Precept 6: Duration models

Soc 504: Advanced Social Statistics

Ian Lundberg

Princeton University

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Outline

- Workflow
- 2 Duration
- 3 Using distributions
- 4 Zelig

We've gotten some questions about our personal project workflows.

Rmarkdown is in some ways ideal:

- Fully reproducible
- Code and results in one place

Problem: If code is slow to run, Rmarkdown is slow to compile each time.

I more often use R and LATEX:

- In RStudio, you can create a new R script. This is your code but does not produce a PDF.
- Save results (see ?save, ggsave, etc.)
- Produce final report in LATEX
 - I use TexShop
 - You can also work in an online platform like Overleaf.
 They also provide great templates!

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- 2 Duration
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Duration models are useful when we are interested in the

time T to an event

but some observations are **censored**: the event has not occurred at the end of data collection

Think, pair, share:

Why can't we do OLS when some observations are censored?

Duration models are useful when we are interested in the

time T to an event

but some observations are **censored**: the event has not occurred at the end of data collection

Think, pair, share:

Why can't we do OLS when some observations are censored? Because for those observations we don't know T!

The time to death T is a random variable. Its distribution is described by four critical functions:

- 1. Density function f(t)
 - Density of death at t
- 2. **CDF** F(t) = P(T < t)
 - Probability of death by t
- 3. Survival function S(t) = P(T > t) = 1 F(t)
 - Probability of survival to t
- 4. Hazard function $h(t) = \frac{f(t)}{S(t)}$
 - Density of death at t given survival up to t

Question: Why isn't the hazard function a probability?



Photo credit: J Zamudio via https://www.nps.gov/yose/planyourvisit/stargazing.htm

PDF

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CDF

$$F(t) = 1 - e^{-\lambda t}$$

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We just proved the memoryless property! How?

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We just proved the memoryless property! How? h(t) is not a function of t. The hazard is constant.

Suppose we want to allow the hazard to vary by some set of predictors.

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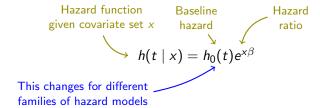
Suppose we want to allow the hazard to vary by some set of predictors.

$$h(t \mid x) = h_0(t)e^{x\beta}$$

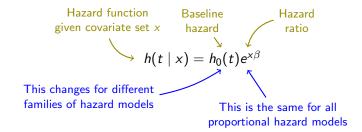
Suppose we want to allow the hazard to vary by some set of predictors.

Hazard function Baseline Hazard given covariate set
$$x$$
 hazard ratio
$$h(t \mid x) = h_0(t)e^{x\beta}$$

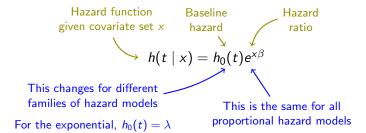
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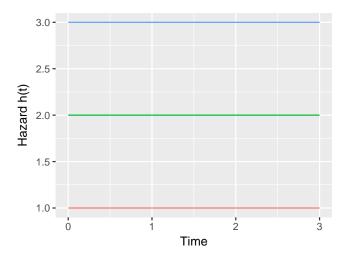
Why add covariates?

Why add covariates? It might be cloudy.



Photo credit: Hannah Lundberg

Exponential hazards



Question: If the green is the baseline hazard $h_0(t)$, what is the hazard ratio that produces the blue line? The red line?

Fitting an Exponential with survreg

```
> library(survival)
> fit <- survreg(Surv(time, event) ~ age + sex,
                dist = "exponential",
+
+
                data = lung)
> summary(fit)
Call:
survreg(formula = Surv(time, event) ~ age + sex, data = lung,
   dist = "exponential")
             Value Std. Error z
(Intercept) 6.3597 0.63547 10.01 1.41e-23
         -0.0156 0.00911 -1.72 8.63e-02
age
            0.4809 0.16709 2.88 4.00e-03
sex
Exponential distribution
Loglik(model) = -1156.1 Loglik(intercept only) = -1162.3
Chisq= 12.48 on 2 degrees of freedom, p= 0.002
Number of Newton-Raphson Iterations: 4
n = 228
```

Interpreting hazard ratios

Q: How would you interpret these?

