

Name \_\_\_\_\_

I will adhere to the Hopkins code of academic ethics.

Signature \_\_\_\_\_

Lecture Section (please check one):

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**Biostatistics 140.623**  
**Third Term, 2014-2015**  
**Final Examination**  
**March 12, 2015**

**Instructions:** You will have two hours for this examination. There are 20 problems. The formula page and Stata output are at the **back** of the exam for your use. Please note that statistical significance is defined by  $p < 0.05$ .

**Questions 1-4 test general knowledge:**

1. What is the main purpose of the Cox regression model? (*Circle only one response*).
  - a) To estimate the survival function for a time-to-event outcome using binned data.
  - b) To estimate the baseline hazard function for a time-to-event outcome under the assumption that the relationship is linear on the log scale.
  - c) To test the assumption of proportional event hazards between risk factor groups.
  - d) To estimate and make inferences about relative event hazards between risk factor groups.
  - e) To determine whether the number at risk relates to covariates.
  
2. Suppose that you were interested in assessing differences in time to death by treatment group (drug versus placebo) and that the calculated log-rank test statistic for treatment equals 0.10, which is approximately a chi-squared statistic with one degree of freedom. The null hypothesis that corresponds to this test statistic is: (*Circle only one response*).
  - a) There are more deaths in the drug group.
  - b) There are more deaths in the placebo group.
  - c) There are equal numbers of deaths in the drug and placebo groups.
  - d) There is a difference in median survival between the drug and placebo groups.
  - e) There is no difference in the overall hazard of death between the drug and placebo groups.

3. The following is a Poisson regression model with 4 (follow-up) time bins (1-4) and treatment covariate defined as  $\text{trt}=1$  for Treatment A; 0 for Treatment B; and indicator variables for time bins 2, 3, and 4.

$$\begin{aligned} &\log(\text{expected events in bin } j) \\ &= \log(\text{person-weeks in bin } j) + \beta_0 + \beta_1(\text{time bin 2}) + \beta_2(\text{time bin 3}) + \beta_3(\text{time bin 4}) + \beta_4 \text{trt} \end{aligned}$$

This model assumes that: (*Circle only one response*)

- a) The hazard of an event is constant within a time bin but varies across time bins.
  - b) The hazard of an event may vary across time bins but increases within a time bin.
  - c) The hazard of an event is constant across time bins.
  - d) The relative hazard of an event for Treatment A versus Treatment B changes across time bin.
  - e) The relative hazard of an event for time bin  $j+1$  versus time bin  $j$  varies by treatment.
4. The AIC (Akaike Information Criterion) is a measure that can be used for: (*Circle only one response*)
- a) Assessing model goodness of fit.
  - b) Comparing observed versus expected outcomes.
  - c) Aiding in model selection based on the model log-likelihood and number of parameters.
  - d) Identifying statistically significant covariates in a model.
  - e) Checking the underlying model assumption of independence of observations.

**Questions 5 through 8** concern data from a study investigating the association between **sleep latency** (the amount of time needed for an individual to fall asleep at night) and **demographic characteristics**. **Models A-D** on pages 11-12 show logistic regression results.

**The outcome  $Y = \text{Slp15} = 1$  if sleep latency  $> 15$  minutes;  $= 0$  if  $\leq 15$  minutes**

**Demographic characteristics are:**

**age in years**

**female** = 1 if female; 0 if male

**smk** = 0 if never; 1 if current; 2 if former smoker

**BMI in  $\text{kg/m}^2$**

**bmicat** - BMI category

1 if  $< 18.5 \text{ kg/m}^2$

2 if  $18.5 - 24.9 \text{ kg/m}^2$

3 if  $25-29.9 \text{ kg/m}^2$

4 if  $\geq 30 \text{ kg/m}^2$

5. If age had been centered at the median age of 61 years in **Model A**, what would be the values of the estimated regression coefficients? (*Circle only one response*).
- a)  $b_0 = -1.27$  and  $b_1 = 0.019$
  - b)  $b_0 = -0.13$  and  $b_1 = 0.019$
  - c)  $b_0 = -1.27$  and  $b_1 = 1.16$
  - d)  $b_0 = 1.16$  and  $b_1 = 0.019$
  - e)  $b_0 = 0.019$  and  $b_1 = 1.16$
6. From **Model B**, we would conclude that the odds ratio for sleep latency > 15 minutes to fall asleep at night: (*Circle only one response*).
- a) Statistically significantly increases with each year of age for all individuals.
  - b) Statistically significantly increases with each year of age for individuals aged 55-65 years but not in younger nor in older individuals.
  - c) Statistically significantly decreases with each year of age for individuals aged > 65 years but not in younger individuals.
  - d) Statistically significantly decreases with each year of age for individuals < 55 years and > 65 years but not in individuals aged 55-65 years.
  - e) Is not statistically significantly associated with age in these individuals.
7. The results of the Likelihood Ratio Test of the Extended **Model D** to the Null **Model C** suggest that: (*Circle only one response*).
- a) BMI category does not contribute to the model of sleep latency beyond what is predicted by smoking status.
  - b) Smoking status does not contribute to the model of sleep latency beyond what is predicted by BMI.
  - c) Neither BMI nor smoking status contributes to the model of sleep latency.
  - d) Taken together, BMI and smoking statistically significantly contribute to the model of sleep latency.
  - e) Taken together, BMI and smoking status statistically significantly contribute to the model of sleep latency beyond what is predicted by age and its spline terms, and sex.

