# UNIVERSITY OF OSLO

# Faculty of mathematics and natural sciences

Exam in:	STK9080 — Survival and event history analysis		
Day of examination:	Thursday 10 June 2021		
Examination hours:	09.00-13.00		
This problem set consists of 5 pages.			
Appendices:	None		
Permitted aids:	Approved calculator		

Please make sure that your copy of the problem set is complete before you attempt to answer anything.

## Problem 1 Hospital length of stay of COVID-19 patients

In the beginning of the corona pandemic, a group of Chinese medical researchers used data from confirmed COVID-19 patients at hospitals in the Sichuan Province. They collected information on demographic, epidemiological, clinical characteristics, and the length of hospital stay for these patients. The study included 538 patients who were admitted in hospitals between January 16 and March 4, 2020. 332 out of these (62%) recovered and were discharged before the end of the study, April 4, 2020. Only 3 patients died in hospital before April 4.

The data used in this Problem are *simulated* based on the reported results from the study (the full underlying data were not reported in the study). They consist of the observed time, **Time** (in days); censoring status D (= 0 or 1); and six binary covariates  $x_1, \ldots, x_6$  (with values 0 and 1 as defined by Table 1 and explained below).

Here, for patients discharged at or before April 4, Time is the *true length* of hospital stay. These patients are given censoring status D = 1. For patients that were alive and still in hospital on April 4, Time is the *observed length* of stay. These patients are given censoring status D = 0. For patients who died during hospital stay, Time is the time from hospital admission to death. These patients are also given censoring status D = 0.

In Table 1, 'time from onset' means time from onset of COVID-19 to admission at the hospital; 'hospital grade' distinguishes between admission to a provincial (rural) and non-provincial (urban) hospitals; 'density of health workers' means number of health workers per 1000 inhabitants; 'clinical grade' is degree of illness.

i	$x_i$	0	1
1	age (years)	< 45	$\geq 45$
2	$\operatorname{gender}$	male	female
3	time from onset	< 5	$\geq 5$
4	hospital grade	non-provincial	provincial
5	density of health workers	< 5.5	$\geq 5.5$
6	clinical grade	mild	severe

Table 1: List of covariates for the length of stay data

The following is an edited output of a Cox-regression in R, using data for all the 538 patients and including all the covariates  $x_1, \ldots, x_6$ .

```
Call:
coxph(formula = Surv(Time, D == 1) ~ x1 + x2 + x3 + x4 + x5 + x6)
 n= 538, number of events= 332
        coef
               se(coef)
                             z \Pr(|z|)
x1 -0.438713
              0.120904 -3.629 0.000285
x2
   0.089060
               0.112596 0.791 0.428963
xЗ
   0.004456
               0.115056 0.039 0.969108
x4 -0.250050
               0.126474 -1.977 0.048032
x5 0.300428
               0.112477 2.671 0.007562
x6 -0.464376
               0.157129 -2.955 0.003123
```

a) Write down the estimated relative risk function for discharge from hospital. Which covariates have a significant effect? (Use significance level 5%).

Compute and give a practical interpretation of the estimated hazard ratios for the covariates for age and clinical grade. Derive an approximate 95% confidence interval for the hazard ratio for age.

What is the estimated relative risk of a patient of age  $\geq 45$  with severe clinical grade compared to a patient of age < 45 with mild clinical grade, when the other covariates are the same for the two?

The covariates for *age* and *clinical grade*, i.e.,  $x_1$  and  $x_6$ , were of particular interest in the study. In the rest of the Problem we therefore disregard the covariates  $x_2, \ldots, x_5$ .

- **b)** A couple of model checks for the application of Cox regression were made for this setting:
  - 1. It was checked whether there is an interaction between the covariates  $x_1$  and  $x_6$ . Discuss briefly how such an interaction should be interpreted. Then explain how you would do a formal check of this interaction. No calculations are asked for here. (No significant interaction was found in the study).
  - 2. It was checked whether the proportional hazards assumption is appropriate for the model with the two covariates  $x_1$  and  $x_6$ . Explain how this could be done in the present case.
- c) An alternative analysis of the data uses *Aalen's additive hazards model*. State this model when  $x_1$  and  $x_6$  are the only covariates.

Figure 1 shows plots from fitting the model in R.

