STAT 6390: Analysis of Survival Data

Textbook coverage: Chapter 2

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- Define T as the random variable of the actual (uncensored, untruncated) survival time of an individual.
- We assume the support of T is non-negative or $(0, \infty)$.
- We call T the *random variable* associated with the survival time, and we define T has a cumulative distribution function given by $F(t) = P(T \le t)$.
- The survival function of T is then defined as

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

Why are we more interested in S(t)?

- The *hazard function* is widely used to survival analysis.
- The hazard function h(t) is defined below

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

- $P(t \le T < t + dt | T \ge t)$ is a conditional probability.
- The conditional probability is then expressed as a probability per unit time by dividing by the time interval, dt, to give a rate.
- The function h(t) is also referred to as the hazard rate, the instantaneous death rate, the intensity rate, or the force of mortality.
- Event rate at time t, conditional on the event not having occurred before t.

- In terms of probability, if t is measured in days, h(t) is the approximate probability that an individual, who is at risk of the event occurring at the start of day t, experiences the event during that day.
 - In this case dt = 1.
 - lim_{dt→0} can be thought of as changing the unit from days to hours, minutes, seconds, milliseconds...
- If the event of interest is not death, h(t) can also be regarded as the expected number of events experienced by an individual in unit time, given that the event has not occurred before then.
 - Think of $E\{I(\cdot)\} = P(\cdot)$.
 - The part "given that the event..." might be ignored if events follow the Poisson process.

 The definition in (1) leads to some useful relationships between survival and hazard functions:

$$(1) = \lim_{\mathrm{d}t \to 0} \frac{\mathrm{P}(t \le T < t + \mathrm{d}t)}{\mathrm{d}t \cdot \mathrm{P}(T \ge t)} = \lim_{\mathrm{d}t \to 0} \frac{F(t + \mathrm{d}t) - F(t)}{\mathrm{d}t} \cdot \frac{1}{\mathrm{P}(T \ge t)} = \frac{\mathrm{d}F(t)}{\mathrm{d}t} \cdot \frac{1}{S(t)}.$$

- h(t) is approximately the probability that an individual experiences an event at this instant (t) given that he/she is risk free up to t.
- If T is a continuous random variable, then we have

$$h(t) = \frac{f(t)}{S(t)}. (2)$$

• This shows that from any one of the three functions, f(t), S(t), and h(t), the other two can be determined.

Equation (2) also implies

$$h(t) = -\frac{\mathrm{d}}{\mathrm{d}t} \{ \log S(t) \}$$
 and $S(t) = e^{-H(t)}$,

where $H(t) = \int_0^t h(u) du$ is the cumulative hazard function.

Similarly, the cumulative hazard function can also be obtained from

$$H(t) = -\log S(t).$$

- The cumulative hazard function is the cumulative risk of an event occurring by time t.
- If the event is death, then H(t) summarizes the risk of death up to time t, given that death has not occurred by t.
- If the event is not death, H(t) can be interpreted as the expected number of events that occur in the interval (0, t).

- It is possible for H(t) > 1, h(t) > 1, or f(t) > 1.
- *F*(*t*), *S*(*t*) are bounded in [0, 1].
- F(t) and H(t) are non-decreasing; S(t) is non-increasing.
- h(t) can go up and down.
- For example, suppose $T \sim \exp(\lambda)$, where λ is the rate. Then
 - $S(t) = e^{-\lambda t}$.
 - $h(t) = \lambda$.
 - $H(t) = \lambda t$.

- The S(t) can be estimated non-parametrically with the *product limit* estimator, which is also known as the *Kaplan-Meier* estimator.
- We first assume none of survival times are censored.
- In this case, the survival probability at t, S(t), is defined as

$$\hat{S}_e(t) = \frac{\text{# individuals with survival times } \ge t}{\text{# individuals in the data set}}.$$
 (3)

- Equation (3) is called empirical survival function.
- Similarly, $\hat{F}_e(t) = 1 \hat{S}_e(t)$ is called the *empirical cumulative distribution* function.

- We illustrate with the first 10 uncensored subjects in the whas 100 data.
- Make sure tidyverse package and whas 100 are properly loaded*.

```
> whas10 <- whas100 %>% filter(fstat > 0) %>% filter(row number() <= 10)</pre>
> whas10
# A tibble: 10 x 9
      id admitdate
                                                        age gender
                     foldate
                                   los lenfol fstat
                                                                      bmi
   <int> <fct>
                     <fct>
                                 <int>
                                         <int> <int> <int>
                                                             <int>
                                                                    <db1>
         3/13/1995
                     3/19/1995
                                                         65
                                                                     31.4
                                      4
       2 1/14/1995
                     1/23/1996
                                           374
                     10/4/2001
       3 2/17/1995
                                          2421
                                                                     27.9
       4 4/7/1995
                     7/14/1995
                                                         81
                                                                     21.5
                     5/29/1998
                                                                     30.7
       5 2/9/1995
                                                         78
6
       6 1/16/1995
                     9/11/2000
                                          2065
                                                         82
                                                                     26.5
       7 1/17/1995
                     10/15/1997
                                                         66
       8 11/15/1994 11/24/2000
                                     56
                                                         81
                                                                     28.3
                     2/23/1996
       9 8/18/1995
                                           189
                                                         76
      12 5/26/1995
                     9/29/1996
                                                                     24.7
                                           492
                                                         8.3
```

* see note 1 for details.

