HANDOUT #7: CENSORED DATA

- 1. Type I Censoring Fixed Censoring Time
- 2. Type II Censoring Fixed Number of Observed Failures
- 3. Random Censoring
- 4. Many other Types of Censoring
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CENSORED DATA

In some situations, the observations from a population or the outcomes from a process are not a complete set of data. Some of the experimental units on whom we planned to make observations or take measurements are removed from the experiment (study) prior to the end of the experiment. There may be partial information about these experimental units but not complete information.

Suppose we have n randomly selected experimental units from a population whose times to occurrence of an event are to be observed:

- Time to failure of an electrical device
- Time to occurrence of a tumor in a laboratory animal injected with a toxic chemical
- Time until patient's blood pressure reaches a specified level after receiving a treatment.
- Amount of stress at which alloy specimen fractures

Several forms of censoring will now be defined.

Suppose n units are placed on test and we want to measure the variable T_i on unit i = 1, ..., n.

1. No Censoring - Complete Data Set

If we observe T_1, \ldots, T_n on all n units and hence we have a complete data set. Standard methods of estimation are then used to estimate population parameters and make inferences about the population.

2. Right Censoring - Incomplete Data Set

If we observe the value of T_i for $i = 1, \dots, m$ but only know that the remaining n - m units have values of T_i greater than their recorded values then we have right censored data. We only have a lower bound on the value of T_i .

• Type I and Type II censoring are special cases of Right Censoring.

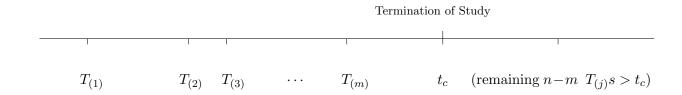
Type I Censoring - Stop Experiment at Time t_c

(The textbook refers to this type of censoring as Type II, see pages 235-236 in textbook)

Suppose n units are placed on test and T_1, T_2, \ldots, T_n are their times to failure. Suppose the reliability study terminates at a preselected time, t_c . All experimental units which have not failed before time t_c have **Type I** censoring.

The experiment has n units on test and the experiment is terminated at a preselected time t_c . Suppose m units have failed before the experiment terminates. Then the ordered times to failure will satisfy:

$$T_{(1)} \le T_{(2)} \le \cdots \le T_{(m)} \le t_c < T_{(m+1)} \le \cdots \le T_{(n)}$$



Because the experiment is terminated at time t_c , we will only observe the times for m of the units. For the remaining n-m units, we will only know that their times satisfy $T_i > t_c$.

In Type I censoring, the time to termination of the experiment, t_c is fixed but the number of observed times, m is random.

Type II Censoring - Stop Experiment When mth Unit Fails

(The textbook refers to this type of censoring as Type I)

An alternatively experimental design has the experiment terminate when the mth (preselected) unit fails. This is referred to as **Type II censoring**.

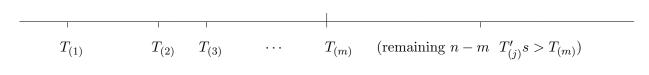
The experiment has n units on test and the experiment is terminated when m units have failed.

The ordered times to failure will satisfy:

$$T_{(1)} \le T_{(2)} \le \dots \le T_{(m)} < T_{(m+1)} \le \dots \le T_{(n)}$$

With the times: $T_{(1)} \leq T_{(2)} \leq \cdots \leq T_{(m)}$ observed but the times for the remaining n-m units unobserved because the experiment was terminated before these units failed.

Termination of Study



The remaining n-m units will not have their failure times recorded. However, we will have the information that $T_{(j)} > T_{(m)}$ for $j = m+1, \ldots, n$.

Type II censoring has the number of observed times, m fixed but the time to termination of the experiment, $T_{(m)}$ random.

3. Left Censoring - Incomplete Data Set

If we observe the value of T_i for $i = 1, \dots, m$ but only know that the remaining n - m units have values of T_i less than their recorded values then we have left censored data. We only have an upper bound on the value of T_i .

• For example, we place n units on test and want to record their time to failure. The units are inspected at specified inspection times and some of the units fail prior to the first inspection time. The failure times of these units would be unknown except we know that they are less than the time of the first inspection, left censoring.

Note, if there are units still functioning after the last inspection time then we would have right censoring.

A second example would be if a measuring device is not sensitive enough to measure observations below a known threshold. For example, if we were measuring ultrasonic noise, the measuring device may not record a value for an sound having frequency less than 20KHz. Thus, any sound having less than 20KHz would not be recorded but we would know that the sound would have a value less than 20KHz.

4. Random Censoring - Incomplete Data

A third type of censoring is Random Censoring in which individual experimental units fail to have their time to the event recorded due to:

- Patient in study just stops returning to medical center
- Machine applying stress to specimens breaks down before specimen fractures
- Patient is removed from study due to side effects of drug
- Operator in Lab stops working
- Emergency budget cuts cause study to be reduced in size resulting in some patients dropping out of study
- Laboratory equipment fails while recording values for a given experimental unit

In this situation, we observe $min\{T_i, C_i\}$, where T_i is the time until event occurs for unit i and C_i is the time at which unit i leaves the experiment(study). Note that Type I censoring is equivalent to random censoring with $C_i \equiv t_c$ for all n units. In most applications, T and C are taken to be independent.

5. Interval Censoring - Incomplete Data Set

- Each of the censored units would have their measured value as being within an interval and no specific value recorded.
- In a failure time study with n units on test, the units are observed at specified inspection times. Therefore, the actual time to failure is unknown. The failure times are recorded as being between two inspection times.

6. Other Types of Censoring or Incomplete Data Set

- Arbitrary Censoring Combinations of left and right censoring and interval censoring with overlapping intervals
- Truncated Data Censoring occurs when there is a bound on all observations for which an exact value is not recorded: lower bound for observations censored on the right, upper bounds for observations censored on the left, and both upper and lower bounds for observations that are interval censored. Truncated data arises when even the existence of a potential observation would be unknown if its value were to lie in a certain range of values, either below a specified point τ_L or above a specified value, τ_U .

Example from Statistical Methods for Reliability Data, Meeker & Escobar Ultrasonic inspection is used to detect flaws in titanium alloys. Ultrasonic signal amplitude is positively correlated with crack size. Titanium grain boundaries reflect signals as well as flaws. Thus below a specified threshold, τ_L it is impossible to be sure whether a signal is a flaw or grain boundary.

In a lab test of the inspection process, specimens with flaws of known size are inspected. The signal's amplitude is measured only when it is above τ_L . All specimens in which a signal is not recorded would have left-censored observations.

However, in a production inspection process, a flaw is not detected when the signal's amplitude is below τ_L . Thus the number of flaws that are present with signal amplitude below τ_L is unknown. The signal amplitudes recorded above τ_L are known as left-truncated observations, or observations from a left-truncated distribution.

Form of Censored Data

The data will have the following structure in each of four situations:

- Let T_1, \ldots, T_n be the times to the event with T_1, \ldots, T_n iid having cdf F.

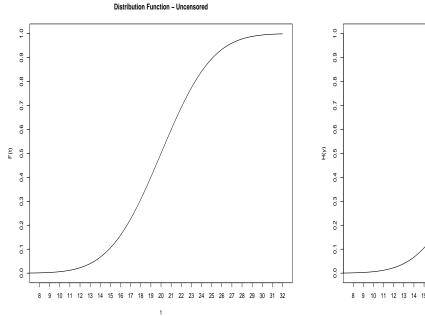
 The values of T_1, \ldots, T_n may or may not be observed depending on whether or not they have been censored.
- Let Y_1, \ldots, Y_n be the observed data with Y_1, \ldots, Y_n iid having cdf H.

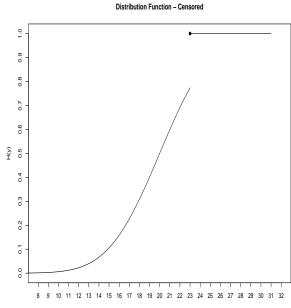
The form of Y_i and its cdf H will depend on the type of censoring in the experiment:

- 1. No Censoring: $Y_i = T_i$ for i = 1, ..., n and $Y_1, ..., Y_n$ iid having cdf H = F.
- 2. Type I Censoring: For all $T_i > t_c$, data is censored, therefore we have

$$Y_i = \begin{cases} T_i & \text{if } T_i \le t_c \\ t_c & \text{if } T_i > t_c \end{cases}$$

The cdf of Y_i , H has a jump at t_c of height p where $p = P[Y_i = t_c] = P[T_i > t_c] = 1 - F(t_c) \neq 0$. This occurs because the distribution of Y_i is truncated at t_c .