8. Suppose that, instead of handling BMI as a categorical variable, that BMI was used as a continuous variable using spline terms with knots at 18.5, 25, and 30 kg/m<sup>2</sup> using the following Stata command:

```
.mkspline bm1 18.5 bm2 25 bm3 30 bm4= bmi, marginal
```

The interpretation of the coefficient for **bm3** would be: (*Circle only one response*).

- The adjusted difference, between individuals with BMI 25-29 kg/m<sup>2</sup> and those with BMI 18.5 – 24.9 kg/m<sup>2</sup>, in the log odds of sleep latency > 15 minutes.
- The difference, between individuals with BMI 25-29.9 kg/m<sup>2</sup> and those with BMI 18.5-24.9 kg/m<sup>2</sup>, in the adjusted change in the log odds of sleep latency > 15 minutes with each kg/m<sup>2</sup> increase in BMI.
- The adjusted change in log odds of sleep latency > 15 minutes with each kg/m<sup>2</sup> increase in BMI among individuals with BMI 25-29.9 kg/m<sup>2</sup>.
- The adjusted log odds of sleep latency > 15 minutes in individuals with BMI ≥ 30 kg/m<sup>2</sup>.
- The adjusted change in average log odds of sleep latency > 15 minutes with each kg/m<sup>2</sup> increase in BMI in individuals with BMI ≥ 30 kg/m<sup>2</sup>.

**Questions 9 -11** reflect Poisson regression models of lung cancer deaths by age groups and population at risk in each age group. Variables are:

**Age** in age categories: < 45, 45-49, 50-54, 55-59, 60-64, 65-69, 70-74, 85-80, > 80

**Smoking category (smoke):**

- 1-smokes **neither** cigarettes nor cigars/pipes
- 2- smokes cigars/pipes **only**
- 3- smokes **both** cigarettes and cigars/pipes
- 4- smokes cigarettes **only**

Poisson regression **Models 1-3** are found on **pages 13-14**.

9. Comparing **Models 1-3**, it can be concluded that: (*Circle only one response*).
- Smoking confounds the relationship between age and lung cancer death rate.
  - Age confounds the relationship between smoking and lung cancer death rate.
  - Smoking modifies the relationship between age and lung cancer death rate.
  - Age modifies the relationship between smoking and lung cancer death rate.
  - Both smoking and age are mediators of the relationship between lung cancer deaths and the population at risk.

10. In **Model 2** which contains only smoking categories, it is assumed that: (*Circle only one response*).
- a) The incidence rate ratio of lung cancer death by smoking status is the same across age categories.
  - b) The incidence rate ratio of lung cancer death by smoking status varies by age category.
  - c) The incidence (hazard) of lung cancer death is not constant across age categories.
  - d) The incidence of lung cancer death changes linearly with age.
  - e) The incidence of lung cancer death is proportional to age.
11. From the **Model 3 output** we can see that, after controlling for age, the incidence rate of lung cancer death is significantly greater in individuals smoking cigarettes only as compared to both cigarettes and cigars/pipe. This is supported by: (*Circle only one response*).
- a)  $\log(\text{IRR})=0.218, Z=5.63, p=0.0$
  - b)  $\log(\text{IRR})=0.417, Z=10.45, p=0.0$
  - c)  $\text{IRR}=1.22, Z=8.39, p=0.0$
  - d)  $\text{IRR}=1.44, Z=9.74, p=0.0$
  - e)  $\text{LR } \chi^2_1 = 4034, p=0.0$

**Questions 12 through 15** concern the results from a randomized clinical trial of percutaneous coronary intervention (PCI) in patients with STEMI (acute ST-segment elevation myocardial infarction).