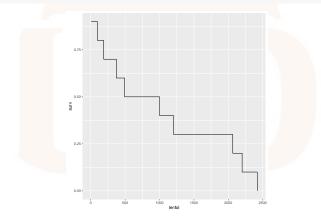
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The empirical estimates can be easily computed with ecdf.

```
whas10 <- whas10 %>% mutate(surv = 1 - ecdf(lenfol)(lenfol))
whas10
A tibble: 10 \times 10
     id admitdate
                    foldate
                                  los lenfol fstat
                                                      age gender
                                                                    bmi
                                                                          surv
  <int> <fct>
                    <fct>
                                       <int> <int> <int>
                                                            <int> <dbl> <dbl>
       3/13/1995
                    3/19/1995
                                                        65
                                                                   31.4 0.9
                                    4
      2 1/14/1995
                    1/23/1996
                                         374
                                                        88
                                                                   22.7 0.6
                    10/4/2001
      3 2/17/1995
                                        2.42.1
                                                                   27.9 0
      4 4/7/1995
                   7/14/1995
                                           98
                                                        81
                                                                   21.5 0.8
      5 2/9/1995
                    5/29/1998
                                                        78
                                                                   30.7 0.3
6
      6 1/16/1995
                    9/11/2000
                                                        82
                                                                   26.5 0.200
      7 1/17/1995
                    10/15/1997
                                                        66
                                                                   35.7 0.4
      8 11/15/1994 11/24/2000
                                                        81
                                                                   28.3 0.100
9
       8/18/1995
                    2/23/1996
                                         189
                                                       76
                                                                   27.1 0.7
     12 5/26/1995
                    9/29/1996
                                         492
                                                        83
                                                                   24.7 0.5
```

• The empirical survival function is a non-increasing step function.

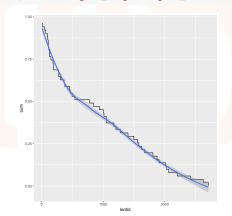
```
> whas10 %>% ggplot(aes(lenfol, surv)) + geom_step(size = 1.2)
```



- The $\hat{S}_e(t)$ is 1 at t=0 and 0 at the final death time.
- The $\hat{S}_e(t)$ is assumed to be constant between adjacent death times.

 Putting everything together, we could plot the empirical survival curve for all the uncensored subjects in whas 100:

```
> whas100 %>% filter(fstat > 0) %>% mutate(surv = 1 - ecdf(lenfol)(lenfol)) %>%
+ gqplot(aes(lenfol, surv)) + geom_step() + geom_smooth()
```



The pipeline between ggplot is "+" instead of "%>%".

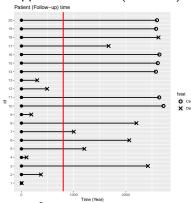
- With censoring, the same idea can be applied with proper adjustment.
- Kaplan-Meier estimator is the default estimator used by many packages.
- The basic idea is to decompose P(T > t) by conditioning on prior times.
- Suppose a sample size of n, P(T > t) can be decomposed as

$$\widehat{S}_{KM}(t) \doteq P(T > t) = P(T > t_{(0)}) \cdot P(T > t_{(1)} | T > t_{(0)}) \cdot P(T > t_{(2)} | T > t_{(1)}) \cdot \dots \cdot P(T > t | T > t_{(i)}),$$

for a series of time intervals $0 \doteq t_{(0)} < t_{(1)} < \ldots < t_{(i)} < t$ for some $i \leq n$.

• In general, the series $\{t_{(1)}, \dots, t_{(m)}\}$ denotes the *m* ordered death times.

Suppose we want P(T > 800) among the first 20 patients in whas 1000.



- There are 6 events before t = 800.
- The events occured at

 t(0) t(1) t(2) t(3) t(4) t(5) t(6)

 0 6 98 189 302 374 492

$$\begin{split} \widehat{S}_{KM}(800) &= P(T > 800) = \\ &= P(T > 0) \times P(T > 6 | T > 0) \times P(T > 98 | T > 6) \times \dots \times P(T > 492 | T > 374) \\ &= 1 \times \frac{19}{20} \times \frac{18}{10} \times \frac{17}{10} \times \frac{16}{17} \times \frac{15}{17} \times \frac{14}{17} = \frac{14}{20} = 70\% \end{split}$$

 $\widehat{S}_{\mathit{KM}}(800) = \widehat{S}_{e}(800)$ here, why?