3. **Type II Censoring:** The first m times to the event are observed and then the experiment is terminated. With $T_{(1)} \leq T_{(2)} \leq \cdots \leq T_{(n)}$, we observe $T_{(1)} \leq T_{(2)} \leq \cdots \leq T_{(m)}$ and then experiment is then terminated. Thus, the largest n-m times to event are not observed. Therefore we have

$$Y_{(i)} = \begin{cases} T_{(i)} & \text{for } i = 1, \dots, m \\ T_{(m)} & \text{for } i = m + 1, \dots, n \end{cases}$$

Using the properties of the order statistics, the joint pdf of the Y_i is

$$f(y_1, y_2, \dots, y_n) = \frac{n!}{(n-m)!} f(y_1) f(y_2) \cdots f(y_m) [S(y_m)]^{n-m}$$

where S(y) = 1 - F(y) is the survival function.

4. Random Censoring: Let C_1, \ldots, C_n be the times at which the n units were censored with $C_i's$ iid with cdf $G(\cdot)$. Let T_1, \ldots, T_n be the times to the occurrence of the event for the n units with $T_i's$ iid with cdf $F(\cdot)$. We observe $(Y_1, \delta_1), (Y_2, \delta_2), \ldots, (Y_n, \delta_n)$ where $Y_i = min(T_i, C_i)$ and

$$\delta_i = I(T_i \leq C_i) = \left\{ \begin{array}{ll} 1 & \text{if Unit i is NOT censored} \\ 0 & \text{if Unit i IS censored} \end{array} \right.$$

With random censoring Y_1, \ldots, Y_n are iid with cdf $H(\cdot)$.

The joint pdf of (Y, δ) is obtained using the independence of T and C:

If $\delta = 1$ (uncensored observation), then $T \leq C$ and Y = T, which yields

$$h(y, \delta) = f(y)P[C \ge y] = f(y)[1 - G(y)]$$

If $\delta = 0$ (censored observation), then $T \geq C$ and Y = C, which yields

$$h(y, \delta) = g(y)P[T \ge y] = g(y)[1 - F(y)] = g(y)S(y)$$

Using δ , we can combine the two situations into a single equation:

$$h(y, \delta) = [f(y)]^{\delta} [1 - G(y)]^{\delta} [g(y)]^{1 - \delta} [S(y)]^{1 - \delta}$$

For data having Type I, Type II, or Random Censoring our goal is to estimate the cdf F and parameters associated with F using the observed data Y_1, \ldots, Y_n .

Parametric Estimation

Suppose we know the family of distributions for which F is a member. For example, suppose the family is a logistic family of distributions with parameters (θ_1, θ_2) , that is, the family of distributions has pdf,

$$f(t; \theta_1, \ \theta_2) = \frac{1}{\theta_2} \frac{e^{-(t-\theta_1)/\theta_2}}{[1 + e^{-(t-\theta_1)/\theta_2}]^2}$$

We want to estimate the parameters associated with F, θ_1 and θ_2 . In order to use MLE techniques we need to specify the likelihood function for the data. The form of the likelihood will depend on the type of censoring:

1. No Censoring:

Let $f(\cdot, \boldsymbol{\theta})$ be the pdf associated with $F(\cdot, \boldsymbol{\theta})$. The likelihood function for the observed data $Y_i's$ is given by

$$L(\boldsymbol{\theta}) = f(Y_1, Y_2, \dots, Y_n; \boldsymbol{\theta}) = \prod_{i=1}^n f(Y_i; \boldsymbol{\theta}) = \prod_{i=1}^n f(T_i; \boldsymbol{\theta})$$

Select the value of $\boldsymbol{\theta}$ to maximize $L(\boldsymbol{\theta})$.

2. Random Censoring

Assume T_i and C_i are independent with $Y_i = min(T_i, C_i)$ and F the cdf of $T_i's$, H the cdf of $Y_i's$, and G the cdf of $C_i's$.

(2a.) If Y has a discrete distribution, then the joint pdf for the pair (Y_i, δ_i) is given by

$$P[Y_i = y, \delta = 1] = P[Y_i = y, C_i \ge T_i] = P[T_i = y, C_i \ge y] = P[T_i = y] P[C_i \ge y] = f(y) [1 - G(y)]$$

$$P[Y_i = y, \delta = 0] = P[Y_i = y, T_i \ge C_i] = P[C_i = y, T_i \ge y] = P[C_i = y] P[T_i \ge y] = g(y) [1 - F(y)]$$

(2b.) For continuous Y, the joint pdf for the pair (Y_i, δ_i) is given by

$$L(Y_i, \delta_i); \boldsymbol{\theta}) = \begin{cases} f(Y_i; \boldsymbol{\theta}) P[C_i \ge T_i] & \text{if } \delta_i = 1 \text{ (uncensored)} \\ g(Y_i) P[C_i < T_i] & \text{if } \delta_i = 0 \text{ (censored)} \end{cases}$$

$$L(Y_i, \delta_i); \boldsymbol{\theta}) = \begin{cases} f(Y_i; \boldsymbol{\theta})[1 - G(Y_i)] & \text{if } \delta_i = 1 \text{ (uncensored)} \\ g(Y_i)[1 - F(Y_i; \boldsymbol{\theta})] & \text{if } \delta_i = 0 \text{ (censored)} \end{cases}$$

$$L(Y_i, \delta_i); \boldsymbol{\theta}) = [f(Y_i; \boldsymbol{\theta})]^{\delta_i} [S(Y_i; \boldsymbol{\theta})]^{(1 - \delta_i)} [1 - G(Y_i)]^{\delta_i} [g(Y_i)]^{(1 - \delta_i)},$$

where $S(Y_i; \boldsymbol{\theta}) = 1 - F(Y_i; \boldsymbol{\theta})$ is the survival function for T_i .

In the likelihood for the full sample we will designate terms involving $[1 - G(Y_i)]$ and $g(Y_i)$ as $K(Y_i, \delta_i)$ because they do not involve the unknown parameters $\boldsymbol{\theta}$. Hence, the likelihood function is given by

$$L(\boldsymbol{\theta}) = \prod_{i=1}^{n} L(Y_i, \delta_i; \boldsymbol{\theta}) = \prod_{i=1}^{n} [f(Y_i; \boldsymbol{\theta})]^{\delta_i} [S(Y_i; \boldsymbol{\theta})]^{(1-\delta_i)} \prod_{i=1}^{n} K(Y_i, \delta_i)$$

which yields

$$L(\boldsymbol{\theta}) = \prod_{i \in U} f(T_i; \boldsymbol{\theta}) \prod_{i \in C} S(C_i; \boldsymbol{\theta}) \left[\prod_{i=1}^n K(Y_i, \delta_i) \right]$$

where U is the set of indices for the uncensored units and C is the set of indices for the censored units.

3. Type I Censoring

The likelihood is obtained from the Random censoring case with $C_i \equiv t_c$ for all $i \in C$:

$$L(\boldsymbol{\theta}) = \prod_{i \in U} f(T_i; \boldsymbol{\theta}) \prod_{i \in C} S(C_i; \boldsymbol{\theta}) \prod_{i=1}^n K(Y_i, \delta_i) = [S(t_c; \boldsymbol{\theta})]^{n-n_U} \prod_{i \in U} f(T_i; \boldsymbol{\theta}) \left[\prod_{i=1}^n K(Y_i, \delta_i) \right]$$

4. Type II Censoring

Observe $T_{(1)} \leq T_{(2)} \leq \ldots \leq T_{(m)}$ with remaining n-m units having $Y_i \equiv T_{(m)}$. Therefore, the likelihood is given by

$$L(\boldsymbol{\theta}) = \binom{n}{m} \left[m! \prod_{i=1}^{m} f(Y_{(i)}; \boldsymbol{\theta}) \right] [S(Y_{(m)}, \boldsymbol{\theta})]^{n-m} \left[\prod_{i=1}^{n} K(Y_{i}, \delta_{i}) \right]$$
$$= \frac{n!}{(n-m)!} \left[\prod_{i=1}^{m} f(Y_{(i)}; \boldsymbol{\theta}) \right] [S(Y_{(m)}, \boldsymbol{\theta})]^{n-m} \left[\prod_{i=1}^{n} K(Y_{i}, \delta_{i}) \right]$$

Once we have established the likelihood function for the data, we then select $\hat{\boldsymbol{\theta}}$ to maximize $L(\boldsymbol{\theta})$ the likelihood function.