The researchers used simple **Cox regression** to measure the association between the primary outcome (a composite of death from cardiac causes, nonfatal myocardial infarction, or refractory angina) and treatment (PCI versus control). The model used is:

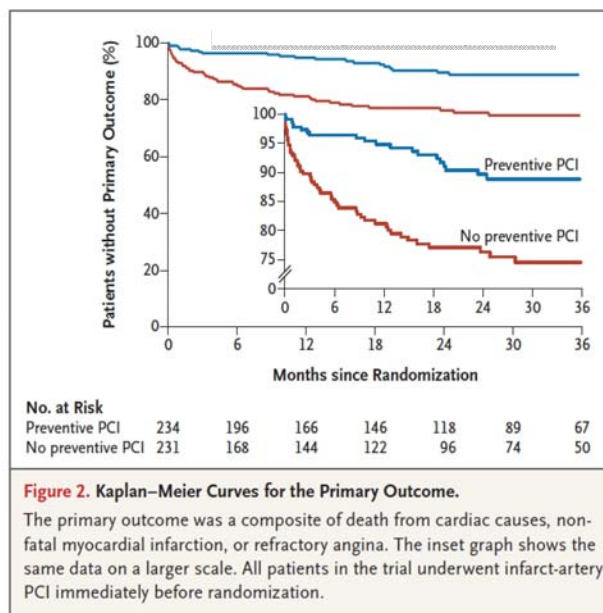
$$\ln(\text{hazard of primary outcome at time } t) = \ln(\lambda_0[t]) + \beta_1 x_1$$

where  $x_1 = 1$  for PCI intervention group and 0 for control group, and  $t$  represents time in the follow-up period (0 – 36 months).

12. What does the function  $\lambda_0(t)$  represent in the Cox regression equation? (*Circle only one response*).
- a) The hazard of the primary outcome in the PCI group at time = 0.
  - b) The hazard of the primary outcome in the control group at time=0.
  - c) The hazard ratio of the primary outcome for the PCI group compared to the control group at time=0.
  - d) The hazard of the primary outcome in the control group as a function of time across the follow-up period.
  - e) The difference in the  $\ln(\text{hazard})$  of primary outcome between the PCI and control groups at an time in the follow-up period.

13. What assumption did the researchers have to make in order to use Cox regression to quantify the relationship between the primary outcome and PCI (versus control)? (*Circle only one response*).
- The relationship between the primary outcome and PCI is statistically significant.
  - The hazard of the primary outcome is constant over time in both the PCI and control groups.
  - PCI will reduce the hazard of the primary outcome by at least 20%.
  - The relationship between the  $\ln(\text{hazard})$  of the primary outcome and time is linear.
  - The ratio of the hazard of the primary outcome for the PCI group compared to the control group is constant over the 36 month follow-up period.

The following shows the Kaplan-Meier curve estimates of the cumulative probability of being without the primary outcome in the **PCI** (Preventive PCI) and **control** groups (No Preventive PCI).



14. Based on the Kaplan-Meier curves above, what can be stated about the estimated value of  $\beta_1$  from the Cox regression model given on page 5? (*Circle only one response*).
- $b_1 > 0$
  - $b_1 < 0$
  - $b_1 = 0$
  - This cannot be answered without being given a specific time, and value of  $\hat{\lambda}_0[t]$  at this specified time.
  - This cannot be answered because there is no relationship between Kaplan-Meier curve estimates and the hazards of the primary outcome.

15. There were 234 patients randomized to the treatment group, and 67 still at risk of mortality at 36 months. In other words, 29% of the treatment group was still at risk of death at 36 months. However, the corresponding Kaplan-Meier curve estimate at 36 months for the treatment group is nearly 90%. How can this have happened? (*Circle only one response*).
- a) Some of the observations in the treatment group were censored (lost to follow-up or completed the study alive) in the Kaplan-Meier estimates.
  - b) The researchers estimated the Kaplan-Meier curve using only the data on patients who died in the follow-up period.
  - c) The researchers do not know how to properly estimate Kaplan-Meier curves.
  - d) The Kaplan-Meier curve estimate at 36 months ( $\hat{S}(36)$ ) is the risk of surviving among only those who were still alive and enrolled in the study at 36 months.
  - e) The researchers grouped the data into one-week time bins prior to plotting the survival curves.

**Questions 16-20** involve data from the UMARU impact study, a randomized trial of 595 subjects between 20 and 50 years old, with a substance abuse issue to assess the relative efficacy of long term versus short term residential drug treatment programs. Subjects were followed for up to 39 months after the start of treatment.

The following are baseline covariates in Cox regression **Models W- Z** which are found on **pages 15-17**.

**treat:** 1 for long-term, 0 for short-term treatment

**age\_cat:** takes on values 1-6 for 5-year age intervals; the age range in each of the intervals are [20, 25), [25, 30), [30, 35), [35,40), [40, 45) and [45, 50].