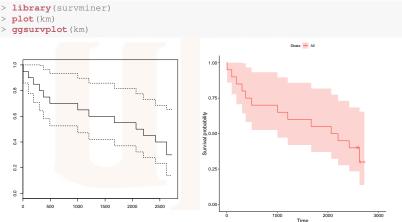


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The Kaplan-Meier estimator can be obtained with the survfit function.

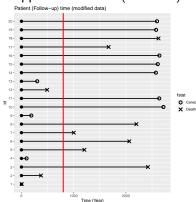
```
> librarv(survival)
> km <- survfit(Surv(lenfol, fstat) ~ 1, data = whas100, subset = id <= 20)
> summary (km)
Call: survfit(formula = Surv(lenfol, fstat) ~ 1, data = whas100, subset = id <=
 time n.risk n.event survival std.err lower 95% CI upper 95% CI
    6
                         0.95
                               0.0487
                                              0.859
                                                           1.000
   98
          19
                         0.90
                              0.0671
                                              0.778
                                                           1.000
  189
         1.8
                         0.85
                               0.0798
                                              0.707
                         0.80 0.0894
                                              0.643
                                                            0.996
  374
          16
                         0.75 0.0968
                                              0.582
                                                           0.966
  492
                         0.70 0.1025
                                              0.525
                                                           0.933
          14
                         0.65
                               0.1067
                                              0.471
                                                           0.897
          13
                         0.60
                               0.1095
                                              0.420
                                                           0.858
1669
                         0.55 0.1112
                                              0.370
                                                           0.818
 2065
          11
                         0.50 0.1118
                                              0.323
                                                           0.775
                         0.45
                               0.1112
                                              0.277
 2421
           9
                         0.40
                                0.1095
                                              0.234
                                                           0.684
 2624
                         0.30
                               0.1194
                                              0.138
                                                            0.654
```

The Kaplan-Meier curve can be plotted with plot or ggsurvplot.



• Since survminer depends on the newest version of survMisc, you might need to update the latter to be able to use ggsurvplot.

Suppose we want P(T > 800) based on the following modified data:



- There are 3 events before t = 800.
- The events occured at $\frac{t_{(0)}}{0}$ $\frac{t_{(1)}}{6}$ $\frac{t_{(2)}}{374}$ $\frac{t_{(3)}}{492}$
- In this modified data, t = 98, 189, 302 are considered as censored.

$$\begin{split} \widehat{S}_{KM}(800) &= P(T > 800) = \\ &= P(T > 0) \times P(T > 6 | T > 0) \times P(T > 374 | T > 6) \times P(T > 492 | T > 374) \\ &= 1 \times \frac{19}{20} \times \frac{15}{16} \times \frac{14}{15} \approx 83.1\% \end{split}$$

The Kaplan-Meier estimator for the whole data is

```
> library(survival)
> km <- survfit(Surv(lenfol, fstat) ~ 1, data = whas100)
> plot (km)
> ggsurvplot (km)
                                                                           Strata - All
                                                     0.75
                                                    probability
05.0
   9.0
                                                     0.25
           500
                         1500
                                2000
                                        2500
                                                                                   2000
                                                                                                 3000
                                                                             Time
```

 If the last observed time corresponds to a censored observation, then the estimate of the survival function does not go to zero.

- Suppose we have a sample of n independent observations $(t_i, c_i), i = 1, 2, ..., n$.
- Suppose there are m deaths and $m \le n$.
- The series $\{t_{(1)}, \ldots, t_{(m)}\}$ are the m ordered death times.
- The Kaplan-Meier estimator has the form

$$\widehat{S}_{KM}(t) = \prod_{t_{(i)} \leq t} \frac{n_i - d_i}{n_i} = \prod_{t_{(i)} \leq t} 1 - \frac{d_i}{n_i},$$

where n_i is the number of individual who are alive at $t_{(i)}$ (at risk), and d_i is the number of individual who died at $t_{(i)}$.

• A potential problem with the Kaplan-Meier estimator is when n_i is small and $n_i = d_i$ occurs at early time.

• An alternative estimate of $\widehat{S}_{KM}(t)$ is the *Nelson-Aalon estimator*.

$$\widehat{S}_{NA}(t) = \prod_{t_{(i)} \le t} \exp\left(-\frac{d_i}{n_i}\right).$$

- The main idea is to see d_i/n_i as the event rate, i.e., $h(t_{(i)}) = d_i/n_i$.
- Recall the relationship h(t) = f(t)/S(t) and think of d_i/n and n_i/n are raw estimates of f(t) and S(t).
- By the similar argument, we have

$$\widehat{H}_{NA}(t) \doteq H(t) = \sum_{t_{(i)} \leq t} d_i/n_i$$
, and $S(t) = e^{-\widehat{H}_{NA}(t)} = \widehat{S}_{NA}(t)$.

- $\widehat{S}_{NA}(t)$ and $\widehat{S}_{KM}(t)$ are derived differently, but both based on d_i and n_i .
- In general $\widehat{S}_{NA}(t) \geq \widehat{S}_{KM}(t)$ but $\widehat{S}_{NA}(t) \approx \widehat{S}_{KM}(t)$.