Example Let T_1, \ldots, T_n be iid exponential: $F(t; \beta) = 1 - e^{-t/\beta}$

$$\Rightarrow f(t; \beta) = \frac{1}{\beta} e^{-t/\beta}$$
 and $S(t; \beta) = 1 - F(t; \beta) = e^{-t/\beta}$

Suppose we have **random censoring** and let n_U be the number of uncensored observations. The likelihood function is given by

$$L(\boldsymbol{\theta}) = \prod_{i \in U} f(T_i; \boldsymbol{\theta}) \prod_{i \in C} S(C_i; \boldsymbol{\theta}) \left[\prod_{i=1}^n K(Y_i, \delta_i) \right]$$

$$= \prod_{i \in U} \frac{1}{\beta} e^{-T_i/\beta} \prod_{i \in C} e^{-C_i/\beta} \left[\prod_{i=1}^n K(Y_i, \delta_i) \right]$$

$$= (\beta)^{-n_U} e^{-\frac{1}{\beta} \left(\sum_{i \in U} T_i + \sum_{i \in C} C_i \right)} \left[\prod_{i=1}^n K(Y_i, \delta_i) \right]$$

$$= (\beta)^{-n_U} e^{-\frac{1}{\beta} \sum_{i=1}^n Y_i} \left[\prod_{i=1}^n K(Y_i, \delta_i) \right]$$

Thus, the log-likelihood is given by

$$\begin{split} l(\beta) &= log(L(\beta)) \; = \; -n_U log(\beta) \; - \; \frac{1}{\beta} \sum_{i=1}^n \; Y_i \; + \; log \left[\prod_{i=1}^n K(Y_i, \delta_i) \right] \quad \Rightarrow \\ & \frac{d}{d\beta} l(\beta) = \frac{-n_U}{\beta} \; + \; \frac{1}{\beta^2} \sum_{i=1}^n Y_i \; + \; 0 \\ & \frac{d}{d\beta} l(\beta)|_{\beta = \hat{\beta}} = 0 \quad \Rightarrow \quad \hat{\beta} = \frac{1}{n_U} \sum_{i=1}^n Y_i \quad \Rightarrow \\ & \hat{\beta} = \frac{\text{Total Time n Units were Operating}}{\text{Total Number of Units Which Observed Event}} \end{split}$$

The same estimator is obtained for Type I censoring.

For Type II censoring, the estimator becomes

$$\hat{\beta} = \frac{1}{n_U} \sum_{i=1}^{n} Y_i = \frac{1}{n_U} \left[\sum_{i=1}^{n_U} T_{(i)} + (n - n_U) T_{(n_U)} \right]$$

 $\hat{\beta}$ is referred to as the Winsorized Mean - replace all extreme values with the largest(smallest) non-extreme value.

Note that the naive estimator, $\hat{\beta}^*$ would satisfy

$$\hat{\beta}^* = \frac{1}{n} \sum_{i=1}^n Y_i = \frac{1}{n} \sum_{i \in U} T_i + \frac{1}{n} \sum_{i \in C} C_i \le \frac{1}{n} \sum_{i=1}^n T_i = \hat{\beta}.$$

Thus, $\hat{\beta}^*$ would on the average underestimate β , because $\hat{\beta}$ is an unbiased estimator of β which would imply

$$E\left[\hat{\beta}^*\right] \leq E\left[\hat{\beta}\right] = \beta$$

Example Let T_1, \ldots, T_n be iid with a Weibull distribution: $F(t; \gamma, \alpha) = 1 - e^{-(t/\alpha)^{\gamma}}$

$$\Rightarrow f(t; \gamma, \alpha) = \frac{\gamma}{\alpha} (t/\alpha)^{\gamma - 1} e^{-(t/\alpha)^{\gamma}} \text{ and } S(t; \gamma, \alpha) = 1 - F(t; \gamma, \alpha) = e^{-(t/\alpha)^{\gamma}}$$

Suppose we have **Type I censoring** and let $m = n_U$ be the number of uncensored observations. The likelihood function is given by

$$L(\boldsymbol{\theta}) = \prod_{i \in U} f(T_i; \boldsymbol{\theta}) \prod_{i \in C} S(C_i; \boldsymbol{\theta}) \left[\prod_{i=1}^n K(Y_i, \delta_i) \right]$$

$$= \prod_{i \in U} \frac{\gamma}{\alpha} (T_i/\alpha)^{\gamma - 1} e^{-(T_i/\alpha)^{\gamma}} \prod_{i \in C} \left[e^{-(t_c/\alpha)^{\gamma}} \right] \left[\prod_{i=1}^n K(Y_i, \delta_i) \right]$$

$$= \prod_{i \in U} \frac{\gamma}{\alpha} (T_i/\alpha)^{\gamma - 1} e^{-(T_i/\alpha)^{\gamma}} \left[e^{-(t_c/\alpha)^{\gamma}} \right]^{n - m} \left[\prod_{i=1}^n K(Y_i, \delta_i) \right]$$

$$= \left(\frac{\gamma}{\alpha} \right)^m \left(\frac{1}{\alpha^{\gamma - 1}} \right)^m \left(\prod_{i \in U} T_i^{\gamma - 1} \right) \left(e^{-\frac{1}{\alpha^{\gamma}} \sum_{i \in U} T_i^{\gamma}} \right) \left(e^{-(n - m)(t_c/\alpha)^{\gamma}} \right) \left[\prod_{i=1}^n K(Y_i, \delta_i) \right]$$

The log-likelihood function is given by

$$l(\gamma, \beta) = log(L(\gamma, \beta))$$

$$\begin{split} l(\gamma,\beta) &= log(L(\gamma,\beta)) \\ &= mlog(\gamma) - m\gamma log(\alpha) + (\gamma-1) \sum_{i \in U} log(T_i) - \frac{1}{\alpha^{\gamma}} \sum_{i \in U} T_i^{\gamma} - \frac{(n-m)}{\alpha^{\gamma}} t_c^{\gamma} \ + \ log \left[\prod_{i=1}^n K(Y_i,\delta_i) \right] \\ &= mlog(\gamma) - m\gamma log(\alpha) + (\gamma-1) \sum_{i \in U} log(Y_i) - \frac{1}{\alpha^{\gamma}} \left(\sum_{i \in U} T_i^{\gamma} + (n-m) t_c^{\gamma} \right) \ + \ log \left[\prod_{i=1}^n K(Y_i,\delta_i) \right] \\ &= mlog(\gamma) - m\gamma log(\alpha) + (\gamma-1) \sum_{i \in U} log(Y_i) - \frac{1}{\alpha^{\gamma}} \left(\sum_{i=1}^n Y_i^{\gamma} \right) \ + \ log \left[\prod_{i=1}^n K(Y_i,\delta_i) \right] \end{split}$$

