect parameters (1 per main effect), $1 = (2-1) \cdot (2-1)$ parameter per second order association, and $1 = (2-1) \cdot (2-1) \cdot (2-1)$ parameter per third order association. Adding the number of baseline, main effect, second order, and third order association parameters, we

find the total number of parameters

$$p(A, C, M) = 1 + 3 + 0 + 0 = 4,$$

$$p(A, CM) = 1 + 3 + 1 + 0 = 5,$$

$$p(C, AM) = 1 + 3 + 1 + 0 = 5,$$

$$p(M, AC) = 1 + 3 + 1 + 0 = 5,$$

$$p(AC, AM) = 1 + 3 + 2 + 0 = 6,$$

$$p(AC, CM) = 1 + 3 + 2 + 0 = 6,$$

$$p(AM, CM) = 1 + 3 + 2 + 0 = 6,$$

$$p(AC, AM, CM) = 1 + 3 + 3 + 0 = 7,$$

$$p(ACM) = 1 + 3 + 3 + 1 = 8$$

of all models.

c. Since (ACM) is the saturated model, Akaike's Information Criterion of each model is

$$AIC(Model) = -2L(Model) + 2p(Model)$$

= $-2[L(Model) - L(ACM)] + 2p(Model) - 2L(ACM)$
= $G^{2}(Model) + 2p(Model) - 2L(ACM),$

where L(Model) and $G^2(Model)$ are the log likelihood and deviance of each model respectively. We select the best model, according to the AIC-criterion, by minimizing AIC(Model), which is equivalent to minimizing $G^2(Model) + 2p(Model)$. We found the number of parameters p(Model) of all models in b). This makes it possible to fill in the second column of the given table, and then add a third column:

Model	G^2	p	$G^2 + 2p$
(A,C,M)	1286.0	4	1294.0
(A,CM)	534.2	5	544.2
(C,AM)	939.6	5	949.6
(M,AC)	843.8	5	853.8
(AC,AM)	497.4	6	509.4
(AC,CM)	92.0	6	104.0
(AM,CM)	187.8	6	199.8
(AC,AM,CM)	0.4	7	14.4
(ACM)	0.0	8	16

Since (AC, AM, CM) minimizes $G^2(Model) + 2p(Model)$, this is the model chosen by the AIC-criterion.

d. In the first step of backward elimination (BE), the largest model among those listed in the table, (ACM), is tested against the model (AC, AM, CM) for which the third order association between the three variables has been removed, by means of a likelihood ratio test. This gives

$$G^{2}(AC, AM, CM | ACM) = -2[L(AC, AM, CM) - L(ACM)]$$

= $G^{2}(AC, AM, CM) - G^{2}(ACM)$
= $0.4 - 0$
 $< \chi^{2}_{8-7}(0.05) = 3.84,$ (4)

where df = p(ACM) - p(AC, AM, CM) = 8 - 7 = 1 is used for the quantile of the χ^2 distribution. Since the deviance does not exceed this quantile, the null hypothesis (AC, AM, CM) is not rejected, and the smaller model is selected in favor of the alternative hypothesis that (ACM) holds but not (AC, AM, CM).

In the second step of the BE scheme we test (AC, AM, CM) against each one of the three models obtained by removing one second order association from (AC, AM, CM). The log likelihood ratios of these three tests are found, as in (4), from the difference in deviance:

$$\begin{array}{rcl} G^2(AC,AM|AC,AM,CM) &=& 497.4-0.4=497.0,\\ G^2(AC,CM|AC,AM,CM) &=& 92.0-0.4=91.6,\\ G^2(AM,CM|AC,AM,CM) &=& 187.8-0.4=187.4, \end{array}$$

Since all these three deviances (by a very large margin) exceed $\chi^2_{7-6}(0.05) = 3.84$, the null hypothesis (the smaller model) is rejected in each test. Therefore, the BE scheme stops after this second step and (AC, AM, CM) is selected, the same model that was chosen with the AIC criterion in c).

Problem 4

a. Let $\pi_{acm} = \mu_{acm}/\mu_{+++}$ be the probability of cell (a, c, m) for multinomial sampling when we condition on the total number of observations of the Poisson model (AC, AM, CM). Regarding M as the outcome variable and A, C as predictor variables of this multinomial model, we find that M|A, C is an ANOVA type logistic regression model, since

