Introduction and Background

Dave Harrington

May 14 - 18, 2018

Administrative Information

Important Definitions

Survival Distributions

Administrative Information

Instructor coordinates

Dave Harrington

- Department of Biostatistics, Harvard T.H. Chan School of Public Health
- Department of Biostatistics and Computational Biology, Dana-Farber Cancer Institute
- Email: dharrington846@gmail.com

Course format

- Lectures, but please ask questions
- Labs (practical exercises)
- Readings (typically, papers from literature)
- Computing (in R)

ACKNOWLEDGEMENTS

Many individuals have contributed to these notes in courses at Harvard, Ghent University, Hasselt University, and other institutions.

Special acknowledgements to

- Rui Wang, Paige Williams, Rebecca Betensky, and Paul Catalano (Harvard)
- Louise Ryan (University of Technology, Sydney)
- Els Goetghebeur (U Ghent)
- Julie Vu (University College London)

Important Definitions

Main ideas

Survival analysis typically focuses on time-to-event data.

Examples:

- Time to death from a chronic disease
- Time to progression of a disease
- Time to onset (or relapse) of a disease
- Length of stay in a hospital or a nursing home

The terms *survival time* and *event time* will be used to mean the time to an event.

Most useful references

- Klein and Moeschberger: Survival Analysis: Techniques for censored and truncated data
- Therneau and Grambsch: Modeling Survival Data: extending the Cox Model (R)
- Collett: Modelling Survival Data in Medical Research
- Hosmer, Lemeshow, and May: Applied Survival Analysis, 2nd ed.
- Kleinbaum: Survival Analysis: A self-learning text
- Cox and Oakes: Analysis of Survival Data

I recommend the K&M text, supplemented by T&G.

A FEW MORE REFERENCES

- Fleming and Harrington: Counting Processes and Survival Analysis
- Kalbfleisch and Prentice: The Statistical Analysis of Failure Time Data
- Allison: Survival Analysis Using the SAS System
- Miller: Survival Analysis

PACKAGES USED

Example: Time to death or hospitalization

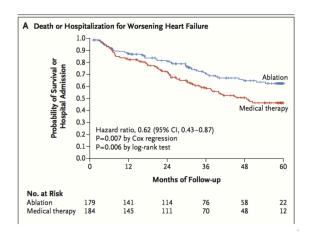


Figure 1: Figure from Marrouche, et al., NEJM 2018

See Marrouche, et al.

STRUCTURE OF EVENT TIME DATA

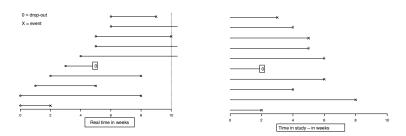


Figure 2: Event time data as observed (L) versus to a data analyst (R) $\,$

Characteristics of event time data

- 'Individuals' do not all enter the study at the same time.
 - This is referred to as staggered entry.
- When the study ends, some individuals still haven't had the event.
- Other individuals drop out or are lost during the study.
 - The last time they were still "free" of the event is all that is known.

The last two features relate to *censoring* of the failure times.

The first of the times until the study ends or the subject drops out is called a *censoring time*.

A HYPOTHETICAL EXAMPLE

Assume 10 subjects assigned to each of four treatments after cancer remission, followed until death or end of study at 36 months.

The times to death (months):

- **Trt 1:** Deaths at 2, 3, 7, 9, 15, 16 (4 alive at end)
- **Trt 2:** Deaths at 1, 1, 2, 4, 4, 6, 7, 9, 11 (1 alive at end)
- **Trt 3:** Deaths at 1, 1, 2, 4, 4, 5 (4 alive at end)
- Trt 4: Deaths at 2, 3, 7, 9, 15, 22, 27, 28, 29 (1 alive at end)
 - For Treatment 1, what is the average time to death?
 - Comparing Treatments 1 and 2, which appears better?
 - Comparing Treatments 1 and 3, which appears better?
 - Comparing Treatments 3 and 4, which appears better?

DESCRIPTIVE COMPARISONS OF "AVERAGE" DEATH TIME

Treatment	Among Deaths:		Median adjusting	
Group	Mean	Median	for Censoring (KM)	
1	8.67	8.00	15.5	
2	5.00	4.00	5.0	
3	2.83	3.00	4.5	
4	15.78	15.00	18.5	

Comparisons of treatments (p-values)

Methods to do these calculations coming in this course.