Taking derivatives of $l(\gamma, \alpha)$ wrt γ and α yield,

$$\begin{split} \frac{d}{d\alpha}l(\gamma,\alpha) &= \frac{d}{d\alpha}\left(mlog(\gamma) - m\gamma log(\alpha) + (\gamma - 1)\sum_{i \in U}log(Y_i) - \frac{1}{\alpha^{\gamma}}\left(\sum_{i=1}^n Y_i^{\gamma}\right) + log\left[\prod_{i=1}^n K(Y_i,\delta_i)\right]\right) \\ &= 0 - \frac{m\gamma}{\alpha} + 0 + \frac{\gamma}{\alpha^{\gamma+1}}\sum_{i=1}^n Y_i^{\gamma} + 0 \end{split}$$

$$\frac{d}{d\gamma}l(\gamma,\alpha) = \frac{d}{d\gamma}\left(mlog(\gamma) - m\gamma log(\alpha) + (\gamma - 1)\sum_{i \in U} log(Y_i) - \frac{1}{\alpha^{\gamma}}\left(\sum_{i=1}^{n} Y_i^{\gamma}\right) + log\left[\prod_{i=1}^{n} K(Y_i, \delta_i)\right]\right)$$

$$= \frac{m}{\gamma} - mlog(\alpha) + \sum_{i \in U} log(Y_i) - \frac{1}{\alpha^{\gamma}}\sum_{i=1}^{n} log(Y_i)Y_i^{\gamma} + \frac{1}{\alpha^{\gamma}}log(\alpha)\left(\sum_{i=1}^{n} Y_i^{\gamma}\right) + 0$$

Then setting the derivatives equal to 0 yield the following equations:

$$\hat{\alpha} = \left(\frac{1}{m} \sum_{i=1}^{n} Y_i^{\hat{\gamma}}\right)^{1/\hat{\gamma}} \tag{1}$$

$$0 = \frac{1}{\hat{\gamma}} + \frac{1}{m} \sum_{i \in U} log(Y_i) - \frac{\sum_{i=1}^{n} Y_i^{\hat{\gamma}} log(Y_i)}{\sum_{i=1}^{n} Y_i^{\hat{\gamma}}}$$
(2)

Notice that equation (2) involves just $\hat{\gamma}$.

A numerical solution can be easily obtained and then the value of $\hat{\alpha}$ can be obtained from equation (1).

A numerical example will be considered next to illustrate the use of SAS and R in obtaining estimators of γ and α in the Weibull model when the data has censoring.

EXAMPLE The following example (slightly modified) is from *Statistical Analysis of Reliability Data* by M.J. Crowder, A.C. Kimber, R.L. Smith, and T.J. Sweeting. In an experiment to determine the strength of a braided cord after weathering, the strengths of 48 pieces of cord that had been weathered for a specified length of time were investigated. The company wanted to estimate the probability that the cord would have strength of at least 53, that is, estimate S(53). Seven cords were damaged during the study which resulted in a decrease in their strength. Therefore, the study produced right censored strength values. The strengths of the remaining 41 cords were determined as shown below:

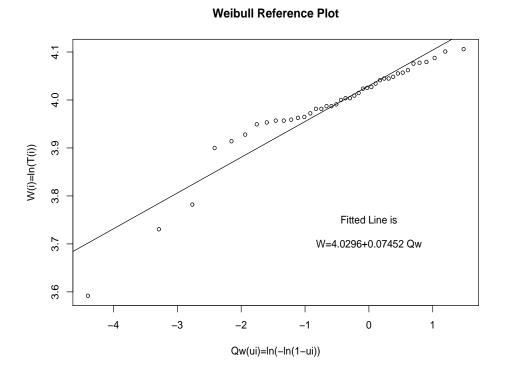
```
36.3
      52.4
             54.8
                    57.1
                          60.7
                                 41.7
                                        52.6
                                               54.8
43.9
      52.7
             55.1
                    57.7
                           49.4
                                 53.1
                                        55.4
                                               57.8
             55.9
50.1
      53.6
                    58.1
                          50.8
                                 53.6
                                        56.0
                                               58.9
51.9
      53.9
             56.1
                    59.0
                          52.1
                                 53.9
                                        56.5
                                               59.1
52.3
      54.1
             56.9
                    59.6
                          52.3
                                 54.6
                                        57.1
                                               60.4
```

The 7 censored strength values from the damaged cords are given next:

```
26.8 29.6 33.4 35.0 40.0 41.9 42.5
```

The true strength values of the 7 cords, T_i are unobservable but we know $T_i > Y_i$, where Y_i are the observed values.

A Weibull reference distribution plot for the 41 uncensored strengths is displayed here:



From the plot, it would appear that the Weibull model is adequate.

The following SAS program will be used to obtain the MLE's of α and γ :

```
*weib_mle_censored.sas in Files/SASCode;
```

```
option ls=75 ps=55 nocenter nodate;
title 'Strength of Braided Cord';
data cords;
input S C @@;
label S = 'Strength of Cord' C = 'Censoring (1=Yes)';
cards;
36.3 0 52.4 0 54.8 0 57.1 0 60.7 0 41.7 0 52.6 0 54.8 0 57.3 0
43.9 0 52.7 0 55.1 0 57.7 0 49.4 0 53.1 0 55.4 0 57.8 0
50.1 0 53.6 0 55.9 0 58.1 0 50.8 0 53.6 0 56.0 0 58.9 0
51.9 0 53.9 0 56.1 0 59.0 0 52.1 0 53.9 0 56.5 0 59.1 0
52.3 0 54.1 0 56.9 0 59.6 0 52.3 0 54.6 0 57.1 0 60.4 0
26.8 1 29.6 1 33.4 1 35.0 1 40.0 1 41.9 1 42.5 1
run;
proc lifereg data=cords;
model S*C(1) = /dist=weibull covb;
run;
```

OUTPUT from SAS Program:

The LIFEREG Procedure in SAS

Number of Observations	48
Noncensored Values	41
Right Censored Values	7
Name of Distribution	Weibull

		Standard	95% Con	fidence	Chi-	
DF	${\tt Estimate}$	Error	Lim	its	Square I	Pr > ChiSq
1	4.0257	0.0100	4.0061	4.0454	161324	<.0001
1	0.0615	0.0077	0.0481	0.0786		
1	56.0223	0.5615	54.9325	57.1337		
1	16.2591	2.0363	12.7202	20.7827		
	1 1 1	1 4.0257 1 0.0615 1 56.0223	DF Estimate Error 1 4.0257 0.0100 1 0.0615 0.0077 1 56.0223 0.5615	DF Estimate Error Lim 1 4.0257 0.0100 4.0061 1 0.0615 0.0077 0.0481 1 56.0223 0.5615 54.9325	DF Estimate Error Limits	DF Estimate Error Limits Square I 1 4.0257 0.0100 4.0061 4.0454 161324 1 0.0615 0.0077 0.0481 0.0786 1 56.0223 0.5615 54.9325 57.1337

The estimators of γ and α are obtained by recalling, $F(t) = 1 - e^{-(t/\alpha)^{\gamma}}$.

Thus, α is the Weibull Scale parameter and γ is the Weibull Shape parameter.

$$\hat{\gamma}$$
 = Weibull Shape = 16.2591; $\hat{\alpha}$ = Weibull Scale = 56.0223

With $S(t) = e^{-(t/\alpha)^{\gamma}}$, the estimate of S(53) would be given by:

$$\widehat{S}(53) = e^{-(53/\widehat{\alpha})^{\widehat{\gamma}}} = e^{-(53/56.0223)^{16.2591}} = .6664$$

Thus, we would estimate that approximately 67% of the cords would have strength of at least 53

From the uncensored data, we have 28 of the 41 values are greater than 53, that is, 68.3%. The two values are very close.

The estimated standard error of the estimator (see STAT 611) is obtained from the inverse of the information matrix: $\widehat{SE}(\hat{S}(53)) = 0.046$

We can use the following R code to obtain the estimates of α and γ for a Weibull model with censored data:

```
R Code in Dostat under the name cords_censoredMLE.R
library(MASS)
library(survival)
st = c(36.3, 52.4, 54.8, 57.1, 60.7, 41.7, 52.6, 54.8, 57.3,
      43.9, 52.7, 55.1, 57.7, 49.4, 53.1, 55.4, 57.8, 50.1,
      53.6, 55.9, 58.1, 50.8, 53.6, 56.0, 58.9, 51.9, 53.9,
      56.1, 59.0, 52.1, 53.9, 56.5, 59.1, 52.3, 54.1, 56.9,
      59.6, 52.3, 54.6, 57.1, 60.4,
     26.8, 29.6, 33.4, 35.0, 40.0, 41.9, 42.5)
0,0,0,0,0,0,0)
cords = survreg(Surv(st, stcens) ~ 1, dist='weibull')
summary(cords)
fitdistr(st,"weibull")
OUTPUT from R Code:
survreg(formula = Surv(st, stcens) ~ 1, dist = "weibull")
          Value Std. Error z
(Intercept) 4.03 0.0100 401.7 0.00e+00
Log(scale) -2.79 0.1252 -22.3 7.82e-110
Scale= 0.0615
Weibull distribution
Loglik(model) = -115.9
                   Loglik(intercept only) = -115.9
Number of Newton-Raphson Iterations: 11
n=48
# MLE's ignoring censoring:
    shape
              scale
  9.4232284 54.5099326
(1.1920947) (0.8643678)
```

From the above R function we must reinterpret the parameters to obtain:

$$\hat{\gamma} = 1/\text{Scale} = 1/.0615 = 16.2602$$

$$\hat{\alpha} = e^{\text{Intercept}} = e^{4.03} = 56.2609$$

and

$$\widehat{S}(53) = e^{-(53/\hat{\alpha})^{\hat{\gamma}}} = e^{-(53/56.2609)^{16.2602}} = .6847$$

 $\widehat{S}(53) = .6664$ using SAS code.