**white:** 1 if subject identifies as white, 0 if non-white.

**iv\_druguse:** 1 if subject was using intravenous (IV) drug at time of enrollment, 0 if not

16. Based on the result from **Model W**, what is the unadjusted hazard ratio (and 95% CI) of relapse for the long-term treatment group compared to the short term treatment group at 24 months after randomization? (*Circle only one response*).
- a) -0.24 (-0.42, -0.06)
  - b) 0.24 (0.06, 0.42)
  - c) 0.79 (0.66, 0.94)
  - d) 1.27 (1.06, 1.52)
  - e) This cannot be answered without being given the value of  $\hat{\lambda}_0[t=24 \text{ months}]$ .

17. Based on the results for **Models W- Y**, which of the following statements is true? (*Circle only one response*).
- a) The proportional hazards assumption with regard to the treatment groups is violated.
  - b) The relationship between time to relapse and treatment group is modified by age at enrollment.
  - c) The relationship between time to relapse and treatment group is substantially confounded by at least one of the following: IV drug use, age, and race.
  - d) The relationship between time to relapse and treatment group is not confounded by IV drug use, age, and race.
  - e) IV drug use is not a statistically significant predictor of time to relapse after accounting for treatment group.
18. Based on the result from **Model Y**, does the relationship between the hazard of relapse and age at enrollment appear to be linear on the log scale (after adjusting for treatment group, IV drug use and race)? (*Circle only one response*).
- a) No, because the AIC value for Model Y is smaller than the AIC values for Models W and X.
  - b) This cannot be answered without seeing the results of a Cox regression that includes age as a continuous predictor (as well as treatment group, IV drug use, and race as predictors)
  - c) This cannot be answered without having the p-value from a Likelihood ratio test comparing model Y to model X.
  - d) No, because the differences in the adjusted  $\ln(\text{hazard})$  are not similar in value for each consecutive pair of age categories (2 vs 1, 3 vs 2, etc.).
  - e) Yes, because some of the age category coefficients are statistically significant.
19. Which of the following is true based on the results from **Model Z**? (*Circle only one response*).
- a) Long-term treatment is more effective than short-term treatment in reducing the hazard of relapse, but only for white subjects (after adjusting for IV drug use and age at enrollment).
  - b) Long-term treatment is more effective than short-term treatment in reducing the hazard of relapse, but only for non-white subjects (after adjusting for IV drug use and age at enrollment).
  - c) The assumption of proportional hazards is violated because the interaction term (white\_treat) is statistically significant.
  - d) There is no difference in the hazards of relapse between the long-term and short term treatment programs after adjusting for race, IV drug use and age.
  - e) The relationship between time-to-relapse and race is modified by IV drug use.



20. Based on the results from **Model Z**, which of the following is the log hazard ratio of relapse at 24 months after randomization for 23- year old white subjects in long term treatment who used IV drugs versus (minus) 42- year old non-white subjects in short-term treatment who used IV drugs? (*Circle only one response*).

- a)  $\ln(\hat{\lambda}_o[24]) - 0.03 + 0.35 - 0.25 + 0.47$
- b)  $\ln(\hat{\lambda}_o[24]) - 0.03 + 0.35 - 0.25 + 0.47 + 0.38$
- c)  $-0.03 + 0.35 - 0.25 + 0.47$
- d)  $-0.03 + 0.35 - 0.25 + 0.47 + 0.38$
- e)  $-0.03 + 0.35 - 0.25 + 23 - 42$

## Biostatistics 140.623

### Final Exam Formula Sheet

Tabled chi-squared values: ( $\alpha=0.05$ )

$$\text{df}=1, \chi^2= 3.84$$

$$\text{df}=2, \chi^2= 5.99$$

$$\text{df}=3, \chi^2= 7.81$$

$$\text{df}=200, \chi^2= 233.99$$

$$Y = \beta_0 + \beta_1 X_1 + \beta_2 X_2 + \dots \varepsilon$$

$$F_{s, n-p-s-1} = \frac{(\text{RSS}_{\text{Null}} - \text{RSS}_{\text{Extended}}) / s}{\text{RSS}_{\text{Extended}} / (n-p-s-1)}$$

$$\text{AIC} = \text{RSS} + 2(\text{model df})$$

$$\ln = \log_e$$

$$\ln\left(\frac{a}{b}\right) = \ln(a) - \ln(b)$$

$$\frac{e^{a+b}}{e^a} = e^b$$

$$\log \text{odds} = \text{logit}[\Pr(Y=1)] = \beta_0 + \beta_1 X_1 + \beta_2 X_2 + \dots \beta_s X_s$$