Interpreting hazard ratios

$$h(t \mid x) = h_0(t)e^{-x\beta}$$

Q: How would you interpret these?

A year increase in age is associated with a 1.6% increase in the hazard, holding sex constant.

There are some things demographers just memorize.

We recommend just looking these up when you need them.

For instance, this fact:

The survival function is e to the minus cumulative hazard.

Hazard function \rightarrow survival function

The derivative of the negative log of the survival function is

$$\frac{\partial}{\partial t} \left(-\log \left[S(t) \right] \right) = \frac{\frac{\partial}{\partial t} \left(-S(t) \right)}{S(t)}$$

$$= \frac{\frac{\partial}{\partial t} \left(-\left[1 - F(t) \right] \right)}{S(t)}$$

$$= \frac{f(t)}{S(t)} = h(t)$$

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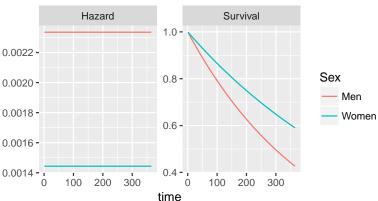
$$= \frac{f(t)}{S(t)} = h(t)$$

Doing the reverse, we can go from h(t) to S(t)

$$\int_0^t \frac{\partial}{\partial t'} \left(-\log \left[S(t') \right] \right) dt = \int_0^t h(t') dt'$$
$$-\log \left[S(t) \right] = \int_0^t h(t') dt'$$
$$S(t) = e^{-\int_0^t h(t') dt'}$$

Plotting survival curves

Exponential survival fits for 50-year-old men and women



Plotting survival curves

How we made the previous slide:

The exponential is almost always parameterized with a rate λ .

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But, it could just as well be defined in terms of a scale $\theta = \frac{1}{\lambda}$

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Using distributions

Zelig

Scales and rates

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$f(T) = \lambda e^{-\lambda x}$	$f(T) = \frac{1}{\theta} e^{-\frac{1}{\theta}x}$

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As rate grows, expected waiting time shrinks

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In general, you have to be careful with the parameterization of survival distributions.

What if we want the hazard to be a function of time?

What if we want the hazard to be a function of time?

Many options.

$$T \sim \mathsf{Weibull}(\alpha, \lambda)$$

PDF ¹

$$f(t) = t^{\alpha - 1} \alpha \lambda^{\alpha} e^{-(\lambda t)^{\alpha}}$$

CDF

$$F(t) = 1 - e^{-(\lambda t)^{\alpha}}$$

Survival function

$$S(t) = P(T > t) =$$

 $^{^{1}}$ I have used the rate parameterization for λ ; lecture slides use the scale parameterization.

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$$h(t \mid \alpha = 1) = t^{\alpha - 1} \alpha \lambda^{\alpha} = t^{1 - 1} 1 \lambda^{1} = \lambda$$

Discussion: If the Weibull contains the Exponential as a special case, why not always use the Weibull?

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If the world is actually Exponential, we gain **efficiency** by making the assumption that the hazard is constant over time.

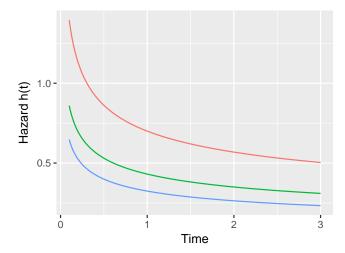
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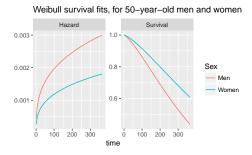
This is a **general theme** of statistics: Modeling assumptions buy us efficiency if they are correct.