- $\widehat{S}_{NA}(t)$ has slightly nicer properties and is more stable.
- If the interest is in estimating the cumulative hazard function, H(t), we can use either the $\widehat{H}_{NA}(t)$, or $\widehat{H}_{KM}(t) = -\log \widehat{S}_{KM}(t)$.
- The $\widehat{H}_{KM}(t)$ follows directly form from $\widehat{S}_{KM}(t)$:

$$\widehat{H}_{KM}(t) = -\sum_{t_{(i)} \le t} \log \left(\frac{n_i - d_i}{n_i} \right).$$

Problems with this estimator?

• $\widehat{S}_{NA}(t)$ can be obtained with coxph of the survival package.

```
> args(coxph)
function (formula, data, weights, subset, na.action, init, control,
    ties = c("efron", "breslow", "exact"), singular.ok = TRUE,
    robust = FALSE, model = FALSE, x = FALSE, y = TRUE, tt, method = ties,
    ...)
NULL
```

coxph refers to "Cox proportional hazard model" that has the form

$$h(t) = h_0(t)e^{X^{\top}\beta}, \tag{4}$$

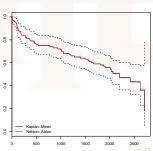
where X is the covariate matrix, β is the regression coefficient, and $h_0(t)$ is called the *baseline hazard* function.

More details will be given in Chapter 3.

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- For now, we will assume $\beta = 0$ in (4), which implies $h(t) = h_0(t)$.
- We will use $h_0(t)$ to obtain $\hat{S}_{NA}(t)$.

```
> cox <- coxph(Surv(lenfol, fstat) ~ 1, data = whas100)
> H0 <- basehaz(cox)
> str(H0)
'data.frame': 95 obs. of 2 variables:
$ hazard: num  0.0201 0.0303 0.0406 0.051 0.0616 ...
$ time : num  6 14 44 62 89 98 104 107 114 123 ...
> plot(km)
> lines(H0$time, exp(-H0$hazard), 's', col = 2)
```



Life-table estimates

- When dataset is large, the $\widehat{S}_{KM}(t)$ and $\widehat{S}_{NA}(t)$ can be obtained with intervals of time, rather than exact time points.
 - The series $\{t_{(1)}, \ldots, t_{(m)}\}$ represents intervals.
 - d_i represents the number of individual who died in $t_{(i)}$.
 - n_i represents the number of individual who are alive in $t_{(i)}$.
- Potential problem with censoring?
- Adjustments under uniform assumption (p25).
- Better to adopt methods for data with interval censoring.

Inference on $\widehat{S}_{KM}(t)$

The 95% confidence interval (CI) does not follow the usual form of

$$PE \pm 1.96 \times SE$$
.

- This is mainly because $\hat{S}_{KM}(t)$ needs to lie between 0 and 1.
- Two common methods to obtain the 95% CI for $\widehat{S}_{KM}(t)$ are the log and log-log transformations.
- The idea is to derive the standard errors on the transformed scale first, then back-transform these back.

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The Delta Method

- We need the Delta method to estimate the standard errors.
- The Delta method states that

$$\operatorname{Var}\{g(X)\} \approx \operatorname{Var}(X) \cdot \{g'(x_0)\}^2$$

where $g'(x_0)$ is the 1st derivative of $g(\cdot)$ evaluates at constant x_0 .

The Delta Method

- A special case of the Delta method is when $g(\cdot) = \log(\cdot)$.
- Setting $g(\cdot) = \log(\cdot)$, we have

$$\operatorname{Var}\{f(X)\}\approx \frac{\operatorname{Var}(X)}{x_0^2}.$$

Inference on $\widehat{S}_{KM}(t)$

- We will first look at the log transformation.
- Recall

$$\widehat{S}_{KM}(t) = \prod_{t_{(i)} \leq t} \frac{n_i - d_i}{n_i}.$$

• The variance of log-transformed $\hat{S}_{KM}(t)$ gives

$$\operatorname{Var}\left\{\log\widehat{S}_{KM}(t)\right\} = \operatorname{Var}\left\{\sum_{t_{(i)} \leq t} \log\left(\frac{n_i - d_i}{n_i}\right)\right\} = \sum_{t_{(i)} \leq t} \operatorname{Var}\left\{\log\left(\frac{n_i - d_i}{n_i}\right)\right\}.$$

- We assume independence between observations in the risk sets.
- For convenience, let's write $p_i = (n_i d_i)/n_i$, and \hat{p}_i when n_i and d_i are known.