$$\Pr(Y=1) = \frac{e^{\beta_0 + \beta_1 X_1 + \beta_2 X_2 + \dots \beta_s X_s}}{1 + e^{\beta_0 + \beta_1 X_1 + \beta_2 X_2 + \dots \beta_s X_s}} = \frac{\text{odds}}{1 + \text{odds}}$$

$$\text{LRT (Likelihood Ratio Test)} = -2 (\text{LL}_{\text{Null}} - \text{LL}_{\text{Extended}})$$

where LL = log likelihood

$$\text{AIC} = -2 \text{LL} + 2(\text{model df})$$

Poisson Regression (LLR) Model:

$$\log(\mu_i) = \log N_i + \beta_1 X_1 + \dots + \beta_p X_p$$

$$\log(\lambda_i) = \beta_1 X_1 + \dots + \beta_p X_p$$

Proportional Hazards Model:

$$\log \lambda(t; X) = \log \lambda_0(t; X) + \beta_1 X_1 + \dots + \beta_p X_p$$

$$\lambda(t; X) = \lambda_0(t; X) e^{\beta_1 X_1 + \dots + \beta_p X_p}$$

$$S(t; X) = [S_0(t)]^{e^{X\beta}}$$

Models A-D concern questions 5-8:

The outcome  $Y = \text{Slp15} = 1$  if sleep latency  $> 15$  minutes;  $= 0$  if  $\leq 15$  minutes

Demographic characteristics are: age in years

female = 1 if female; 0 if male      smk = 0 if never; 1 if current; 2 if former smoker

BMI in  $\text{kg/m}^2$

bmicat - BMI category: 1 if  $< 18.5 \text{ kg/m}^2$ ; 2 if  $18.5 - 24.9 \text{ kg/m}^2$ ; 3 if  $25-29.9 \text{ kg/m}^2$

Model A

```
. logit slp15 age
```

Logistic regression

```
Number of obs   =      821
LR chi2(1)      =       4.27
Prob > chi2     =      0.0387
Pseudo R2      =      0.0038
```

Log likelihood = -565.09366

slp15	Coef.	Std. Err.	z	P> z	[95% Conf. Interval]	
age	.018698	.0090718	2.06	0.039	.0009176	.0364784
_cons	-1.272739	.5571782	-2.28	0.022	-2.364788	-.18069

```
. est store A
```

Model B

```
. mkspline age1 55 age2 65 age3 = age, marginal
```

```
. logit slp15 age1 age2 age3
```

Logistic regression

```
Number of obs   =      821
LR chi2(3)      =       9.25
Prob > chi2     =      0.0261
Pseudo R2      =      0.0082
```

Log likelihood = -562.60413

slp15	Coef.	Std. Err.	z	P> z	[95% Conf. Interval]	
age1	-.0128556	.0361538	-0.36	0.722	-.0837158	.0580046
age2	.0819046	.0547896	1.49	0.135	-.0254811	.1892903
age3	-.1132878	.0509072	-2.23	0.026	-.213064	-.0135115
_cons	.2584022	1.886719	0.14	0.891	-3.439499	3.956303

```
. est store B
```

```
. lincom age1 +age2
```

```
( 1) [slp15]age1 + [slp15]age2 = 0
```

slp15	Coef.	Std. Err.	z	P> z	[95% Conf. Interval]	
(1)	.069049	.0258665	2.67	0.008	.0183515	.1197465

```
. lincom age1 +age2+ age3
```

```
( 1) [slp15]age1 + [slp15]age2 + [slp15]age3 = 0
```

slp15	Coef.	Std. Err.	z	P> z	[95% Conf. Interval]	
(1)	-.0442388	.0310547	-1.42	0.154	-.1051049	.0166273

Model C

```
. logit slp15 age1 age2 age3 female
```

```
Logistic regression
```

```
Number of obs = 821
```

```
LR chi2(4) = 9.59
```

```
Prob > chi2 = 0.0479
```

```
Pseudo R2 = 0.0085
```

```
Log likelihood = -562.43374
```

slp15	Coef.	Std. Err.	z	P> z	[95% Conf. Interval]	
age1	-.0129809	.0361621	-0.36	0.720	-.0838574	.0578956
age2	.0828356	.0548255	1.51	0.131	-.0246204	.1902917
age3	-.1144016	.050957	-2.25	0.025	-.2142754	-.0145278
female	.0823713	.1411278	0.58	0.559	-.1942341	.3589768
_cons	.2214594	1.888178	0.12	0.907	-3.479301	3.92222

```
. est store C
```