The values are slightly different possibly due to round-off error or precision differences in the calculations.

If the censoring of seven values had been ignored, the mle's based on 48 uncensored values are

$$\hat{\gamma} = 9.4232$$

$$\hat{\alpha} = 54.5099$$

and

$$\widehat{S}(53) = e^{-(53/\hat{\alpha})^{\hat{\gamma}}} = e^{-(53/54.5099)^{9.4232}} = .4642$$

A substantial change in the estimated proportion of cords that would have strength greater than 53 units.

Thus confirming the importance of taking into account whether or not a data value is censored.

Distribution-Free Estimators

Suppose it is **unknown** to which family of distributions the cdf of T_i belongs. We want to use the data to estimate the survival function: S(t) = Pr[T > t] and parameters associated with the survival function, such as, μ , $\tilde{\mu}$, and σ .

• No Censoring

1. Estimator of μ based on $\hat{S}(t)$

With no censoring, $\hat{S}(t) = \frac{1}{n} \sum_{i=1}^{n} I(T_i > t) = \text{proportion of } T_i's \text{ greater than } t.$

$$E[T] = \mu = \int_{0}^{\infty} t dF(t) = tF(t)|_{0}^{\infty} - \int_{0}^{\infty} F(t) dt = \int_{0}^{\infty} [1 - F(t)] dt = \int_{0}^{\infty} S(t) dt \Rightarrow$$

$$\hat{\mu} = \int_{0}^{\infty} \hat{S}(t)dt = \frac{1}{n} \sum_{i=1}^{n} \int_{0}^{\infty} I(T_i \ge t)dt = \frac{1}{n} \sum_{i=1}^{n} \int_{0}^{T_i} dt = \frac{1}{n} \sum_{i=1}^{n} T_i = \bar{T}$$

2. Estimator of median, M based on $\hat{S}(t)$

Recall, S(t) = 1 - F(t), so we have

$$Q(u) = \inf(t: F(t) \ge u) = \inf(t: S(t) \le 1 - u)$$

Therefore, the median is defined by

$$M = Q(.5) = inf(t: F(t) > .5) = inf(t: S(t) < .5)$$

Thus, we can estimate the median by

$$\hat{M} = \inf\{t : \hat{S}(t) < .5\}$$

In general the pth quantile is estimated by

$$\widehat{Q}(p) = in f(t : \widehat{S}(t) < 1 - p)$$

Kaplan-Meier Product Limit Estimator of S(t)

Suppose we have n homogeneous units placed on test.

Let $t_1 < t_2 < \cdots < t_k$ be the times at which units fail

Let d_j units fail at time t_j for j = 1, 2, ..., k

Let m_j be the number of units that are censored in the interval $[t_j, t_{j+1})$ for j = 0, 1, ..., k, with $t_o = 0$ and $t_{k+1} = \infty$

Let $n_j = (m_j + d_j) + (m_{j+1} + d_{j+1}) + \cdots + (m_k + d_k)$ be number of units at risk for failure just prior to time t_j

Note: $n_1 = n$, $n_2 = n - (d_1 + m_1)$

The Kaplan-Meier product limit estimator of S(t) = P[T > t], where T is the time to failure, is given by

For
$$t \in [t_i, t_{i+1})$$
, $\widehat{S}(t) = \prod_{j=1}^{i} \frac{n_j - d_j}{n_j}$

The estimator $\widehat{S}(t)$ is undefined for $t > t_k$

An heuristic justification of the estimator is as follows.

For $t \in [0, t_1)$, $S(t) = P[T > t] = P[\text{unit survives beyond time t}] \Rightarrow$

 $\widehat{S}(t) = 1$ because there are no failures in $[0, t_1)$

For $t \in [t_1, t_2)$,

 $S(t) = P[T > t] = P[\text{unit survives in } [t_1, t] \mid \text{survives in } [0, t_1)] P[\text{unit survives in } [0, t_1)] \Rightarrow$

$$\widehat{S}(t) = \left(\frac{n_1 - d_1}{n_1}\right) \cdot 1$$

For $t \in [t_2, t_3)$,

 $S(t) = P[T > t] = P[\text{unit survives in } [t_1, t_1] \mid \text{survives in } [t_1, t_2)] P[\text{unit survives in } [t_1, t_2)] \Rightarrow$

$$\widehat{S}(t) = \left(\frac{n_2 - d_2}{n_2}\right) \left(\frac{n_1 - d_1}{n_1}\right)$$

For $t \in [t_3, t_4)$,

 $S(t) = P[T > t] = P[\text{unit survives in } [t_3, t] \mid \text{survives in } [t_2, t_3)] P[\text{unit survives in } [t_2, t_3)] \Rightarrow$

$$\widehat{S}(t) = \left(\frac{n_3 - d_3}{n_3}\right) \left[\left(\frac{n_2 - d_2}{n_2}\right) \left(\frac{n_1 - d_1}{n_1}\right) \right]$$

From the Kaplan-Meier estimator we can then obtain estimators of the mean and median similarly as was done for uncensored data:

$$\hat{\mu} = \int_{0}^{\infty} \hat{S}(t)dt \Rightarrow$$

$$\hat{\mu} = \text{area under } \hat{S}(\cdot) \text{ curve } = \sum_{i=1}^k [T_{(i)} - T_{(i-1)}] S(T_{(i-1)})$$

where $T_{(i)}$ are the observed times to the event for the k uncensored units, with $T_{(0)} = 0, S(0) = 1$.

The estimated standard error of $\widehat{S}(t)$ can be obtained from the Greenwood Formula:

Case 1: No Censoring

 $\widehat{S}(t) = \frac{n_t}{n}$, where n_t is the number of n units still working at time t

$$Var(\widehat{S}(t)) = \frac{S(t)[1-S(t)]}{n} \quad \Rightarrow \quad \widehat{SE}[\widehat{S}(t)] = \sqrt{\frac{\widehat{S}(t)[1-\widehat{S}(t)]}{n}}$$

Case 2: Censoring with Tied Observations

$$\widehat{SE}[\widehat{S}(t)] = \widehat{S}(t) \sqrt{\sum_{j:t_j \le t} \frac{d_j}{n_j(n_j - d_j)}}$$

The courses STAT 643-644 or STAT 645 will extensively cover the topic of censored data and survival analysis.

The following books are good references for the analysis of censored data and survival analysis.