$$\begin{aligned} \log it P(M = 1 | A = a, C = c) \\ &= \log[P(M = 1 | A = a, C = c) / P(M = 0 | A = a, C = c)] \\ &= \log[(\pi_{ac1}/\pi_{ac+})/(\pi_{ac0}/\pi_{ac+})] \\ &= \log((\pi_{ac1}/\pi_{ac0})) \\ &= \log((\mu_{ac1}/\mu_{ac0})) \\ &= \log((\mu_{ac1}) - \log((\mu_{ac0}))) \\ &= \lambda + \lambda_a^A + \lambda_c^C + \lambda_1^M + \lambda_{ac}^{AC} + \lambda_{a1}^{AM} + \lambda_{c1}^{CM}) \\ &- (\lambda + \lambda_a^A + \lambda_c^C + \lambda_0^M + \lambda_{ac}^{AC} + \lambda_{a0}^{AM} + \lambda_{c0}^{CM}) \\ &= \alpha + \beta_a^A + \beta_c^C, \end{aligned}$$
(5)

with

$$\begin{aligned} \alpha &= \lambda_1^M - \lambda_0^M = \lambda_1^M, \\ \beta_a^A &= \lambda_{a1}^{AM} - \lambda_{a0}^{AM} = \lambda_{a1}^{AM}, \\ \beta_c^C &= \lambda_{c1}^{CM} - \lambda_{c0}^{CM} = \lambda_{c1}^{CM}. \end{aligned}$$

In the last step we assumed that a = c = m = 0 are baseline levels, putting to zero all loglinear parameters with at least one 0 index. Then all effect parameters $\beta_0^A = \beta_0^C = 0$ vanish, and the remaining three nonzero parameters of the logistic regression model, are $\boldsymbol{\beta} = (\alpha, \beta_1^M, \beta_1^C)$.

b. The conditional odds ratio of marijuana use between those that use alcohol and those that do not, conditional on cigarette use, is

$$\theta_{AM(c)} = \frac{P(M=1|A=1, C=c)/P(M=0|A=1, C=c)}{P(M=1|A=0, C=c)/P(M=0|A=0, C=c)}.$$
(6)

It follows from (5) that

$$\log \theta_{AM(c)} = \log it P(M = 1 | A = 1, C = c) - \log it P(M = 1 | A = 0, C = c) \\ = \alpha + \beta_1^A + \beta_c^C - (\alpha + \beta_0^A + \beta_c^C) \\ = \beta_1^A - \beta_0^A \\ = \beta_1^A \\ = \lambda_{11}^{AM}$$

when a = m = 0 are chosen as baseline levels of alcohol and marijuana use. Equivalently,

$$\theta_{AM(c)} = \exp(\lambda_{11}^{AM}). \tag{7}$$

- c. There is homogeneous association between alcohol use A and marijuana use M if the conditional odds ratio $\theta_{AM(c)}$ does not depend on the level c of smoking. It follows from (7) that model (AC, AM, CM) has homogeneous association, since the right hand side of this equation does not depend on c. Similarly, one shows that all loglinear models for which A and M are not involved in the same third order association, have homogeneous association between A and M. Hence, among the loglinear models listed in the table of Problem 3, all models except the saturated model (ACM) have homogeneous association between A and M.
- d. Since M and C are conditionally independent given A for model (AM, AC), it follows that P(M|A, C) = P(M|A). Inserting this relation into (6), we find that the conditional odds ratio

$$\theta_{AM(c)} = \frac{P(M=1|A=1)/P(M=0|A=1)}{P(M=1|A=0)/P(M=0|A=0)} = \theta_{AM}$$
(8)

for model (AM, AC) equals the marginal odds ratio θ_{AM} , by the definition of the latter. We estimate the marginal odds ratio from the marginal twoway table of A, M, by replacing each P(M = m | A = a) in (8) with $\hat{P}(M = m | A = a) = n_{a+m}/n_{a++}$. Since all n_{0++} and n_{1++} cancel out, it follows that

$$\hat{\theta}_{AM} = \frac{n_{0+0}n_{1+1}}{n_{0+1}n_{1+0}} = \frac{(43+279)\cdot(911+44)}{(3+2)\cdot(538+456)} = 61.9.$$

This is very different from the estimate $\hat{\theta}_{AM(c)} = 19.8$ of the conditional odds ratio between A and M for model (AC, AM, CM). However, we know from Problems 3c) and 3d) that (AM, AC) fits data much worse than (AC, AM, CM). For this reason the estimated (conditional) odds ratio of model (AC, AM, CM) is more trustworthy. Therefore, the odds of using marijuana is about 20 times higher for those that use alcohol, compared to those that don't.