Model D

```
. logit slp15 age1 age2 age3 female i.smk i.bmicat
```

```
Logistic regression
```

```
Number of obs = 821
```

```
LR chi2(9) = 22.84
```

```
Prob > chi2 = 0.0066
```

```
Pseudo R2 = 0.0201
```

```
Log likelihood = -555.8081
```

slp15	Coef.	Std. Err.	z	P> z	[95% Conf. Interval]	
age1	-.0289882	.0371392	-0.78	0.435	-.1017796	.0438032
age2	.0963857	.0557898	1.73	0.084	-.0129603	.2057317
age3	-.0963259	.0516255	-1.87	0.062	-.1975101	.0048582
female	.0921986	.1426108	0.65	0.518	-.1873135	.3717106
smk						
Current	.2092384	.2036174	1.03	0.304	-.1898445	.6083212
Former	-.1891028	.1599506	-1.18	0.237	-.5026002	.1243945
bmicat						
2	.1040141	.9355023	0.11	0.911	-1.729537	1.937565
3	-.0053195	.9291496	-0.01	0.995	-1.826419	1.81578
4	.4902065	.9266137	0.53	0.597	-1.325923	2.306336
_cons	.8475727	2.131638	0.40	0.691	-3.330362	5.025507

```
. est store D
```

```
. lrtest D C
```

```
Likelihood-ratio test
```

```
LR chi2(5) = 13.25
```

```
(Assumption: C nested in D)
```

```
Prob > chi2 = 0.0211
```

```
. est stats *
```

```
Akaike's information criterion and Bayesian information criterion
```

Model	Obs	ll(null)	ll(model)	df	AIC	BIC
A	821	-567.2302	-565.0937	2	1134.187	1143.608
B	821	-567.2302	-562.6041	4	1133.208	1152.05
C	821	-567.2302	-562.4337	5	1134.867	1158.42
D	821	-567.2302	-555.8081	10	1131.616	1178.721

**Models 1-3** concerns questions 9 -11. **Variables** are:

**dead** (number of deaths) in each age group;

**population** (number at risk) in each age group.

**Age** in age categories: < 45, 45-49, 50-54, 55-59, 60-64, 65-69, 70-74, 85-80, > 80 years

Smoking category (**smoke**):

1-smokes **neither** cigarettes nor cigars/pipes

2- smokes cigars/pipes **only**

3- smokes **both** cigarettes and cigars/pipes

4- smokes cigarettes **only**

#### Model 1

. poisson dead i.age, exposure(pop)

$$\log(\lambda_j) = \beta_0 + \beta_1 \text{age}_2 + \beta_2 \text{age}_3 + \beta_3 \text{age}_4 + \beta_4 \text{age}_5 + \beta_5 \text{age}_6 + \beta_6 \text{age}_7 + \beta_7 \text{age}_8 + \beta_8 \text{age}_9$$

Poisson regression	Number of obs	=	36
	LR chi2(8)	=	3864.26
	Prob > chi2	=	0.0000
Log likelihood = -215.8728	Pseudo R2	=	0.8995

dead	Coef.	Std. Err.	z	P> z	[95% Conf. Interval]	
age						
45-49	.5560324	.0799878	6.95	0.000	.3992593	.7128056
50-54	.9881493	.0768149	12.86	0.000	.8375948	1.138704
55-59	1.371452	.0652555	21.02	0.000	1.243553	1.49935
60-64	1.628995	.0625358	26.05	0.000	1.506427	1.751563
65-69	1.957145	.0626921	31.22	0.000	1.834271	2.080019
70-74	2.205774	.0641042	34.41	0.000	2.080132	2.331416
75-79	2.457785	.0671346	36.61	0.000	2.326204	2.589367
80+	2.687489	.0708023	37.96	0.000	2.548719	2.826259
_cons	-3.395722	.0584206	-58.13	0.000	-3.510224	-3.281219
ln(pop)	1	(exposure)				

#### Model 2

. poisson dead i.smoke, exposure(pop)  $\log(\lambda_j) = \beta_0 + \beta_1 \text{smoke}_2 + \beta_2 \text{smoke}_3 + \beta_3 \text{smoke}_4$

Poisson regression	Number of obs	=	36
	LR chi2(3)	=	145.28
	Prob > chi2	=	0.0000
Log likelihood = -2075.3636	Pseudo R2	=	0.0338

dead	Coef.	Std. Err.	z	P> z	[95% Conf. Interval]	
smoke						
2.smoke	.3667831	.0466918	7.86	0.000	.2752688	.4582974
3.smoke	-.0633457	.038233	-1.66	0.098	-.1382809	.0115896
4.smoke	.054597	.0392158	1.39	0.164	-.0222646	.1314587
_cons	-1.839969	.0349215	-52.69	0.000	-1.908414	-1.771524
ln(pop)	1	(exposure)				