- ullet The Statistical Models and Methods for Lifetime Data by Lawless
- \bullet Survival Analysis $Using S\:$ by Tableman and Kim
- \bullet Statistical Methods for Reliability Data by Meeker and Escobar

The Kaplin-Meier PL Estimator will now be obtained for the data in the cord strength example using the following SAS code:

```
cord_censored_KM.sas
```

```
option ls=75 ps=55 nocenter nodate;
title 'Strength of Braided Cord'
data cords;
input S C @@;
label S = 'Strength of Cord' C = 'Censoring (1=Yes)';
cards;
36.3 0 52.4 0 54.8 0 57.1 0 60.7 0 41.7 0 52.6 0 54.8 0 57.3 0
43.9 0 52.7 0 55.1 0 57.7 0 49.4 0 53.1 0 55.4 0 57.8 0
50.1 0 53.6 0 55.9 0 58.1 0 50.8 0 53.6 0 56.0 0 58.9 0
51.9 0 53.9 0 56.1 0 59.0 0 52.1 0 53.9 0 56.5 0 59.1 0
52.3 0 54.1 0 56.9 0 59.6 0 52.3 0 54.6 0 57.1 0 60.4 0
26.8 1 29.6 1 33.4 1 35.0 1 40.0 1 41.9 1 42.5 1
run;
proc lifetest data=cords outsurv=a plots=(s);
time S*C(1) ;
proc print data=a;
run;
```

OUTPUT FROM LIFETEST:

The LIFETEST Procedure

Product-Limit Survival Estimates

a	a	.	Survival Standard	Number	Number
S	Survival	Failure	Error	Failed	Left
0.0000	1.0000	0	0	0	48
26.8000*				0	47
29.6000*				0	46
33.4000*				0	45
35.0000*				0	44
36.3000	0.9773	0.0227	0.0225	1	43
40.0000*	•			1	42
41.7000	0.9540	0.0460	0.0318	2	41
41.9000*				2	40
42.5000*				2	39
43.9000	0.9295	0.0705	0.0393	3	38
49.4000	0.9051	0.0949	0.0452	4	37
50.1000	0.8806	0.1194	0.0502	5	36
50.8000	0.8562	0.1438	0.0544	6	35
51.9000	0.8317	0.1683	0.0581	7	34
52.1000	0.8072	0.1928	0.0613	8	33
52.3000	•	•		9	32
52.3000	0.7583	0.2417	0.0667	10	31
52.4000	0.7338	0.2662	0.0688	11	30
52.6000	0.7094	0.2906	0.0708	12	29
52.7000	0.6849	0.3151	0.0724	13	28
53.1000	0.6605	0.3395	0.0739	14	27
53.6000	•			15	26
53.6000	0.6115	0.3885	0.0761	16	25
53.9000				17	24
53.9000	0.5626	0.4374	0.0774	18	23
54.1000	0.5382	0.4618	0.0778	19	22
54.6000	0.5137	0.4863	0.0781	20	21
54.8000	0 4649		. 0770	21	20
54.8000	0.4648	0.5352	0.0779	22	19
55.1000	0.4403	0.5597	0.0776	23	18
55.4000 55.9000	0.4158 0.3914	0.5842 0.6086	0.0770 0.0763	24 25	17 16
56.0000	0.3669	0.6331	0.0753	26	15
56.1000	0.3425	0.6575	0.0742	27	14
56.5000	0.3180	0.6820	0.0728	28	13
56.9000	0.2935	0.7065	0.0712	29	12
57.1000	0.2000	0.11000	0.0112	30	11
57.1000	0.2446	0.7554	0.0672	31	10
57.3000	0.2202	0.7798	0.0648	32	9
57.7000	0.1957	0.8043	0.0620	33	8
57.8000	0.1712	0.8288	0.0589	34	7
58.1000	0.1468	0.8532	0.0553	35	6
58.9000	0.1223	0.8777	0.0512	36	5
59.0000	0.0978	0.9022	0.0465	37	4
59.1000	0.0734	0.9266	0.0408	38	3
59.6000	0.0489	0.9511	0.0337	39	2
60.4000	0.0245	0.9755	0.0242	40	1
60.7000	0	1.0000	0	41	0

NOTE: The marked survival times are censored observations.

Summary Statistics for Time Variable ${\tt S}$

Quartile Estimates

	Point	95% Confi	dence Interval
Percent	Estimate	[Lower	Upper)
75	57.1000	55.9000	58.9000
50	54.8000	53.6000	56.1000
25	52.4000	50.8000	53.9000

Mean Standard Error 54.1824 0.7316

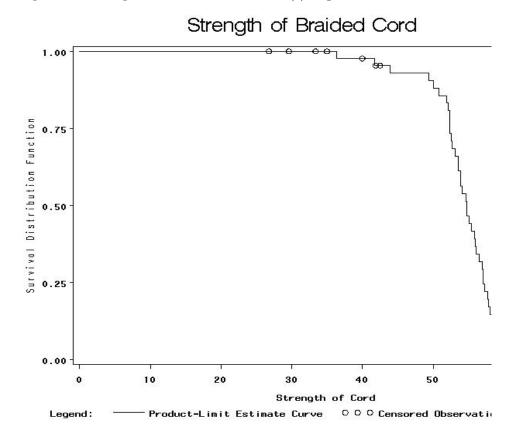
Summary of the Number of Censored and Uncensored Values

			Percent
Total	Failed	Censored	Censored
48	41	7	14.58

Strength of Braided Cord

0bs	S	_CENSOR_	SURVIVAL	SDF_LCL	SDF_UCL
1	0.0	:	1.00000	1.00000	1.00000
2	26.8	0	1.00000	•	•
3	29.6	0	1.00000	•	•
4	33.4	0	1.00000	•	•
5	35.0	0	1.00000		
6	36.3	1	0.97727	0.84941	0.99677
7	40.0	0	0.97727		
8	41.7	1	0.95400	0.82832	0.98830
9	41.9	0	0.95400	•	•
10	42.5	0	0.95400	. 70700	
11	43.9	1	0.92954	0.79702	0.97675
12	49.4	1	0.90508	0.76630	0.96332
13	50.1	1	0.88062	0.73639	0.94855
14	50.8	1	0.85616	0.70725	0.93274
15	51.9	1	0.83170	0.67882	0.91607
16	52.1	1	0.80723	0.65102	0.89867
17	52.3	1	0.75831	0.59709	0.86206
18	52.4	1	0.73385	0.57087	0.84298
19	52.6	1	0.70939	0.54510	0.82342
20	52.7	1	0.68493	0.51975	0.80344
21	53.1	1	0.66046	0.49480	0.78305
22	53.6	1	0.61154	0.44606	0.74114
23	53.9	1	0.56262	0.39877	0.69781
24	54.1	1	0.53816	0.37565	0.67563
25	54.6	1	0.51369	0.35288	0.65313
26	54.8	1	0.46477	0.30837	0.60712
27	55.1	1	0.44031	0.28665	0.58362
28	55.4	1	0.41585	0.26528	0.55979
29	55.9	1	0.39139	0.24428	0.53562
30	56.0	1	0.36692	0.22366	0.51109
31	56.1	1	0.34246	0.20344	0.48620
32	56.5	1	0.31800	0.18363	0.46094
33	56.9	1	0.29354	0.16426	0.43528
34	57.1	1	0.24462	0.12696	0.38266
35	57.3	1	0.22015	0.10911	0.35563
36	57.7	1	0.19569	0.09188	0.32806
37	57.8	1	0.17123	0.07533	0.29988
38	58.1	1	0.14677	0.05959	0.27101
39	58.9	1	0.12231	0.04479	0.24133
40	59.0	1	0.09785	0.03115	0.21067
41	59.1	1	0.07338	0.01900	0.17881
42	59.6	1	0.04892	0.00889	0.14542
43	60.4	1	0.02446	0.00193	0.11054
44	60.7	1	0.00000	•	•

A plot of the Kaplan-Meier Estimator of S(t) is given below.



A comparison of the KM estimator to the values obtained from the Weibull model are displayed next:

$$\hat{S}_W(53.1) = e^{-(53.1/56.0223)^{16.2591}} = .6508 \qquad \hat{S}_{KM}(53.1) = .6605$$

$$\hat{\mu}_{KM} = \sum_{i=1}^{n_{\nu}} \left[T_{(i)} - T_{(i-1)} \right] S(T_{(i-1)})$$

$$= (36.3 - 0)(1.0) + (41.7 - 36.3)(.97727) + (43.9 - 41.7)(.954) + \cdots + (60.7 - 60.4)(.02446)$$

$$= 54.1824$$

$$\hat{\mu}_W = \hat{\alpha}\Gamma\left(\frac{1+\hat{\gamma}}{\hat{\gamma}}\right) = \Gamma\left(\frac{17.2602}{16.2602}\right) = 54.4630$$

$$\hat{Q}_{KM}(.5) = \hat{S}_{K-M}(.5) = 54.8000$$

$$\hat{Q}_{W}(.5) = \hat{\alpha}(-\log(.5))^{1/\hat{\gamma}} = (56.2609)(-\log(.5))^{1/16.2602} = 55.0069$$

The estimates based on the Kaplan-Meier distribution free procedure and the MLE's based on a Weibull model are very close. From this example, it would appear that the distribution-free estimator yields values very close to the MLE's when there is a reasonable sample size.