Problem 5

a. This is a loglinear model with t_i as an offset. Let $\boldsymbol{\lambda} = (\lambda_0, \lambda_1)^T$ be the parameter vector. Then the likelihood function is

$$l(\boldsymbol{\lambda}) = \prod_{i=0}^{3} e^{-\mu_i} \frac{\mu_i^{y_i}}{y_i!},$$

and the log likelihood

$$L(\boldsymbol{\lambda}) = \log l(\boldsymbol{\lambda}) = \sum_{i=0}^{3} [y_i \log(\mu_i) - \mu_i - \log(y_i!)] = \operatorname{constant} + \sum_{i=0}^{3} [y_i(\lambda_0 + \lambda_1 i) - t_i \exp(\lambda_0 + \lambda_1 i)],$$
(9)

where

constant =
$$\sum_{i=0}^{3} \left[y_i \log(t_i) - \log(y_i!) \right]$$

does not depend on the two parameters λ_0 and λ_1 .

b. Since

$$\mu_i = t_i \exp(\lambda_0 + \lambda_1 i), \tag{10}$$

we find that

$$\frac{d\mu_i}{d\boldsymbol{\lambda}} = \begin{pmatrix} \partial u_i / \partial \lambda_0 \\ \partial u_i / \partial \lambda_1 \end{pmatrix} = \mu_i \begin{pmatrix} 1 \\ i \end{pmatrix}.$$

From this and (9) it follows that the likelihood score vector equals

$$\boldsymbol{u}(\boldsymbol{\lambda}) = \begin{pmatrix} \frac{\partial L(\boldsymbol{\lambda})}{\partial \lambda_0} \\ \frac{\partial L(\boldsymbol{\lambda})}{\partial \lambda_1} \end{pmatrix} = \sum_{i=0}^{3} (y_i - \mu_i) \begin{pmatrix} 1 \\ i \end{pmatrix}.$$
(11)

The likelihood equations are obtained by solving

$$oldsymbol{u}(oldsymbol{\lambda})_{oldsymbol{\lambda}=(\hat{\lambda}_0,\hat{\lambda}_1)}=\left(egin{array}{c} 0 \ 0 \end{array}
ight)$$

with respect to $\hat{\lambda}_0$ and $\hat{\lambda}_1$, which is equivalent to solving

$$\sum_{i=0}^{3} y_i \begin{pmatrix} 1\\i \end{pmatrix} = \sum_{i=0}^{3} t_i \exp(\hat{\lambda}_0 + \hat{\lambda}_1 i) \begin{pmatrix} 1\\i \end{pmatrix}.$$

c. We first find the Hessian matrix

$$\boldsymbol{H}(\boldsymbol{\lambda}) = \frac{d^2 L(\boldsymbol{\lambda})}{d^2 \boldsymbol{\lambda}} = \begin{pmatrix} \frac{\partial^2 L(\boldsymbol{\lambda})}{\partial^2 \lambda_0} & \frac{\partial^2 L(\boldsymbol{\lambda})}{\partial^2 \lambda_0} \\ \frac{\partial^2 L(\boldsymbol{\lambda})}{\partial^2 \lambda_0} & \frac{\partial^2 L(\boldsymbol{\lambda})}{\partial^2 \lambda_1} \end{pmatrix}$$

of the log likelihood by differentiating (11) with respect to λ_0 and λ_1 . This gives

$$oldsymbol{H}(oldsymbol{\lambda}) = -\sum_{i=0}^{3} \mu_i \left(egin{array}{cc} 1 & i \ i & i^2 \end{array}
ight).$$

Since $H(\lambda)$ does not depend on data it is non-stochastic. Therefore the Fisher information matrix equals

$$\boldsymbol{J}(\boldsymbol{\lambda}) = -E\left[\boldsymbol{H}(\boldsymbol{\lambda})\right] = -\boldsymbol{H}(\boldsymbol{\lambda}) = \sum_{i=0}^{3} \mu_i \left(\begin{array}{cc} 1 & i\\ i & i^2 \end{array}\right).$$
(12)

d. The covariance matrix $\text{Cov}(\hat{\lambda})$ of the parameter vector is approximately $J(\lambda)^{-1}$. It is estimated by

$$\widehat{\text{Cov}}(\hat{\boldsymbol{\lambda}}) = \hat{\boldsymbol{J}}^{-1} = \begin{pmatrix} \hat{J}_{00}^{(-1)} & \hat{J}_{01}^{(-1)} \\ \hat{J}_{10}^{(-1)} & \hat{J}_{11}^{(-1)} \end{pmatrix},$$

where

$$\hat{J} = J(\hat{\boldsymbol{\lambda}}) = -\boldsymbol{H}(\hat{\boldsymbol{\lambda}}) = \sum_{i=0}^{3} \hat{\mu}_i \begin{pmatrix} 1 & i \\ i & i^2 \end{pmatrix}.$$

is the observed (expected) Fisher information matrix, and $\hat{\mu}_i = t_i \exp(\hat{\lambda}_0 + \hat{\lambda}_1 i)$ the estimated expected number of heart attacks within each group *i* of patients. The one-sided Wald test, with approximate significance level α , rejects H_0 in favor of H_a when

$$z_W = \frac{\lambda_1}{\sqrt{\hat{J}_{11}^{(-1)}}}$$

exceeds the $(1 - \alpha)$ -quantile z_{α} of a standard normal distribution.