Comparison	Log-Rank Test	Wilcoxon Test	Exponential Model
1 vs 2	0.045	0.048	0.014
3 vs 4	0.62	0.67	0.56
1 vs 3	0.63	0.37	0.73
2 vs 4	0.15	0.06	0.09
1 vs 4	0.37	0.68	0.34

More key features of survival data

- Survival times are often right-skewed, so the median is usually a better measure of center than the mean.
- The median can be estimated from data that include censored observations (not always possible with the mean).
- Calculating summary statistics and comparing survival distributions must account for cases without events.
- Comparisons between survival distributions may yield different conclusions depending on assumptions.

Types of censoring

Right-censoring

Let T_i be the time to event and U_i be the time to censoring for an individual i. Only the r.v. $X_i = \min(T_i, U_i)$ is observed, due to

- loss to follow-up
- drop-out
- study termination ('administrative censoring')

Called *right*-censoring because the true unobserved event is to the right of the censoring time (i.e., after the censoring time).

In addition to X_i , the *failure indicator* δ_i is observable:

$$\delta_i = \begin{cases} 1 & \text{if} \quad T_i \le U_i \\ 0 & \text{if} \quad T_i > U_i \end{cases}$$

Types of censoring ...

Left-censoring

The r.v.'s observed are $Y_i = \max(T_i, U_i)$ and the failure indicator ϵ_i :

$$\epsilon_i = \left\{ \begin{array}{ll} 1 & \text{if} \quad U_i \le T_i \\ 0 & \text{if} \quad U_i > T_i \end{array} \right.$$

Ex.: study of age at which African children learn a task (Miller).

- Some already knew the task (left-censoring)
- Some learned during study (exact)
- Some had not yet learned by end of study (right-censoring)

Types of censoring...

Interval-censoring

Observe (L_i, R_i) where $T_i \in (L_i, R_i)$

Examples:

- Time to prostate cancer, observe longitudinal PSA measurements
- Time to undetectable viral load in AIDS studies, based on measurements of viral load taken at each clinic visit
- Detect recurrence of colon cancer after surgery. Follow patients every 3 months after resection of primary tumor.

This short course is restricted to right-censoring.

Independent versus informative censoring

Censoring is **independent** if U_i is independent of T_i .

Examples:

- If U_i is the planned end of the study (say, 2 years after the study opens), then it is usually independent of the event times.
 - What if there is a trend over calendar time in the survival times?
- If U_i is the time that a patient drops out of the study because they have become much sicker and/or had to discontinue taking the study treatment, then U_i and T_i are probably not independent.

An individual censored at U should be *representative* of all subjects who survive to U.

Censoring is considered **informative** if the distribution of U_i contains information about the parameters characterizing the distribution of T_i .

Survival Distributions

SOME MATHEMATICAL DEFINITIONS

There are several equivalent ways to characterize the probability distribution of a survival random variable.

- The density function f(t)
- The survivor function S(t)
- The hazard function $\lambda(t)$
- The cumulative hazard function $\Lambda(t)$

Some are special to survival analysis.

Density function

For a discrete random variable

Suppose that T takes values in a_1, a_2, \ldots, a_J .

$$f(t) = P(T = t)$$

$$= \begin{cases} f_j & \text{if} \quad t = a_j, j = 1, 2, \dots, J \\ 0 & \text{if} \quad t \neq a_j, j = 1, 2, \dots, J \end{cases}$$

For a continuous random variable

$$f(t) = \lim_{\Delta t \to 0} \frac{1}{\Delta t} P(t \le T < t + \Delta t)$$

Survivorship function S(t)

In other settings, the cumulative distribution function

$$F(t) = P(T \le t) = 1 - S(t)$$

is of interest.

In survival analysis, interest tends to focus on S(t) = P(T > t), the survival or survivorship function.¹

The survival function measures the probability an individual will experience the event beyond time t.

¹Be careful: some books use the definition $S(t) = P(T \ge t)$.

Survivorship function S(t) ...

For a continuous random variable:

$$S(t) = \int_{t}^{\infty} f(u) du$$

For a discrete random variable:

$$S(t) = \sum_{u>t} f(u) = \sum_{a_j>t} f(a_j) = \sum_{a_j>t} f_j$$

HAZARD FUNCTION $\lambda(t)$

The hazard function measures the probability of death at time t, conditional on having survived until that time.

This is sometimes called the instantaneous failure rate.

For a *continuous* random variable T:

$$\lambda(t) = \lim_{\Delta t \to 0} \frac{1}{\Delta t} P(t \le T < t + \Delta t | T \ge t)$$

$$= \lim_{\Delta t \to 0} \frac{1}{\Delta t} \frac{P([t \le T < t + \Delta t] \cap [T \ge t])}{P(T \ge t)}$$

$$= \lim_{\Delta t \to 0} \frac{1}{\Delta t} \frac{P(t \le T < t + \Delta t)}{P(T \ge t)}$$

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

HAZARD FUNCTION $\lambda(t)$...