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- The key is to estimate Var {log (p_i)} with the Delta method.
- For each $t_{(i)}$, n_i is fixed but d_i is random.
- $n_i d_i$ can be assumed to follow the binomial distribution with parameters n_i and $1 d_i/n_i$. Then

$$\operatorname{Var}(p_i) = \frac{\operatorname{Var}(n_i - d_i)}{n_i^2} = \frac{\frac{d_i}{n_i} \cdot \left(1 - \frac{d_i}{n_i}\right)}{n_i}.$$

With the Delta method, we have

$$\operatorname{Var}\{\log(p_i)\} \approx \frac{\operatorname{Var}(p_i)}{\hat{p}_i} = \frac{d_i}{n_i \cdot (n_i - d_i)}.$$

Inference on $\widehat{S}_{KM}(t)$

From the above result, we have

$$\operatorname{Var}\left\{\log\widehat{S}_{\mathsf{KM}}(t)\right\} \approx \sum_{t_{(i)} \leq t} \frac{d_i}{n_i \cdot (n_i - d_i)}.$$

By the Delta method (again),

$$\operatorname{Var}\left\{\log\widehat{S}_{KM}(t)\right\} \approx \operatorname{Var}\left\{\widehat{S}_{KM}(t)\right\} \cdot \frac{1}{\widehat{S}_{KM}^2(t)}.$$

Altogether, this gives

$$\operatorname{Var}\{\widehat{S}_{KM}(t)\} \approx \widehat{S}_{KM}^{2}(t) \cdot \sum_{t_{(i)} \leq t} \frac{d_{i}}{n_{i} \cdot (n_{i} - d_{i})}.$$

- This result is known as the Greenwood's formula, or the log transformation.
- This estimator can be obtained from a counting process approach.

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- The Greenwood formula is the default method for survfit.
- With the Greenwood formula, the $100(1 \alpha)\%$ condifdence interval of $\widehat{S}_{KM}(t)$ can be obtained using the usual form of PE $\pm Z_{\alpha/2} \times$ SE.
- The bounds can still be outside of [0, 1].
- An alternative approach is to consider the log-log transformation.

By the Delta method, we have

$$\operatorname{Var}\left[\log\left\{-\log\widehat{\widehat{S}}_{\mathsf{KM}}(t)\right\}\right] \approx \frac{1}{\{-\log\widehat{\widehat{S}}_{\mathsf{KM}}(t)\}^2} \cdot \sum_{t_{(i)} \leq t} \frac{d_i}{n_i \cdot (n_i - d_i)}.$$

• This implies that the $100(1-\alpha)\%$ condifidence interval can be constructed by inversing

$$\log\{-\log \widehat{S}_{\mathit{KM}}(t)\} \pm Z_{lpha/2} imes \mathrm{SE}\left[\log\{-\log \widehat{S}_{\mathit{KM}}(t)\}
ight]$$

• Since – log of a survival function gives the cumulative hazard function, e.g., – log S(t) = H(t), the log-log approach called the "log hazard" approach.

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Types of CI can be specified with conf.type in survfit.

```
> ?survfit.coxph
```

• Some options are available for conf . type depending on $g(\cdot)$ used in the Delta method.

plain
$$g(x) = x$$

 $\log g(x) = \log(x)$
 $\log - \log g(x) = \log\{-\log(x)\}$
 $\log it g(x) = \log\left(\frac{x}{1-x}\right)$
 $\arcsin g(x) = \arcsin \sqrt(x)$

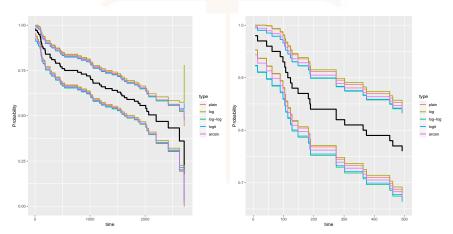
• In addition, Peto et al. (1977) proposed to estimate $\operatorname{Var}\{\widehat{S}_{KM}(t)\}$ from

$$\operatorname{Var}\{\widehat{S}_{KM}(t)\} = \frac{\widehat{S}_{KM}(t) \cdot (1 - \widehat{S}_{KM}(t))}{n_i}$$

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Inference on $\widehat{S}_{KM}(t)$

- The following depict $\widehat{S}_{KM}(t)$ with the five conf.type's for whas 100.
- The CI's are quite close to each other.



*Inference on the median and percentiles

• Given a $\widehat{S}(t)$ ($\widehat{S}_{KM}(t)$ or $\widehat{S}_{NA}(t)$), the estimate of the pth percentile is

$$\hat{t}_p = \min\{t : \hat{S}(t) \le (p/100)\}.$$

- The Delta method can be used again to obtain $Var(\hat{t}_p)$.
- Setting $g(\cdot) = S(\cdot)$, we have the relationship

$$\operatorname{Var}\{\widehat{S}(\hat{t}_p)\} \approx \operatorname{Var}(\hat{t}_p) \cdot \{f(\hat{t}_p)\}^2,$$

where $f(t) = d\hat{S}(t)/dt$.

 The only unknown in the above equation is f(t), which can be approximated by linear interpolation:

$$\hat{f}(\hat{t}_p) \approx \frac{\widehat{S}(\hat{u}_p) - \widehat{S}(\hat{l}_p)}{\hat{l}_p - \hat{u}_p},$$

where $\hat{u}_p < \hat{t}_p < \hat{l}_p$, $\hat{u}_p = \max\{t : \widehat{S}(t) \ge p/100 + \epsilon\}$ and $\hat{l}_p = \min\{t : \widehat{S}(t) \le p/100 - \epsilon\}$, for some small constant ϵ .