An analysis of the above data will be conducted using the following R code:

censoredcords_KM.R

```
library(MASS)
library(survival)
st = c(36.3, 52.4, 54.8, 57.1, 60.7, 41.7, 52.6, 54.8, 57.3,
     43.9, 52.7, 55.1, 57.7, 49.4, 53.1, 55.4, 57.8, 50.1,
     53.6, 55.9, 58.1, 50.8, 53.6, 56.0, 58.9, 51.9, 53.9,
     56.1, 59.0, 52.1, 53.9, 56.5, 59.1, 52.3, 54.1, 56.9,
     59.6, 52.3, 54.6, 57.1, 60.4,
     26.8, 29.6, 33.4, 35.0, 40.0, 41.9, 42.5)
0,0,0,0,0,0)
Surv(st, stcens)
cords.surv <- survfit(Surv(st, stcens) ~ 1,conf.type="log-log")</pre>
summary(cords.surv)
print(cords.surv,print.rmean=TRUE)
postscript("u:/meth1/psfiles/cords_surv.ps",height=8,horizontal=F)
plot(cords.surv,conf.int=F,log=TRUE,
main="Kaplan-Meier Estimator of Survival Function", xlab="Strength of Cord",
ylab="Survival Function")
graphics.off()
```

Estimator from R Code:

Call: survfit(formula = Surv(st, stcens) ~ 1, conf.type = "log-log") time n.risk n.event survival std.err lower 95% CI upper 95% CI 36.3 44 0.9773 0.0225 0.84941 0.997 1 41.7 42 1 0.9540 0.0318 0.82832 0.988 0.79702 43.9 39 0.977 1 0.9295 0.0393 49.4 38 1 0.9051 0.0452 0.76630 0.963 50.1 37 1 0.8806 0.0502 0.73639 0.949 50.8 36 0.8562 0.70725 0.933 1 0.0544 51.9 35 1 0.8317 0.67882 0.916 0.0581 0.65102 52.1 34 1 0.8072 0.0613 0.899 52.3 2 0.862 33 0.7583 0.0667 0.59709 52.4 31 1 0.7338 0.0688 0.57087 0.843 52.6 30 0.7094 0.823 1 0.0708 0.54510 52.7 29 0.6849 0.0724 0.51975 0.803 1 53.1 28 0.6605 0.0739 0.783 1 0.49480 53.6 27 2 0.6115 0.0761 0.44606 0.741 53.9 25 2 0.5626 0.698 0.0774 0.39877 54.1 23 1 0.5382 0.0778 0.37565 0.676 54.6 0.5137 0.653 22 1 0.0781 0.35288 54.8 2 0.4648 0.607 21 0.0779 0.30837 55.1 19 1 0.4403 0.0776 0.28665 0.584 55.4 18 1 0.4158 0.0770 0.26528 0.560 55.9 17 1 0.3914 0.0763 0.24428 0.536 56.0 16 1 0.3669 0.0753 0.22366 0.511 56.1 0.3425 0.20344 0.486 15 1 0.0742 56.5 14 0.3180 0.461 1 0.0728 0.18363 0.2935 0.435 56.9 13 1 0.0712 0.16426 57.1 12 2 0.2446 0.0672 0.12696 0.383 57.3 0.2202 0.356 10 1 0.0648 0.10911 57.7 9 1 0.1957 0.0620 0.09188 0.328 57.8 8 1 0.1712 0.0589 0.07533 0.300 7 58.1 0.1468 0.05959 0.271 1 0.0553 58.9 6 1 0.1223 0.0512 0.04479 0.241 59.0 5 1 0.0978 0.0465 0.03115 0.211 0.0408 59.1 4 0.0734 0.01900 0.179 1 59.6 3 1 0.0489 0.0337 0.00889 0.145 60.4 2 1 0.0245 0.0242 0.00193 0.111

Call: survfit(formula = Surv(st, stcens) ~ 1, conf.type = "log-log")

NaN

```
records
          n.max
                   n.start
                             events
                                     *rmean
                                              *se(rmean)
                                                          median
                                                                   0.95LCL
                                                                              0.95UCL
         48.000
                   48.000
                                                           54.800
                                                                                56.100
 48.000
                             41.000
                                      54.182
                                                0.723
                                                                    53.100
* restricted mean with upper limit = 60.7
```

NA

NA

A graph of the survival is produced next:

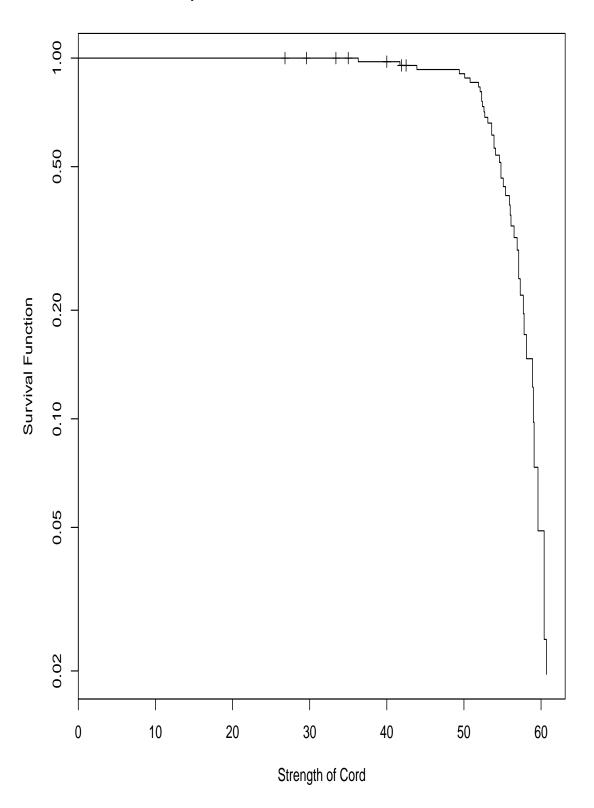
1

0.0000

60.7

1

Kaplan-Meier Estimator of Survival Function



Comparing Two Survival Curves from Censored Data

EXAMPLE from The Statistical Analysis of Failure Time Data

The following table gives the times from insult with the carcinogen DMBA to mortality from vaginal cancer in rats. Two groups were distinguished by a pretreatment regime. Four times to mortality were randomly censored and are denoted by an *. For these rats, we can see that their times to mortality exceed 216, 244, 204, and 344 days, respectively, but we do not know the times exactly. The random censoring occurred because the four rats died of causes unrelated to the application of the carcinogen and they were free of tumor at death, or they may simply not have developed tumor at the time of data analysis. There are two separate groups of survival times data in this example.

		Days to Vaginal Cancer Mortality in Rats								
Group 1	143	164	188	188	190	192	206	209	213	216
	220	227	230	234	246	265	304	216*	244*	
Group 2	142	156	163	198	205	232	232	233	233	233
	233	239	240	261	280	280	296	296	323	204*
	344*									

The following SAS program was used to obtain a Kaplan-Meier estimator for each of the two groups of rats.

longneck/reliability/kmest_rat.sas;

```
option ls=75 ps=55 nocenter nodate;
title 'Cancer Treatment-Estimated S(t)';
data cancer;
input T ST G @@;
LGT=log(T);
label ST = 'Censoring Indicator';
label T = 'Time to Death';
label G = 'Treatment Group';
143 1 1 164 1 1 188 1 1 188 1 1 190 1 1 192 1 1 206 1 1
209 1 1 213 1 1 216 1 1 220 1 1 227 1 1 230 1 1 234 1 1
246 1 1 265 1 1 304 1 1 216 0 1 244 0 1
142 1 2 156 1 2 163 1 2 198 1 2 205 1 2 232 1 2 232 1 2
233 1 2 233 1 2 233 1 2 233 1 2 239 1 2 240 1 2 261 1 2
280 1 2 280 1 2 296 1 2 296 1 2 323 1 2 204 0 2 344 0 2
run:
proc lifetest data=cancer outsurv=a plots=(s);
time T*ST(0);
strata G;
proc print data=a;
run;
```