Weibull hazards



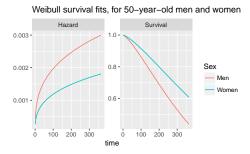
Fitting a Weibull model

Weibull results



Common question: The gap between those hazards clearly changes over time! Is this a violation of a modeling assumption?

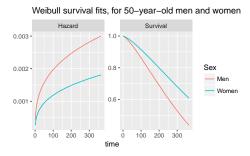
Weibull results



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Weibull results



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A: No, they are still proportional!

(Also since these are fitted values, they necessarily agree with the modeling assumptions, so this was a trick question.)

You can fit a survival model using any distribution for which the support is all positive numbers.

There are a huge number of options.

$$T \sim \mathsf{LogNormal}(\mu, \sigma^2) \sim e^Z$$
 (where $Z \sim \mathit{N}(\mu, \sigma^2)$

$$f(t) = \frac{1}{Y_i \sigma \sqrt{2\pi}} \exp\left(-\frac{(\ln(Y_i) - \mu)^2}{2\sigma^2}\right)$$

CDF

$$F(t) = \int_0^t f(x)dx = \text{ugly formula}$$

Survival function

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Hazard function: Risk of event at t given survival up to t

$$T \sim \mathsf{LogNormal}(\mu, \sigma^2) \sim \mathsf{e}^Z \; (\mathsf{where} \; Z \sim \mathsf{N}(\mu, \sigma^2))$$

$$f(t) = \frac{1}{Y_i \sigma \sqrt{2\pi}} \exp\left(-\frac{(\ln(Y_i) - \mu)^2}{2\sigma^2}\right)$$

CDF

$$F(t) = \int_0^t f(x)dx = \text{ugly formula}$$

Survival function

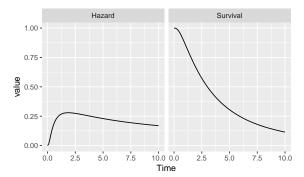
$$S(t) = P(T > t) = 1 - P(T < t) = 1 - F_T(t) = \text{ugly formula}$$

Hazard function: Risk of event at t given survival up to t

$$h(t) = \frac{f(t)}{S(t)} = \text{ugly formula}$$

Fitting a Lognormal

Note: This figure doesn't correspond to the model above - just an example of a LogNormal



$$f(t) = b\eta e^{bt} e^{\eta} \exp(-\eta e^{bt})$$

$$f(t) = b\eta e^{bt} e^{\eta} \exp(-\eta e^{bt})$$

$$F(t) = 1 - \exp(-\eta \left(e^{bt} - 1\right))$$

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$$h(t) = \frac{f(t)}{S(t)}$$

$$= \frac{b\eta e^{bt} e^{\eta} \exp(-\eta e^{bt})}{\exp(-\eta\left(e^{bt} - 1\right))}$$

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 $= \frac{b\eta e^{bt} e^{\eta} \exp(-\eta e^{bt})}{\exp\left(-\eta\left(e^{bt} - 1\right)\right)}$
 $= b\eta e^{bt} e^{\eta}$
 $\log[h(t)] = \underbrace{(\log(b) + \log(\eta) + \eta)}_{\text{Intercept}} + \underbrace{b}_{\text{Slope}} t$

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$$= \alpha + \beta t$$

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The log of the hazard function is linear in time!

This is why people like the Gompertz.

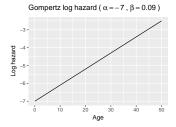
Gompertz hazard with $\alpha = -7, \beta = .09$

$$\log[h(t)] = \alpha + \beta t, \quad h(t) = \exp(\alpha + \beta t)$$

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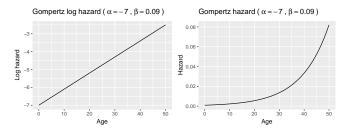
Q: If the $\log[h(t)]$ increases linearly with t, what does h(t) look like?