For a discrete random variable T:

$$\lambda(a_j) \equiv \lambda_j = P(T = a_j | T \ge a_j) = \frac{P(T = a_j)}{P(T \ge a_j)}$$
$$= \frac{f(a_j)}{S(a_j -)} = \frac{f(t)}{\sum_{k: a_k \ge a_j} f(a_k)}$$

The form of the denominator for both continuous and discrete variables is the reason some books use $P(T \ge t)$ as the definition of the survivor function.

Cumulative hazard function $\Lambda(t)$

For a *continuous* random variable *T*:

$$\Lambda(t) = \int_0^t \lambda(u) du$$

For a *discrete* random variable *T*:

$$\Lambda(t) = \sum_{k: a_k \le t} \lambda_k$$

Relationship between S(t) and $\lambda(t)$

For a continuous random variable,

$$\lambda(t) = \frac{f(t)}{S(t)}.$$

For a right-continuous survivor function S(t),

$$f(t) = -S'(t) \text{ or } S'(t) = -f(t).$$

These relationships can be used to show another way to write $\lambda(t)$:

$$\lambda(t) = \frac{f(t)}{S(t)} = -\frac{f(t)}{S(t)} = -\left(\frac{1}{S(t)}\right)S'(t) = -\frac{d}{dt}[\log S(t)]$$

$$\lambda(t) = -\frac{d}{dt}[\log S(t)]$$

Relationship between S(t) and $\Lambda(t)$

For a continuous random variable:

$$\Lambda(t) = \int_0^t \lambda(u) du$$

$$= \int_0^t -\frac{d}{du} \log S(u) du$$

$$= -\log S(t) + \log S(0)$$

Thus,
$$S(t) = e^{-\Lambda(t)}$$
.

Relationship between S(t) and $\Lambda(t)$...

For a discrete random variable:

Suppose that $a_j \leq t < a_{j+1}$. Then

$$S(t) = P(T > t) = P(T > a_1, T > a_2, \dots, T > a_j)$$

$$= P(T > a_1) \times P(T > a_2 | T > a_1) \times \cdots$$

$$\cdots \times P(T > a_j | T > a_{j-1})$$

$$= (1 - \lambda_1) \times \cdots \times (1 - \lambda_j)$$

$$= \prod_{k: a_k < t} (1 - \lambda_k)$$

RELATIONSHIPS: AN OVERVIEW

$$f(t)\Delta t pprox P(t \le T < t + \Delta t)$$
 $\lambda(t)\Delta t pprox P(t \le T < t + \Delta t | T \ge t)$
 $S(t) = P(T > t) = \int_{t}^{\infty} f(u)du$
 $f(t) = -\frac{d}{dt}S(t)$
 $\lambda(t) = \frac{f(t)}{S(t)}$
 $\lambda(t) = -\frac{d}{dt}[\log S(t)]$

$$S(t) = e^{-\Lambda(t)}$$

 $S(t) \approx 1 - \Lambda(t)$ while the cumulative hazard is small.

Some parametric survival distributions

The exponential distribution

- Simplest distribution, only one unknown parameter
- Plays a role similar to that of the normal distribution in linear regression

$$f(t) = \lambda e^{-\lambda t}$$
 for $t \ge 0$
 $S(t) = \int_t^\infty f(u) du = e^{-\lambda t}$
 $\lambda(t) = \frac{f(t)}{S(t)} = \lambda$ constant hazard
 $\Lambda(t) = \int_0^t \lambda(u) du = \int_0^t \lambda du = \lambda t$

Some parametric survival distributions...

The Weibull distribution generalizes the exponential, and has two parameters

- λ: the scale parameter
- γ : the *shape* parameter

$$S(t) = e^{-\lambda t^{\gamma}}$$

$$f(t) = \frac{-d}{dt}S(t) = \gamma \lambda t^{\gamma - 1}e^{-\lambda t^{\gamma}}$$

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

$$\Lambda(t) = \int_0^t \lambda(u)du = \lambda t^{\gamma}$$

THE WEIBULL DISTRIBUTION...

The Weibull distribution is convenient because of simple forms. It includes several hazard shapes:

- $\gamma = 1 \rightarrow \text{constant hazard}$
- $0 < \gamma < 1 \rightarrow \text{ decreasing hazard}$
- $\gamma > 1
 ightarrow \, {
 m increasing \ hazard}$