Model 3

```
. poisson dead i.age i.smoke, exposure(pop)
```

$$\log(\lambda_j) = \beta_0 + \beta_1 \text{age}_2 + \beta_2 \text{age}_3 + \beta_3 \text{age}_4 + \beta_4 \text{age}_5 + \beta_5 \text{age}_6 + \beta_6 \text{age}_7 + \beta_7 \text{age}_8 + \beta_8 \text{age}_9 + \\ + \beta_9 \text{smoke}_2 + \beta_{10} \text{smoke}_3 + \beta_{11} \text{smoke}_4$$

```
Poisson regression
```

```
Number of obs   =      36
LR chi2(11)     =    4034.50
Prob > chi2     =      0.0000
Pseudo R2      =      0.9391
```

```
Log likelihood = -130.75483
```

dead	Coef.	Std. Err.	z	P> z	[95% Conf. Interval]	
age						
45-49	.5538766	.0799886	6.92	0.000	.3971019	.7106514
50-54	.9803869	.0768183	12.76	0.000	.8298257	1.130948
55-59	1.379458	.0652606	21.14	0.000	1.25155	1.507367
60-64	1.654229	.0625688	26.44	0.000	1.531596	1.776861
65-69	1.998171	.0627875	31.82	0.000	1.87511	2.121232
70-74	2.271406	.0643537	35.30	0.000	2.145275	2.397537
75-79	2.558575	.0677844	37.75	0.000	2.42572	2.69143
80+	2.846925	.0724225	39.31	0.000	2.704979	2.98887
smoke						
2.smoke	.0478065	.0469926	1.02	0.309	-.0442972	.1399103
3.smoke	.2179552	.0386942	5.63	0.000	.142116	.2937945
4.smoke	.4169596	.0399121	10.45	0.000	.3387333	.4951859
_cons	-3.680024	.0682382	-53.93	0.000	-3.813769	-3.54628
ln(pop)	1	(exposure)				

```
. lincom 4.smoke - 3.smoke, irr
( 1) - [dead]3.smoke + [dead]4.smoke = 0
```

dead	IRR	Std. Err.	z	P> z	[95% Conf. Interval]	
(1)	1.220187	.028954	8.39	0.000	1.164738	1.278276

```
. lincom 4.smoke - 2.smoke, irr
( 1) - [dead]2.smoke + [dead]4.smoke = 0
```

dead	IRR	Std. Err.	z	P> z	[95% Conf. Interval]	
(1)	1.446509	.0548414	9.74	0.000	1.342918	1.558091

```
. lincom 3.smoke - 2.smoke, irr
( 1) - [dead]2.smoke + [dead]3.smoke = 0
```

dead	IRR	Std. Err.	z	P> z	[95% Conf. Interval]	
(1)	1.185481	.0431857	4.67	0.000	1.10379	1.273218

**Models W-Z** concern questions 16-20:

**treat:** 1 for long-term, 0 for short-term treatment

**age\_cat:** takes on values 1-6 for 5-year age intervals; the age range in each of the intervals are [20, 25), [25, 30), [30, 35), [35, 40), [40, 45) and [45, 50].

**white:** 1 if subject identifies as white, 0 if non-white.

**iv\_druguse:** 1 if subject was using intravenous (IV) drug at time of enrollment, 0 if not

**Model w:**  $\ln(\text{hazard of relapse at time } t) = \ln(\lambda_0[t]) + \beta_1 x_1$

```
. stcox treat, nohr
      failure _d:  censor == 1
      analysis time _t:  time
Cox regression -- Breslow method for ties
No. of subjects =          585          Number of obs   =          585
No. of failures =          471
Time at risk    =         141923
Log likelihood   =    -2710.1336          LR chi2(1)      =          6.86
                                          Prob > chi2      =          0.0088
-----+-----
      _t |          Coef.   Std. Err.      z    P>|z|     [95% Conf. Interval]
-----+-----
      treat |    -.2419827   .0923941    -2.62   0.009    -.4230717    -.0608936
-----+-----
. est store W
```

**Model x:**  $\ln(\text{hazard of relapse at time } t) = \ln(\lambda_0[t]) + \beta_1 x_1 + \beta_2 x_2$

```
. stcox treat iv_druguse, nohr
      failure _d:  censor == 1
      analysis time _t:  time
Cox regression -- Breslow method for ties
No. of subjects =          585          Number of obs   =          585
No. of failures =          471
Time at risk    =         141923
Log likelihood   =    -2704.3199          LR chi2(2)      =         18.49
                                          Prob > chi2      =          0.0001
-----+-----
      _t |          Coef.   Std. Err.      z    P>|z|     [95% Conf. Interval]
-----+-----
      treat |    -.2302159   .0924643    -2.49   0.013    -.4114426    -.0489893
      iv_druguse |   .3255089   .0967982     3.36   0.001     .135788     .5152299
-----+-----
. est store X
```