Gompertz hazard with $\alpha = -7, \beta = .09$

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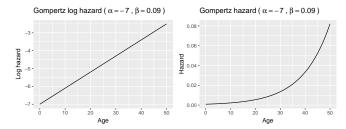
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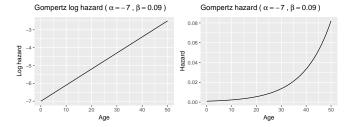


Q: For what questions would this be a good choice?

Gompertz hazard with $\alpha = -7, \beta = .09$

$$\log[h(t)] = \alpha + \beta t, \quad h(t) = \exp(\alpha + \beta t)$$

Q: If the $\log[h(t)]$ increases linearly with t, what does h(t) look like?



Q: For what questions would this be a good choice? Mortality

Note: Example motivated by U.S. mortality; see German Rodriguez's example

here.

Time between breaks while hiking out of this valley. You don't need a rest right away...



Donahue Pass, Yosemite. Photo credit: Riley Brian

...but after going for a while your hazard of resting increases. Gompertz.

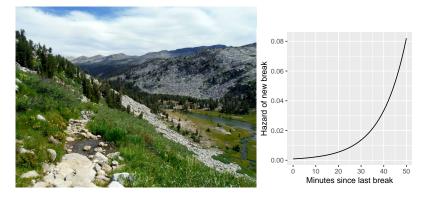
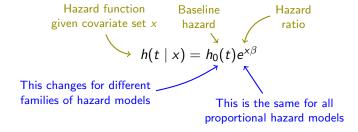


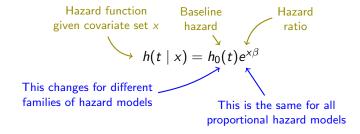
Photo credit: Riley Brian

As I said at the beginning, all of the survival models above have the form:



Different models allow different kinds of flexibility in the baseline hazard $h_0(t)$.

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Different models allow different kinds of flexibility in the baseline hazard $h_0(t)$.

Can we model hazard ratios without any assumptions about $h_0(t)$?

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The Cox model is fit based on the order at which people die, rather than the times, so it does not assume a baseline hazard.

You can fit one with coxph()

Outline

- 1 Workflow
- 2 Duration
- 3 Using distributions
- 4 Zelig

Using distributions

Most common question we are asked:

How do I know when to use a given distribution for a given problem?

Using distributions

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How do I know when to use a given distribution for a given problem?

When you know the **story of the distributions**, you can find one that **maps onto** your current problem.

Suppose someone says to you, "I ran 10 hypothesis tests. What's the probability that the at least 1 p-values is less than 0.05 if all the null hypotheses are true?"