Explain how the plots should be interpreted. What is in particular the interpretation of the Intercept plot to the left in Figure 1?

(Continued on page 3.)

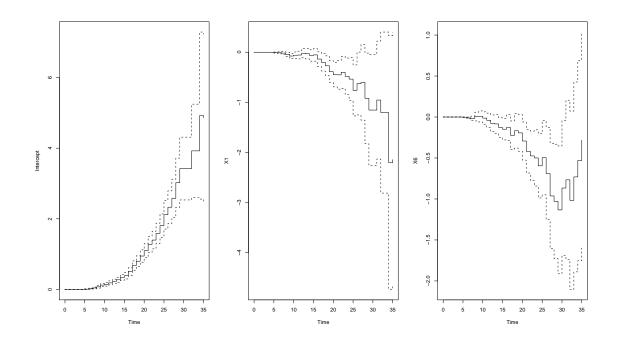


Figure 1: Plots from fitting Aalen's model to the data with covariates  $x_1$  and  $x_6$ .

#### Problem 2 The Gehan-Breslow test

Consider two counting processes  $N_1(t)$  and  $N_2(t)$  with intensity processes of the multiplicative form

$$\lambda_h(t) = Y_h(t)\alpha_h(t); \quad h = 1, 2$$

Here, the  $Y_h(t)$  are predictable processes, while the  $\alpha_h(t)$  are deterministic positive functions. It is assumed that the two processes do not make simultaneous jumps.

We want to test the null hypothesis

$$H_0: \ \alpha_1(t) = \alpha_2(t) \ \text{ for } 0 \le t \le t_0$$

where  $t_0$  is the upper time limit of the study. If  $H_0$  holds, then the common (but unknown) version of the  $\alpha_h(t)$  will be called  $\alpha(t)$ . A subscript • means the sum over 1 and 2.

The test statistic of the Gehan-Breslow test for  $H_0$  is

$$Z(t_0) = \int_0^{t_0} Y_1(s) Y_2(s) \left( d\hat{A}_1(s) - d\hat{A}_2(s) \right)$$
(1)

Here  $\hat{A}_h(t)$  is the Nelson-Aalen estimator for  $A_h(t) = \int_0^t \alpha_h(s) ds$  (h = 1, 2).

a) Looking at equation (1) and comparing to the corresponding expression for the *log-rank* test, how would you describe the practical difference between the two tests? (You should be brief here).

Show that the test statistic  $Z(t_0)$  can be written

$$Z(t_0) = \int_0^{t_0} Y_2(s) \, dN_1(s) - \int_0^{t_0} Y_1(s) \, dN_2(s)$$

(Continued on page 4.)

b) Show that under  $H_0$  we have

$$Z(t_0) = \int_0^{t_0} Y_2(s) \, dM_1(s) - \int_0^{t_0} Y_1(s) \, dM_2(s)$$

where  $M_1(t)$  and  $M_2(t)$  are mean zero martingales.

How are the martingales  $M_1(t)$  and  $M_2(t)$  defined here?

c) Assume that  $H_0$  holds.

Show that Z(t) is a mean zero martingale (as a function of t) with predictable variation

$$\langle Z \rangle (t) = \int_0^t Y_1(s) Y_2(s) Y_{\bullet}(s) \alpha(s) ds$$
<sup>(2)</sup>

(*Hint:* You may use without proof that for counting process martingales  $M_1, M_2$  we have  $\langle M_1 + M_2 \rangle (t) = \langle M_1 \rangle (t) + \langle M_2 \rangle (t)$ ).

d) Assume that the counting processes  $N_1(t)$  and  $N_2(t)$  are obtained by aggregating, respectively,  $n_1$  and  $n_2$  individual processes. Let  $n = n_1 + n_2$  and assume that there are deterministic functions  $y_h(t)$  such that  $Y_h(t)/n \xrightarrow{P} y_h(t)$ , h = 1, 2.