*Inference on the median and percentiles

- Replacing the unknown quantities with their empirical estimates, the $100(1-\alpha)\%$ CI can be obtained through $\hat{t}_p \pm Z_{\alpha/2} \times \text{SE}(\hat{t}_p)$.
- Clinicians are specifically interested in median survival (follow-up) time, because survival time data often tend to skewed to the right.
- Other quantity of interest is the semi-interquartiles range (SIQR):

SIQR =
$$\frac{t_{75} - t_{25}}{2}$$
.

 Like the variance, the larger the SIQR, the more dispersed is the survival time distribution.

*Inference on the median and percentiles

Median survival time is printed by survfit.

```
> survfit(Surv(lenfol, fstat) ~ 1, data = whas100)
Call: survfit(formula = Surv(lenfol, fstat) ~ 1, data = whas100)
    n events median 0.95LCL 0.95UCL
    100    51    2201    1806    NA
```

Median follow-up time does not always exists.

A more practical approach?

- As is seen from before, the focus in survival analysis is on observing the occurrence of events over time.
- Such occurrences constitute point (counting) process; counting number of events as they come along.
- There is a very neat collection of theories for counting processes.
- More in-depth details can be found in Fleming and Harrington (2011);
 Kalbfleisch and Prentice (2011).

- Appendix 2 gives a short introduction of counting processes, in the case of right censoring.
- We start by focusing on a single type of event without censoring.
- For a given time t, let N(t) be the number of events over the time period (0, t], then N(t) is a counting process.
- The counting process N(t) is continuous from the right, with jump size 1.

Poisson process

- A well-known example of a counting process is the homogeneous Poisson process.
- Jumps occur randomly and independently of each other (independent increment property).
- A homogeneous Poisson process is described by its *intensity* λ .
- The λ is the probability of occurrence of an event in a small interval divided by the length of the interval.
- We will apply this idea to modeling counting process.

- Suppose the intensity function that describes N(t) is $\lambda(t)$.
- Under our assumption that a subject can experience at most one event, consider a small time interval [t, t + dt), $\lambda(t)$ is

$$\lambda(t)dt = P(dN(t) = 1|past),$$

where dN(t) denotes the # of events in [t, t + dt), or N(t + dt) - N(t).

 Formally speaking, the "past" represents the *filtration* of the process up to but not including time t.

Under our assumption, dN(t) is binary, and

$$\lambda(t)dt = E\{I(dN(t) = 1)|past\} = E\{dN(t)|past\}.$$

Then it follows

$$E(dN(t) - \lambda(t)dt|past) = 0.$$
 (5)

If we define a new process

$$M(t) = N(t) - \int_0^t \lambda(s) ds,$$
 (6)

then we have E(dM(t)|past) = 0.

• It turns out M(t) is a zero-mean martingale.

Definition (6) can be written as

$$dN(t) = \lambda(t)dt + dM(t), \tag{7}$$

reflecting the relationship *observation* = *signal* + *noise*.

- Think of M(t) as the sum of random errors.
- Let

$$\Lambda(t) = \int_0^t \lambda(s) \mathrm{d}s,$$

- (5) and (6) implies $\Lambda(t)$ is the cumulative expected # of events in (0, t], or $\Lambda(t) = \mathbb{E}\{N(t)\}.$
- Λ(t) versus H(t)?

- Now we will look at the scenario when there is only one event type and a subject can experience at most one event (no ties in event times).
- In this case, the counting process $N(t) = I(T \le t)$, where T is the survival time with hazard h(t).
- The N(t) defined above has a jump size 1 at T.
- We then have

$$P(dN(t) = 1|past) = \begin{cases} h(t)dt & \text{if } T \ge t \\ 0 & \text{if } T < t \end{cases} = h(t)I(T \ge t)dt.$$

• The relationship above implies the intensity process $\lambda(t) = h(t) \cdot I(T \ge t)$.

- Building onto our assumption, now assume we have *n* independent subjects, each with T_i , i = 1, ..., n, and hazard $h_i(t)$.
- Suppose $h_i(t) = h(t)$ for all i.
- From the last example, we have

$$N_i(t) = I(T_i \le t)$$
 and $\lambda_i(t) = h_i(t)I(T_i \ge t)$.

 Define the <u>aggregated</u> process by adding together the individual processes:

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

- This process counts the # of individuals who experienced the event by t.
- Is N(t) here a proper counting process?

 Assuming continuous survival times, and the aggregated processs jumps one unit at a time,

$$E(dN(t)|past) = \sum_{i=1}^{n} E(dN_i(t)|past)$$

implies the aggregated intensity satisfies $\lambda(t)dt = \sum_{i=1}^{n} \lambda_i(t)dt$ and

$$\lambda(t) = h(t) Y(t),$$

where $Y(t) = \sum_{i=1}^{n} I(T_i \ge t)$ is the number of individuals at risk right before t, e.g., $Y(t) = n - N(t^-)$.