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You draw this picture.

```
Want the distribution of this order statistic
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You reply:

"You want to know the distribution of the order statistic $U_{(3)}$.

Suppose someone says to you, "I ran 10 hypothesis tests. What's the probability that the at least 1 p-values is less than 0.05 if all the null hypotheses are true?"

You draw this picture.

You reply:

"You want to know the distribution of the order statistic $U_{(3)}$. Let me take you to the wilderness. We will count shooting stars."



 $PC: \ http://wilderness.org/30-prettiest-lakes-wildlands$

We will count shooting stars and record the times we see them.²



²Thanks to William Chen for the shooting stars example. See more at http://www.wzchen.com/probability-cheatsheet/□ → ←② → ←② → ←② → ←② → ◆② ←

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$$\begin{array}{c} & \longrightarrow \\ 0 & \text{Time} \end{array}$$

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$$\begin{array}{c} & & \\ & \downarrow \\ \end{array}$$

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$$\begin{array}{c} & & \\ & \times \times \times \\ & & \\ \end{array}$$

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Shooting stars come at a **constant rate**.

The times between the arrivals are $X_1, X_2, \ldots \stackrel{\mathsf{iid}}{\sim} \mathsf{Exponential}(\lambda)$.

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Q: What is the distribution of X_1 given this information?

$$X_1$$

$$X_1 + X_2 = 1$$

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Q: If I run one hypothesis test, what is the probability under the null that it falls below 0.05?

A:
$$P(U < .05) = P\left(\frac{X_1}{X_1 + X_2} < .05\right) = 0.05$$

Now suppose we observe X_1, \ldots, X_{11} and we rescale so their sum is 1.

Let's re-label the imes marks with U values with arbitrary indexes.

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There is a connection between *p*-values and shooting stars.

Order statistics

Let's denote the k-th order statistic by $U_{(k)}$.

$$U_{(k)} = \frac{\sum_{i=1}^{k} X_i}{\sum_{i=1}^{11} X_i}$$

A new distribution: The Beta

If
$$X_1,\ldots,X_n\sim$$
 Exponential, then $\frac{\sum_{i=1}^k X_i}{\sum_{i=1}^n X_i}\sim$ Beta(k,n - k - 1)

So
$$U_{(1)} \sim \text{Beta}(1, 10)$$
.

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So
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Q: Can you reason about the expected value of a Beta(1,10)?

A: There are 11 white space that we would expect to be of equal size, so we might expect that $E(U_{(1)}) = \frac{1}{11}$. This is right!

Q: Given what we know about shooting stars, what distribution do you think the smallest *p*-value takes?

$$p_{(1)} \sim \mathsf{Beta}(1,9)$$

Q: What is the probability that the smallest *p*-value is less than 0.05?

$$P(p_{(1)} < .05) = P(\text{Beta}(1,10) < .5) = F_{\text{Beta}(1,9)}(.05) = 0.37$$

It is very easy to get a false positive by running 10 hypothesis tests!

Key takeaways

We've taught you the stories of many distributions.

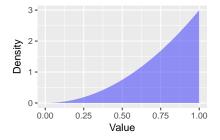
To use them, try to fit your problem into one of these **known stories**!



Photo credit: Hannah Lundberg



In my own research, I wanted to choose a prior distribution on a correlation that I expected to be near 1. I chose Beta(3,1).



I chose that by thinking:

- I want the distribution of the highest of 3 uniform draws.
- I want the distribution of the proportion of time spent waiting for 3 shooting stars, out of a total time spend waiting for 4.

Plugging your problem into a **known story** can help you find a solution.



Generalizing that story

Suppose someone says to you, "I ran 100 hypothesis tests. What's the probability that the 7th-smallest p-value is less than 0.05 if the null hypotheses are true?"

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That is the proportion of time spent waiting for the 7th shooting star:

$$U_{(7)} \sim \mathsf{Beta}(7,93)$$
 $P(U_{(7)} < .05) = F_{\mathsf{Beta}(7,93)}(.05) = 0.23$

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So, it's not that strange to see 7 p-values less than 0.05. And we learned this all from shooting stars!

One other story you might use

What if we wanted a distribution for the time until the *k*th star comes?

$$X_1, \ldots, X_k \stackrel{\mathsf{iid}}{\sim} \mathsf{Exponential}(\lambda)$$

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 $G_k \sim X_1 + \dots + X_k$

Then we say

$$G_k \sim \mathsf{Gamma}(k, \lambda)$$

The **Gamma distribution** characterizes the wait time until the *k*th star.

Outline

- 1 Workflow
- 2 Duration
- 3 Using distributions
- 4 Zelig

Side note: Zelig

Zelig is an R package designed to make everything we do in class easier.

Note the Zelig workflow overview.

We will use the Zelig-Exponential.

Zelig example: Lung cancer survival

We will walk through the example using data on lung cancer survival

```
> library(survival)
```

- > data(lung)
 > head(lung)
- > head(lung)

