

# Mean survival time by ordered fractions of population with censored data

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## Mean survival by ordered fractions

- Let  $T$  be a non-negative random variable with  $E[T] < \infty$ .
- Expectation of  $T$  can be expressed as

$$\mu = E[T] = \int_0^\infty S(t)dt = \int_0^1 Q(p)dp, \quad (1)$$

where  $S(\cdot)$  and  $Q(\cdot)$  denote the survival and quantile functions, respectively.

- Given  $\{\lambda_0, \lambda_1, \dots, \lambda_K\}$  grid of proportions, we divide  $\mu$  into separate components

$$\mu = \sum_{k=1}^K \mu_k, \text{ where } \mu_k = \int_{\lambda_{k-1}}^{\lambda_k} Q(p)dp, \quad \lambda_0 = 0 \text{ and } \lambda_K = 1, \quad (2)$$

with  $\lambda_{k-1} < \lambda_k, \forall k \in \{1, \dots, K\}$ .

- Weighting each  $\mu_k$  by its corresponding inverse proportion

$$\bar{\mu}_k = \frac{\mu_k}{\lambda_k - \lambda_{k-1}},$$

we obtain the mean survival time for the fraction of population defined by  $(\lambda_{k-1}, \lambda_k)$ .

### Easy interpretation

If  $(\lambda_0, \lambda_1) = (0, 0.5)$ ,  $\bar{\mu}_1$  quantifies mean survival time for the first half of the population to experience the event of interest.

- In the presence of a censoring variable  $C$ , when  $Y = \min(T, C)$  is observed,  $\lambda_K$  in (2) can be set to the largest proportion of observed events (corresponding to the last observed quantile).

## Estimation and simulation results

- We estimate  $\mu_k$  via the Kaplan-Meier estimator of the underlying survival function.
- Given  $\hat{S}(\cdot)$  and the grid of proportions  $\{\gamma_0, \gamma_1, \dots, \gamma_K\} = \{1 - \lambda_0, 1 - \lambda_1, \dots, 1 - \lambda_K\}$ , from (1) and (2)

$$\begin{aligned} \hat{\mu}_k &= \sum_{j=1}^{J_k} y_j [\min\{\hat{S}(y_{j-1}), \gamma_{k-1}\} - \max\{\hat{S}(y_j), \gamma_k\}] \\ &= \sum_{j=1}^{J_k} \hat{Q}(p_j)(p_j - p_{j-1}), \end{aligned}$$

where  $y_j$  are observed event times such that  $\hat{S}(y_j) \in [\gamma_k, \gamma_{k-1}]$ ,  $\forall j \in \{1, \dots, J_k\}$ , and  $\hat{S}(y_0) \geq \gamma_{k-1}$  and  $\hat{S}(y_{J_k}) \leq \gamma_k$ .

**Table 1:** Average over 5,000 samples with 200 observations from a log-logistic model with scale  $\alpha = 1$  and shape  $\beta = 2$ , and censoring variable uniform in  $(0, 7/3)$  (average censoring rate of 50%).

$k$	$\lambda_k$	$\mu_k$	$\hat{\mu}_k$	$\hat{\mu}_k^L - \hat{\mu}_k^U$	nsim <sub>k</sub>	d <sub>k</sub>
1	0.20	0.064	0.064	0.044–0.086	100%	34
2	0.40	0.131	0.132	0.101–0.175	100%	29
3	0.60	0.201	0.202	0.156–0.264*	100%	23
4	0.80	0.311	0.304	0.226– $\infty$	70.7%	14
5	0.95	0.420	0.307	0.239– $\infty$	5.80%	4

Where

- $\mu_k$  true mean survival time for fraction  $(\lambda_{k-1}, \lambda_k)$ .
- $\hat{\mu}_k$  estimated mean survival time for fraction  $(\lambda_{k-1}, \lambda_k)$ .
- $\hat{\mu}_k^L$  and  $\hat{\mu}_k^U$  lower and upper bounds integrating over equal precision confidence bands for the Kaplan-Meier estimator [1].
- nsim<sub>k</sub> % of simulations where  $\hat{\mu}_k$  could be computed.
- d<sub>k</sub> number of observed events.

### Precision

Estimates' precision decreases with increasing  $k$ , as the proportion of censored observations increases with  $k$  and, on average, fewer events are observed.