• Plug $\lambda(t)$ into Equation (7) gives the Nelson-Aalen estimator for H(t).

Derivation of $\widehat{H}_{NA}(t)$

• Plug $\lambda(t)$ into Equation (7), we have

$$dN(t) = h(t)Y(t)dt + dM(t)$$

Intergrating both side after diving by Y(t) gives

$$\int_0^t \frac{1}{Y(t)} \mathrm{d}N(t) - \int_0^t h(t) \mathrm{d}t = \int_0^t \frac{1}{Y(t)} \mathrm{d}M(t)$$

- The right-hand side is a stochastic integral with respect to a zero-mean martingale.
- The first term in the left-hand side gives $\widehat{H}_{NA}(t) = \sum_{t_{(i)} \le t} \frac{1}{n_i}$.
- Then $E\left\{\widehat{H}_{NA}(t)-\widehat{H}(t)\right\}=0$ and $\widehat{H}_{NA}(t)$ is an unbiased estimator of $\widehat{H}(t)$.
- The variance estimator can be derived using the variation processes or with transformation through the Delta methods.

- Consider the same assumptions but allowing ties in event times.
- Two common approaches to deal with tied event times depend on how tied event times arise.
 - 1 Two event times coincide due to rounding.

$$\widehat{H}_{NA}(t) = \sum_{t_{(i)} \le t} \sum_{k=1}^{d_i} \frac{1}{n_i - k}.$$

Event times are genuinely discrete, so that tied event times are real and not due to rounding.

$$\widehat{H}_{NA}(t) = \sum_{t_{(i)} \leq t} \frac{d_i}{n_i}.$$

 More discussion on large number properties with tied event times can be found in Anderson et al. (1993).

- For nonparametric or semi-parametric methods, we do not need to have a distributional assumption on the censoring time, *C*.
- Rather, we will assume independent censoring:

$$P(t \le \tilde{T}_i < t + dt, \Delta_i = 1 | \tilde{T}_i \ge t, past) = P(t \le T_i < t + dt | T_i \ge t),$$

where $\tilde{T}_i = \min(T_i, C_i)$ and $\Delta_i = I(T_i < C_i)$, for i = 1, ..., n.

In the presense of right censoring, the counting process is specified as

$$N_i(t) = I\{\tilde{T}_i \leq t, \Delta_i = 1\}, i = 1, \ldots, n,$$

and $\lambda_i(t)dt = P(dN_i(t) = 1|past)$.

• Applying the independent censoring assumption, the intensity process for $N_i(t)$ takes the form $\lambda_i(t) = h_i(t)Y_i(t)$, where $Y_i(t) = I(\tilde{T}_i \ge t)$ is the at risk indicator for individual i.

As in the uncensored example, the aggregated process is

$$N(t) = \sum_{i=1}^{n} N_i(t) = \sum_{i=1}^{n} I(\tilde{T}_i \leq t, \Delta_i = 1),$$

and the aggregated intensity process is

$$\lambda(t) = \sum_{i=1}^n \lambda_i(t) = \sum_{i=1}^n h_i(t) Y_i(t).$$

- In the case where $h_i(t) = h(t)$ for all i, the intensity process takes the form $\lambda(t) = h(t)Y(t)$, where $Y(t) = \sum_{i=1}^{n} Y_i(t)$.
- The Nelson-Aalen estimator can be derived similarly.

- Counting processes are non-decreasing.
- Martingales associated with M(t) are sub-martingales.
- Doob-Meyer decomposition guarantees the relationship

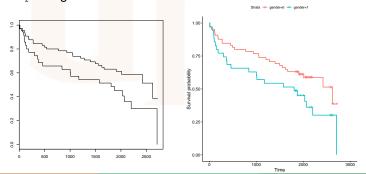
$$N(t) = h(t)Y(t) + M(t).$$

Martingale central limit theorem.

Comparison of survival functions

 The simplest way of comparing the survival functions is to plot them on the same axe.

ggsurvplot gives more informative output.



Comparison of survival functions

- In the whas 100 study, the survival function for female patients (gender
 0) is always greater than that for male patients.
- Two explanations for the observed difference:
 - A real difference between the survival times of the two groups.
 - The difference has been observed is the result of chance variation.
- Procedures such as the hypothesis test is needed to distinguish the two possible explanations.

Hypothesis testing

- As the hypothesis tests commonly encountered, the testing procedure here consists of the following major steps:
 - 1 State the *null hypothesis* and the *alternative hypothesis*.
 - 2 Formulate a *test statistic* that measures the extent to which the observed data depart from the null hypothesis.
 - 3 Calculate the p-value; the probability of obtaining a value as extreme or more extreme than the observed value, when the null hypothesis is true.