OUTPUT FROM SAS:

Cancer Treatment-Estimated S(t)

The LIFETEST Procedure

Stratum 1: G = 1

Product-Limit Survival Estimates

			Survival		
			Standard	Number	Number
T	Survival	Failure	Error	Failed	Left
0.000	1.0000	0	0	0	19
143.000	0.9474	0.0526	0.0512	1	18
164.000	0.8947	0.1053	0.0704	2	17
188.000				3	16
188.000	0.7895	0.2105	0.0935	4	15
190.000	0.7368	0.2632	0.1010	5	14
192.000	0.6842	0.3158	0.1066	6	13
206.000	0.6316	0.3684	0.1107	7	12
209.000	0.5789	0.4211	0.1133	8	11
213.000	0.5263	0.4737	0.1145	9	10
216.000	0.4737	0.5263	0.1145	10	9
216.000*				10	8
220.000	0.4145	0.5855	0.1145	11	7
227.000	0.3553	0.6447	0.1124	12	6
230.000	0.2961	0.7039	0.1082	13	5
234.000	0.2368	0.7632	0.1015	14	4
244.000*	•		•	14	3
246.000	0.1579	0.8421	0.0934	15	2
265.000	0.0789	0.9211	0.0728	16	1
304.000	0	1.0000	0	17	0

 $\ensuremath{\mathsf{NOTE}}\xspace$. The marked survival times are censored observations.

Summary Statistics for Time Variable $\ensuremath{\mathsf{T}}$

Quartile Estimates

	Point	95% Confide	ence Interval
Percent	Estimate	[Lower	Upper)
75	234.000	216.000	265.000
50	216.000	192.000	234.000
25	190.000	188.000	216.000

Mean Standard Error 218.757 9.403

Stratum 2: G = 2

Product-Limit Survival Estimates

			Survival Standard	Number	Number
Т	Survival	Failure	Error	Failed	Left
0.000	1.0000	0	0	0	21
142.000	0.9524	0.0476	0.0465	1	20
156.000	0.9048	0.0952	0.0641	2	19
163.000	0.8571	0.1429	0.0764	3	18
198.000	0.8095	0.1905	0.0857	4	17
204.000*				4	16
205.000	0.7589	0.2411	0.0941	5	15
232.000				6	14
232.000	0.6577	0.3423	0.1053	7	13
233.000	•	•	•	8	12
233.000	•	•	•	9	11
233.000				10	10
233.000	0.4554	0.5446	0.1114	11	9
239.000	0.4048	0.5952	0.1099	12	8
240.000	0.3542	0.6458	0.1072	13	7
261.000	0.3036	0.6964	0.1031	14	6
280.000				15	5
280.000	0.2024	0.7976	0.0902	16	4
296.000				17	3
296.000	0.1012	0.8988	0.0678	18	2
323.000	0.0506	0.9494	0.0493	19	1
344.000*				19	0

NOTE: The marked survival times are censored observations.

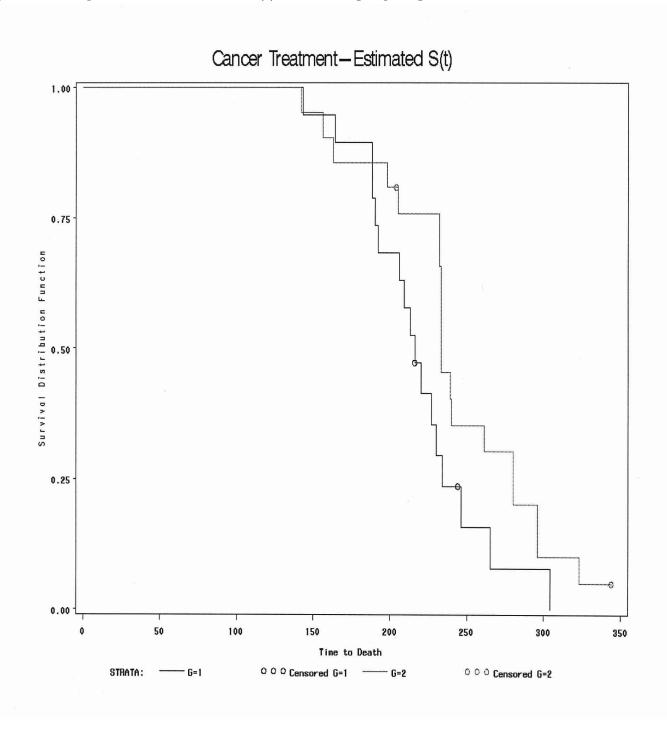
Summary Statistics for Time Variable ${\tt T}$

Quartile Estimates

	Point	95% Confi	dence Interval
Percent	Estimate	[Lower	Upper)
75	280.000	233.000	296.000
50	233.000	232.000	280.000
25	232.000	163.000	233.000

Mean Standard Error

240.795 11.206



The following R code will produce similar results to what was achieved using the SAS code:

```
carcinogen_2G-KM.R
library(survival)
T = c(143,164,188,188,190,192,206,209,213,216,
     220,227,230,234,246,265,304,216,244,
     142,156,163,198,205,232,232,233,233,233,
     233,239,240,261,280,280,296,296,323,204,
     344)
1,1,1,1,1,1,1,1,1,1,1,1,1,1,1,1,1,1,0,0)
out = cbind(T,ST,G)
Surv(T, ST)
carcin <- survfit(Surv(T, ST) ~ G)</pre>
summary(carcin)
print(carcin, print.rmean=TRUE,rmean="individual")
postscript("u:/meth1/psfiles/carcin_surv.ps",height=8,horizontal=F)
plot(carcin,ylab="Survival Function",xlab="Time to Death",
main="Cancer Treatment - Estimated S(t)",lty=1:2 )
legend(25,.8,c("Group 1","Group 2"),lty=1:2,lwd=2)
graphics.off()
```

OUTPUT FROM R:

records n.max n.start events *rmean *se(rmean) median 0.95LCL 0.95UCL G=1 9.12 G=2 11.35

Call: survfit(formula = Surv(T, ST) ~ G, conf.type = "log-log")

		G=1				
time	n.risk	${\tt n.event}$	survival	${\tt std.err}$	lower 95% CI	upper 95% CI
143	19	1	0.947	0.0512	0.68119	0.992
164	18	1	0.895	0.0704	0.64079	0.973
188	17	2	0.789	0.0935	0.53191	0.915
190	15	1	0.737	0.1010	0.47893	0.881
192	14	1	0.684	0.1066	0.42794	0.844
206	13	1	0.632	0.1107	0.37899	0.804
209	12	1	0.579	0.1133	0.33208	0.763
213	11	1	0.526	0.1145	0.28720	0.719
216	10	1	0.474	0.1145	0.24438	0.673
220	8	1	0.414	0.1145	0.19616	0.621
227	7	1	0.355	0.1124	0.15191	0.566
230	6	1	0.296	0.1082	0.11168	0.509
234	5	1	0.237	0.1015	0.07578	0.447
246	3	1	0.158	0.0934	0.03143	0.374
265	2	1	0.079	0.0728	0.00567	0.288
304	1	1	0.000	NaN	NA	NA

G=2

time	n.risk	n.event	${\tt survival}$	std.err	lower 95% CI	upper 95% CI
142	21	1	0.9524	0.0465	0.70721	0.993
156	20	1	0.9048	0.0641	0.67005	0.975
163	19	1	0.8571	0.0764	0.61972	0.952
198	18	1	0.8095	0.0857	0.56891	0.924
205	16	1	0.7589	0.0941	0.51394	0.892
232	15	2	0.6577	0.1053	0.41232	0.820
233	13	4	0.4554	0.1114	0.23531	0.652
239	9	1	0.4048	0.1099	0.19615	0.605
240	8	1	0.3542	0.1072	0.15914	0.556
261	7	1	0.3036	0.1031	0.12446	0.506
280	6	2	0.2024	0.0902	0.06327	0.397
296	4	2	0.1012	0.0678	0.01719	0.275
323	2	1	0.0506	0.0493	0.00349	0.207

Cancer Treatment – Estimated S(t)