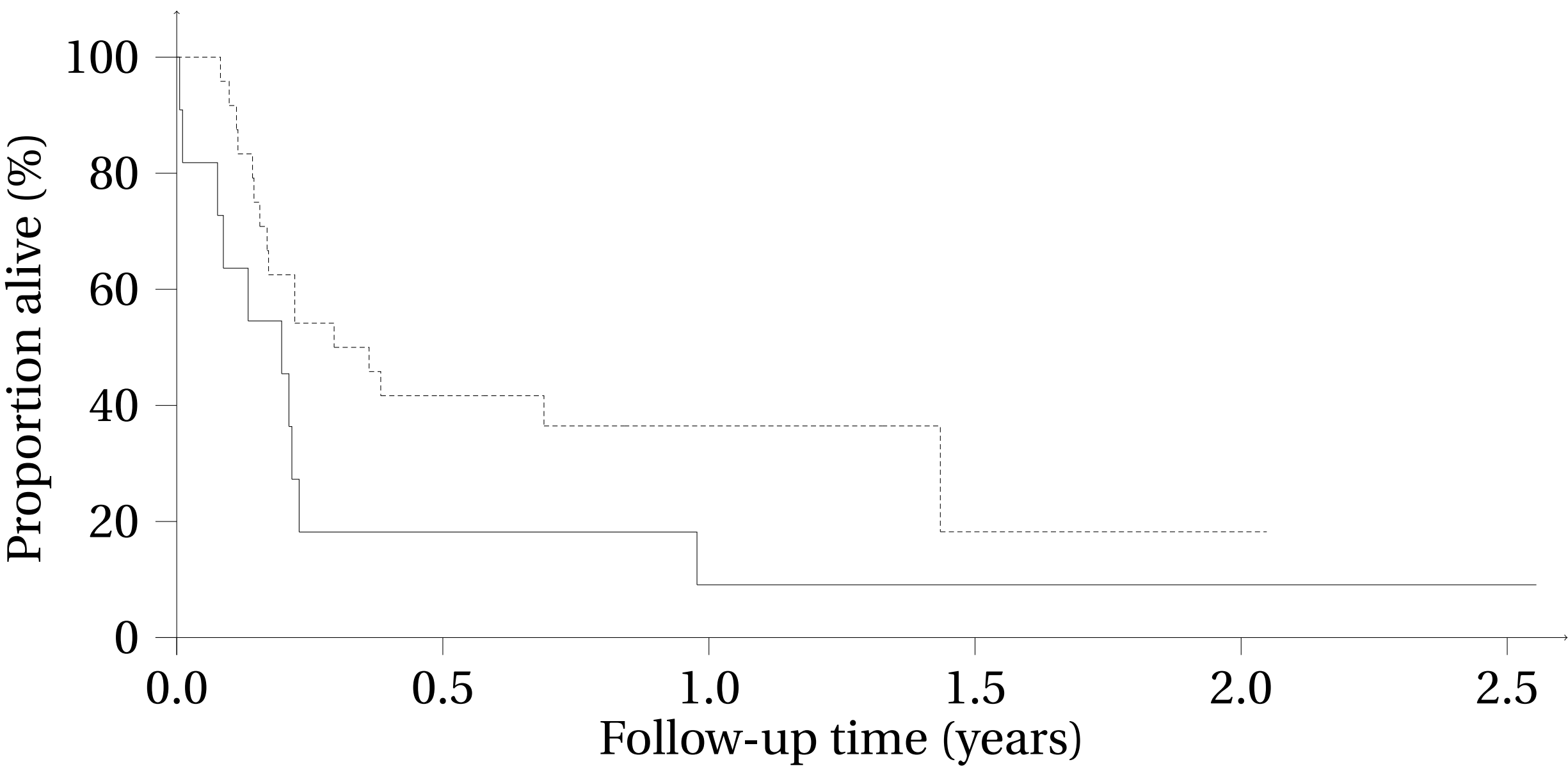
### Mean survival by ordered fractions vs. Restricted mean

When  $\lambda_K < 1$ , the mean survival time for the  $\lambda_K$ -th fraction of the population observed to experience the event, does not correspond to the restricted mean [2] computed up to the last observed quantile  $y^* = Q(\lambda_K)$ , i.e.,

$$\bar{\mu}_K = \frac{1}{\lambda_K} \int_0^{\lambda_K} Q(p)dp \neq \int_0^{y^*} S(y)dy = \mu^*.$$

## Application example: Survival after bone marrow transplant in lymphoma patients

- Data on 35 patients with lymphoma that received either an allogenic or an autologous bone marrow transplant [3].
- Aim of the study: finding differences in lymphoma-free survival after having received either type of transplant.



**Figure 1:** Estimated Kaplan-Meier survival curves after bone marrow transplant for lymphoma patients that received allogenic (solid line) or autologus (dashed line) transplant.

### Results' comparison

- Restricted mean survival estimates do not detect significant differences between groups  
146.5 days, 95% CI  $(-29.71, 322.7)$ .
- Mean survival by ordered fractions estimates detect significant differences among earlier failures  
32.15 days, 95% CI  $(13.98, 50.31)$ ,  
among the weakest 10% of the patients (first 10% to die or relapse in each group after receiving the transplant).

**Table 2:** Estimates for mean survival differences (in days) between allogenic ( $\hat{\mu}_k^1$ ) and autologus ( $\hat{\mu}_k^0$ ) bone marrow transplants with bootstrapped 95% confidence intervals by ordered deciles of population up to the last commonly observed fraction (80th percentile).

$k$	$\lambda_k$	$\hat{\mu}_k^1 - \hat{\mu}_k^0$	95% CI
1	0.1	32.15	13.98–50.31
2	0.2	36.72	2.843–70.60
3	0.3	26.23	–19.94–72.43
4	0.4	28.98	–40.51–98.48
5	0.5	32.80	–124.5–190.1
6	0.6	80.60	–1283–1444
7	0.7	349.4	–446.4–1145
8	0.8	441.4	–130.3–1013

- We detected an improvement on lymphoma-free survival for the autologus transplant group amongst the weakest 20% of patients, that is, amongst those who first died or relapsed.
- No difference between the groups was detected when using restricted mean estimates.

## References

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