Hypothesis testing

- p-value can be interpreted as a measure of the strength of evidence against the null hypothesis.
- A p-value of 0.000 should be interpreted as p < 0.001.
- A one-sided hypothesis test is only appropriate when there is no interest in departures from the null hypothesis in the opposite direction.
- We will look at non-parametric procedures including the log-rank test and the Wilcoxon test.

- If we are willing to assume survival data follow a particular parametric distribution, we can construct a test based on likelihood theory.
- We will derive a nonparametric test using a rank-based procedure.
- Suppose the population survival curves are $S_1(t)$ and $S_0(t)$.
- Ideally, we would like to test $H_o: S_1(t) = S_0(t)$ versus $H_a: S_1(t) > S_0(t)$, $S_1(t) < S_0(t)$, or $S_1(t) \neq S_0(t)$.
- Unfortunately, this is difficult because H_a can take a wide range of forms.
- Instead, it is more feasible to consider the Lehman alternative:

$$H_a: S_1(t) = [S_0(t)]^{\theta}, \text{ for } \theta > 0.$$
 (8)

• The hypothesis in (8) can be view in terms of the hazard functions:

$$H_0: \lambda_1(t) = \lambda_0(t)$$
 versus $H_a: \lambda_1(t) = \theta \lambda_0(t)$.

Either way, we could rewrite the hypothesis test as

$$H_o: \theta = 1 \text{ versus } H_a: \theta > 1, \text{ or } \theta < 1, \text{ or } \theta \neq 1.$$

- One can interpret the results solely based on θ too.
- For example, under H_a : $\theta > 1$, then
 - $S_1(t)$ will be uniformly lower than $S_0(t)$.
 - $h_1(t)$ will be uniformly higher than $h_0(t)$.

- Following the notations defined previously, we assume
 - the two independent random samples are of size N_0 and N_1 .
 - D is the total number of events in the combined sample (size $N_0 + N_1$).
 - $\{t_{(1)}, \ldots, t_{(D)}\}$ is the ordered event times.
 - Let d_{0i} and d_{1i} are the number of events in group 0 and 1, respectively.
 - Let n_{0i} and n_{1i} are the number of at risk in group 0 and 1, respectively.

• For the *i*th event time, we construct the following two-by-two table:

	Group 1	Group 0	Total
Failure	d_{1i}	d_{0i}	di
Non-failure	$n_{1i} - d_{1i}$	$n_{0i} - d_{0i}$	$n_i - d_i$
At risk	n _{1i}	n _{0i}	n _i

- Suppose the numbers of failures in two groups are independent, holding the margins fixed, then the distribution of d_{1i} (or d_{0i}) follows a hypergeometric distribution.
- This gives

$$E(d_{1i}) = \frac{n_{1i} \cdot d_i}{n_i}, \text{ and } Var(d_{1i}) = \frac{n_{1i} \cdot n_{0i} \cdot d_i \cdot (n_i - d_i)}{n_i^2 \cdot (n_i - 1)}.$$

• Using the idea from the χ^2 tests, the test statistic has the form

$$Q = \frac{\left[\sum_{i=1}^{D} \omega_{i} \{d_{1i} - E(d_{1i})\}\right]^{2}}{\sum_{i=1}^{D} \omega_{i}^{2} Var(d_{1i})},$$
(9)

where ω_i is a possibly data-dependent weight.

- When there is no tied failure times, $Var(d_{1i}) = \frac{n_{1i} \cdot n_{0i}}{n_i^2}$.
- Under the null, Q follows the χ^2 distribution with one degree of freedom.
- The log-rank test sets $\omega_i = 1$ (Peto and Peto, 1972).

- Different selections of ω_i were studied.
- Most common expression has the form of

$$\omega_i = \left\{\hat{S}(t_{(i)})\right\}^{\rho}, \text{ for } \rho \in [0, 1],$$

where $\hat{S}(t)$ is the survival estimator from the combined sample, ignoring group.

- This is called the Fleming-Harrington $G(\rho)$ test (e.g. Cox and Oakes, 1984; Fleming et al., 1987).
 - $\rho = 0$ gives the log-rank test
 - $\rho = 1$ gives the Wilcoxon test.
 - $\rho = 0.5$ gives the Tarone and Ware (1977) test.

• The $G(\rho)$ test is implemented in survdiff of the survival package

• The log-rank test shows gender is marginally significant at $\alpha = 0.05$.

- Varying ρ shows similar results.
- In general, when the tests give different results, then more than one result should be reported.
- It is important to also consult the plot to ascertain the directional effect of stratification or treatment.

• As in the Pearson's χ^2 test, a continuity correction could be considered when sample sizes are too small:

$$Q^* = \frac{\left[\sum_{i=1}^{D} \omega_i \{ |d_{1i} - \mathrm{E}(d_{1i})| - 0.5 \} \right]^2}{\sum_{i=1}^{D} \omega_i^2 \mathrm{Var}(d_{1i})}.$$

- The tests discussed here can be extended to comparison more than two survival functions simultaneously.
- K-by-K tables instead of two-by-two tables at each event time.
- When there are many groups to compare, it might be more convenient to consider regression models.

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