```
inst time status age sex ph.ecog ph.karno pat.karno meal.cal wt.loss
     3 306
                 2 74
                                                    100
                                                            1175
                                                                      NA
                                          90
        455
                 2 68
                                                     90
                                                            1225
                                                                      15
                                          90
     3 1010
                 1 56
                                          90
                                                     90
                                                              NΑ
                                                                      15
4
     5 210
                 2 57 1
                                          90
                                                     60
                                                            1150
                                                                      11
5
        883
                 2 60
                                         100
                                                     90
                                                              NΑ
    12 1022
                    74
                                          50
                                                    80
                                                             513
```

lung <- mutate(lung, event = as.numeric(status == 2))</pre>

Variable definitions: Lung cancer survival

?lung

inst: Institution code
time: Survival time in days

status: censoring status 1=censored, 2=dead

age: Age in years

sex: Male=1 Female=2

ph.ecog: ECOG performance score (0=good 5=dead)

ph.karno: Karnofsky performance score (bad=0-good=100) rated by physician

pat.karno: Karnofsky performance score as rated by patient

meal.cal: Calories consumed at meals
wt.loss: Weight loss in last six months

Zelig step 1: Fit a model

Zelig step 1: Fit a model

```
> summary(fit)
Model:
Call:
z5$zelig(formula = Surv(time, event) ~ age + sex, data = lung)
             Value Std. Error z
(Intercept) 6.3597 0.63547 10.01 1.41e-23
         -0.0156 0.00911 -1.72 8.63e-02
age
           0.4809 0.16709 2.88 4.00e-03
sex
Scale fixed at 1
Exponential distribution
Loglik(model) = -1156.1 Loglik(intercept only) = -1162.3
Chisq= 12.48 on 2 degrees of freedom, p= 0.002
Number of Newton-Raphson Iterations: 4
n = 228
Next step: Use 'setx' method
```

Zelig step 2: Use setx to set covariates of interest

```
men <- setx(fit, age = 50, sex = 1)
women <- setx(fit, age = 50, sex = 2)</pre>
```

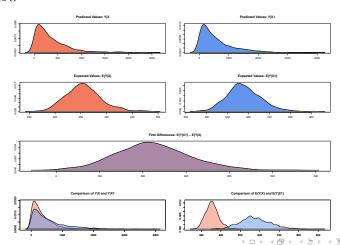
Zelig step 2: Use setx to set covariates of interest

Zelig step 3: Use sim to simulate quantities of interest

```
> sims <- sim(obj = fit, x = men, x1 = women)
> summary(sims)
 sim x :
 ----
ev
                sd
                        50%
                                2.5%
     mean
1 355.086 33.63733 353.5258 296.6169 428.758
pv
                          50%
                                  2.5%
                                           97.5%
        mean
                   sd
[1,] 351.414 361.6174 242.511 7.082744 1357.005
 sim x1:
 ----
ev
                       50%
                               2.5%
1 577.5684 78.5113 571.178 438.4341 743.9957
pν
                            50%
                                   2.5%
[1,] 562.8317 550.6102 382.9658 11.5627 2016.61
fd
      mean
                sd
                        50%
                                2.5%
1 222 4824 85 0493 217 0278 61 08082 396 5632
```

Zelig step 4: Use graph to plot simulation results

```
pdf("ZeligFigures.pdf",
    height = 5, width = 7)
plot(sims)
dev.off()
```



Estimate your model:

Set your covariates:

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```
men <- setx(fit, sex = 1, fn = mean)
women <- setx(fit, sex = 2, fn = mean)</pre>
```

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```
men <- setx(fit, sex = 1, fn = mean)
women <- setx(fit, sex = 2, fn = mean)</pre>
```

Simulate your QOI:

```
#install.packages("Zelig")
require(Zelig)
fit <- zelig(Surv(time, event) ~ age + sex,
             model = "exp",
              data = lung)
Set your covariates:
men <- setx(fit, sex = 1, fn = mean)
women <- setx(fit, sex = 2, fn = mean)
Simulate your QOI:
sims \leftarrow sim(obj = fit, x = men, x1 = women)
```

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```

After break: expectation maximization, missing data

Cards! Questions?



Photo credit: Riley Brian