Model Y:

$\ln(\text{hazard of relapse at time } t) =$

$$\ln(\lambda_0[t]) + \beta_1 x_1 + \beta_2 x_2 + \beta_3 x_3 + \beta_4 x_4 + \beta_5 x_5 + \beta_6 x_6 + \beta_7 x_7 + \beta_8 x_8$$

```
. stcox treat white iv_druguse i.age_cat, nohr
```

```
      failure _d:  censor == 1
analysis time _t:  time
```

```
Iteration 0:  log likelihood = -2713.5637
Iteration 1:  log likelihood = -2696.0106
Iteration 2:  log likelihood = -2695.9678
Iteration 3:  log likelihood = -2695.9678
Refining estimates:
Iteration 0:  log likelihood = -2695.9678
```

Cox regression -- Breslow method for ties

```
No. of subjects =          585          Number of obs   =          585
No. of failures =          471
Time at risk    =         141923
Log likelihood   =   -2695.9678          LR chi2(8)      =         35.19
                                          Prob > chi2      =         0.0000
```

_t	Coef.	Std. Err.	z	P> z	[95% Conf. Interval]	
treat	-.2227576	.0926809	-2.40	0.016	-.4044088	-.0411064
white	.2096038	.112931	1.86	0.063	-.0117368	.4309444
iv_druguse	.386416	.1053305	3.67	0.000	.179972	.5928599
age_cat						
25-29	-.0605183	.1692869	-0.36	0.721	-.3923145	.271278
30-34	-.2179493	.1678659	-1.30	0.194	-.5469603	.1110618
35-39	-.1843114	.1770157	-1.04	0.298	-.5312558	.1626329
40-44	-.4944738	.2132287	-2.32	0.020	-.9123943	-.0765533
45-50	-.7889832	.3270934	-2.41	0.016	-1.430074	-.1478919

```
.est store Y
```

Model Z:

$\ln(\text{hazard of relapse at time } t) =$

$$\ln(\lambda_0[t]) + \beta_1 x_1 + \beta_2 x_2 + \beta_3 x_3 + \beta_4 x_4 + \beta_5 x_5 + \beta_6 x_6 + \beta_7 x_7 + \beta_8 x_8 + \beta_9 x_9$$

```
.gen white_treat = white*treat
```

```
. stcox treat white white_treat iv_druguse i.age_cat, nohr
```

```
      failure _d:  censor == 1
analysis time _t:  time
```

```
Iteration 0:  log likelihood = -2713.5637
Iteration 1:  log likelihood = -2695.3932
Iteration 2:  log likelihood = -2695.3293
```



Iteration 3: log likelihood = -2695.3293  
 Refining estimates:  
 Iteration 0: log likelihood = -2695.3293

Cox regression -- Breslow method for ties

No. of subjects =	585	Number of obs =	585
No. of failures =	471		
Time at risk =	141923		
Log likelihood =	-2695.3293	LR chi2(9) =	36.47
		Prob > chi2 =	0.0000

_t	Coef.	Std. Err.	z	P> z	[95% Conf. Interval]	
treat	-.0282404	.1965929	-0.14	0.886	-.4135554	.3570745
white	.347035	.1691883	2.05	0.040	.015432	.6786379
white_treat	-.2518304	.2236506	-1.13	0.260	-.6901775	.1865166
iv_druguse	.3809726	.1053208	3.62	0.000	.1745477	.5873975
age_cat						
25-29	-.0466745	.1695572	-0.28	0.783	-.3790005	.2856515
30-34	-.2019194	.1682786	-1.20	0.230	-.5317394	.1279006
35-39	-.1649912	.1775157	-0.93	0.353	-.5129156	.1829332
40-44	-.4664019	.2142489	-2.18	0.029	-.886322	-.0464817
45-50	-.782055	.3270817	-2.39	0.017	-1.423123	-.1409866

. lincom treat+ white\_treat

( 1) treat + white\_treat = 0

_t	Coef.	Std. Err.	z	P> z	[95% Conf. Interval]	
(1)	-.2800709	.1058858	-2.65	0.008	-.4876031	-.0725386

. est stats \*

Akaike's information criterion and Bayesian information criterion

Model	Obs	ll(null)	ll(model)	df	AIC	BIC
W	585	-2713.564	-2710.134	1	5422.267	5426.639
X	585	-2713.564	-2704.32	2	5412.64	5421.383
Y	585	-2713.564	-2695.968	8	5407.936	5442.908
Z	585	-2713.564	-2695.329	9	5408.659	5448.003-----

Note: N=Obs used in calculating BIC; see [R] BIC note