## Biostats 653 - Final

NAME	 	 	

Date: December 15, 2016 Instructor: Xiang Zhou

Time: 120 minutes (4:00pm – 6:00pm)

Note that all sub-questions of Question 1 can be answered in a sentence or two. For Questions 2 and 3, you can directly use formulas derived in class. Use the blank pages attached at the back if needed. Try not to leave empty space even if you do not know the answer.

## Question 1 (40 pt)

A study was conducted to investigate the effects of an antidepressant drug used to treat individuals who suffer from debilitating panic attacks. Panic attacks are temporary periods (on the order of 10 to 15 minutes) of intense fear and distress that can terrify the sufferer and interfere with his or her day-to-day life. A total of m=300 subjects confirmed to suffer from such attacks were recruited and randomized to three groups:

Group 1 – low-dose antidepressant therapy (100 subjects)

Group 2 - high-dose antidepressant therapy (100 subjects)

Group 3 – placebo (100 subjects)

Before starting his/her assigned treatment, each subject was asked whether or not she/he had suffered at least one panic attack in the previous week (0=no, 1=yes). This was taken as the subject's baseline response (week 0). All subjects then started on their assigned therapies. At 1, 2, 3, and 4 weeks thereafter, each subject visited the clinic and was asked to report whether or not she/he had suffered at least one attach since the last visit (0=no, 1=yes). The gender of each subject (0=female, 1=male) was also recorded.

The following table shows the proportions of subjects reporting suffering at least one attach in the previous week at each time point:

Group	Baseline(week 0)	Week 1	Week 2	Week 3	Week 4	
1 (low dose)	0.70	0.61	0.65	0.59	0.54	
2 (high dose)	0.60	0.57	0.54	0.51	0.39	
3 (placebo)	0.67	0.59	0.64	0.66	0.66	

Let  $Y_{ij}$  be the indicator of whether or not subject i reported at least one panic attack,  $i=1,\cdots,300$ , in week  $t_{ij}=0,1,2,3,4$ , and let

 $\delta_{1i}=1$  if subject i was in Group 1 (low dose), = 0 otherwise

 $\delta_{2i}=1$  if subject i was in Group 2 (high dose), = 0 otherwise

 $\delta_{3i}=1$  if subject i was in Group 3 (placebo), = 0 otherwise

The study team first considered the following marginal model for the probability of at least one panic attack:

$$E(Y_{ij}) = \frac{\exp(\beta_0 + \beta_1 t_{ij} \delta_{1i} + \beta_2 t_{ij} \delta_{2i} + \beta_3 t_{ij} \delta_{3i})}{1 + \exp(\beta_0 + \beta_1 t_{ij} \delta_{1i} + \beta_2 t_{ij} \delta_{2i} + \beta_3 t_{ij} \delta_{3i})}$$

1). (5 pt) In words, state what the above model assumes about the pattern of change of the logarithm of the odds of having at least one panic attack in each group, with respect to time.
2). (5 pt) The first question the study team wished to address was whether or not the pattern of change of the log odds of having at least one panic attack once treatment is initiated is possibly different for at least one of the three groups. State the null hypothesis and explain how you would test this null in the marginal model.
3). (5 pt) To complete the model specification, the study team needs to make an assumption on the marginal variance $V(Y_i)$ (an n by n matrix). What assumptions do you want to make on $V(Y_i)$ ?
4). (5 pt) During the study, some subjects failed to come in week 4. The study team found out that whether a subject came in week 4 only depends on his/her baseline measurement and does not depend on the $\beta$ 's in the model (i.e. a patient tends not to come if he/she has at least one attack in the week before week 0). What is the missing data mechanism?

5). (5 pt) Under the missing data mechanism of 1.4, can we ignore the missing data and use only the observed data to fit the marginal model (i.e. is the missingness ignorable)?
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6). (5 pt) The study team then considered a conditional model with a random intercept, with the conditional mean assumed as:
$E(Y_{ij} b_i) = \frac{\exp(\beta_0 + \beta_1 t_{ij}\delta_{1i} + \beta_2 t_{ij}\delta_{2i} + \beta_3 t_{ij}\delta_{3i} + b_i)}{1 + \exp(\beta_0 + \beta_1 t_{ij}\delta_{1i} + \beta_2 t_{ij}\delta_{2i} + \beta_3 t_{ij}\delta_{3i} + b_i)}$
What is the interpretation of $eta_1$ ? Is the interpretation different from the $eta_1$ in the above marginal model?
7). (5 pt) Generally, what are the advantage(s) and disadvantage(s) of a conditional model compared with a marginal model?
8). (5 pt) Under the same missing data mechanism of 1.4, can we ignore the missing data and use only the observed data to fit the conditional model (i.e. is the missingness ignorable)?