Show that  $Z(t)n^{-3/2}$  under  $H_0$  converges in distribution as  $n \to \infty$  to a mean zero Gaussian martingale with predictable variation process

$$\int_0^t y_1(s) y_2(s) y_{\bullet}(s) \alpha(s) ds$$

(*Hint:* See Section 3.3.5 in Aalen, Borgan & Gjessing).

e) Use equation (2) to derive an unbiased estimator  $V(t_0)$  for  $Var(Z(t_0))$  under  $H_0$ . (*Hint:* You may use that  $N_{\bullet}(t) = \int_0^t Y_{\bullet}(s)\alpha(s)ds + M_{\bullet}(t)$  under  $H_0$ .) Verify that the following test statistic for large n is approximately chi-square distributed with 1 degree of freedom under  $H_0$ :

$$X^{2}(t_{0}) = \frac{(Z(t_{0}))^{2}}{V(t_{0})}$$

### Problem 3 Poisson processes with unobserved heterogeneity

Suppose that n independent homogeneous Poisson processes

$$N_1(t),\ldots,N_n(t)$$

are under study, where the *i*th process,  $N_i(t)$ , has constant rate  $\theta_i > 0$  and is observed in the time interval  $[0, \tau_i]$  for fixed  $\tau_i > 0$  (i = 1, 2, ..., n).

a) What is in general meant by the *intensity* of a counting process N(t)? Explain that the intensity of the process  $N_i(t)$  is

$$\lambda_i(t) = I(t \le \tau_i)\theta_i; \quad t \ge 0, \quad i = 1, \dots, n$$

Then use the general expression for a parametric likelihood function (as derived in the course) to explain that the likelihood contribution for the the process  $N_i(t)$ can be written

$$\theta_i^{N_i(\tau_i)} \exp\{-\theta_i \tau_i\}$$

(Continued on page 5.)

Suppose now that each process  $N_i(t)$  has a separate unobserved frailty variable  $Z_i$  such that, conditional on  $Z_i$ , the process  $N_i(t)$  is a homogeneous Poisson process with rate given by

$$\theta_i = Z_i \theta$$

Here  $\theta > 0$  is a common basic rate for the *n* processes, while  $Z_1, \ldots, Z_n$  are independent and gamma-distributed with expected value 1 and variance  $\delta > 0$ .

The definition and some results for the gamma-distribution are given at the end of this Problem.

- b) Find an expression for the *unconditional* likelihood contribution for the process  $N_i(t)$ .
- c) Show that the log-likelihood of the data from all the n processes can be written

$$\ell(\theta, \delta) = N \log \theta + \sum_{i=1}^{n} \log \left\{ \frac{\Gamma\left(N_i(\tau_i) + \delta^{-1}\right)}{\Gamma(\delta^{-1})} \right\}$$
$$- n\delta^{-1} \log \delta - \sum_{i=1}^{n} (N_i(\tau_i) + \delta^{-1}) \log(\theta\tau_i + \delta^{-1})$$

where  $N = \sum_{i=1}^{n} N_i(\tau_i)$ .

Derive the score function  $U_1(\theta, \delta) = \frac{\partial}{\partial \theta} \ell(\theta, \delta)$ .

Show that the maximum likelihood estimator  $\hat{\theta}$  for  $\theta$  can be found from this score function alone if the  $\tau_i$  are all equal, i.e.,  $\tau_i = \tau$  for  $i = 1, \ldots, n$ .

Find an expression for  $\hat{\theta}$  in this case, and give it a suitable interpretation.

How would you next find the maximum likelihood estimator for  $\delta$ ? (You need not do any calculations here).

d) For the shared frailty model, studied in the course, one considers how to recover (i.e., "estimate") the value of the frailty  $Z_i$  for each cluster *i*.

How would you, in the present Problem, compute recovered values for  $Z_i$  for i = 1, 2, ..., n based on the data from all the *n* processes? Find explicit expressions for the recovered  $Z_i$ .

#### The gamma distribution

If  $Z \sim \text{gamma}(k, \gamma)$ , for  $k, \gamma > 0$ , then

$$f_{Z}(t) = \frac{\gamma^{k}}{\Gamma(k)} t^{k-1} e^{-\gamma t} \text{ for } t > 0$$

$$E(Z) = \frac{k}{\gamma}$$

$$Var(Z) = \frac{k}{\gamma^{2}}$$

$$E\left[Z^{a} e^{-bZ}\right] = \frac{\Gamma(a+k)}{\Gamma(k)} \frac{\gamma^{k}}{(b+\gamma)^{a+k}} \text{ for } a, b \ge 0$$