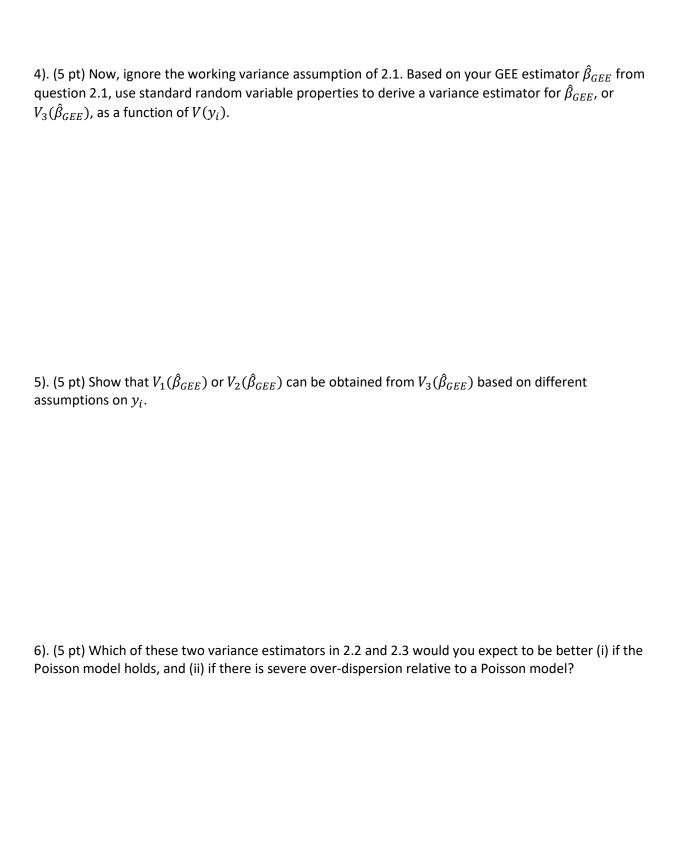
## Question 2 (30 pt)

Assume we have observed longitudinal data  $y_{ij}$ ,  $i=1,\cdots,N,j=1,\cdots,n_i$ . For all sub-questions in Questions 2 and 3, we will consider a special case when the number of repeated measurement is 1, or  $n_i=1$ , for all I (i.e. no repeated measurements). We define  $y_i\equiv y_{i1}$ . Now, suppose we have count data  $y_1,\cdots,y_N$  which we assume are independently distributed with mean  $\beta$ , or  $E(y_i)\equiv \mu_i=\beta$  (i.e. identity link with constant mean).

1). (5 pt) Suppose we fit this model under a working variance assumption that  $V(y_i) = \mu_i = \beta$ , as is the case for Poisson data. Obtain the GEE estimator  $\hat{\beta}_{GEE}$ .

2). (5 pt) Under the assumption of 2.1, obtain the model-based variance estimator for  $\hat{\beta}_{GEE}$ , or  $V_1(\hat{\beta}_{GEE})$ .

3). (5 pt) Under the assumption of 2.1, obtain the robust variance estimator for  $\hat{\beta}_{GEE}$ , or  $V_2(\hat{\beta}_{GEE})$ .



## Question 3 (30 pt)

1). (5 pt) Settings are identical to Question 2. Now suppose you decide to use a generalized linear mixed model (GLMM) with mean  $\mu$  and a random intercept  $b_i$  to model  $y_i$ . Specifically, you assume that  $y_i$ , conditional on  $\beta$ ,  $b_i$ , follows a Poisson distribution, with the conditional mean  $\log \left( E(y_i|b_i) \right) = \beta + b_i$ . You further assume that  $b_i$  follows a normal distribution with mean 0 and variance  $\sigma^2$ . Write down your likelihood. (Note that the probability mass function for a Poisson random variable x is  $P(x;\lambda) = \frac{\lambda^x e^{-\lambda}}{x!}$ .)

2). (10 pt) Compute the marginal mean  $E(y_i)$  and the marginal variance  $V(y_i)$  of the GLMM. Show that  $V(y_i) > E(y_i)$ ; that is, GLMM naturally accounts for over-dispersion.

3). (5 pt) Explain how you would estimate  $\beta$  and its standard error in the above GLMM.

4). (5 pt) In class, we have focused on four different methods to deal with the intractable integration in GLMM. Here, you will develop an alternative approach for inference in GLMM. You notice that the integral of GLMM is often intractable because we assume that the random effects  $b_i$  follow a normal distribution. You reason that, instead of normal, perhaps a different assumption on the distribution of the random effects can sometimes lead to a tractable integral. To see this, you decide to assume that the exponential of the random effects, which are now positive, follow a gamma distribution, or  $a_i \equiv e^{b_i} \sim Gamma(\alpha, \theta)$ , with  $\alpha$ ,  $\theta$  known. With this assumption, re-write the likelihood function in terms of the new random effects  $a_i$ . (Note that the probability density function for a gamma distribution is

$$P(x|\alpha,\theta) = \frac{\theta^{\alpha}x^{\alpha-1}e^{-x\theta}}{\Gamma(\alpha)}.$$

5). (5 pt) Inside the integral, you look that the part that involves  $a_i$  and you notice that this part looks very familiar. In fact, you notice that this part belongs to some distribution (i.e. is the kernel of that distribution). What distribution is this? Based on this observation, and the fact that any distribution integrates to one, solve the integration in GLMM analytically.