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### **Definitions**

- **Censoring**: Incomplete information about a patient's survival time, occurring when the study ends before the event (like death) happens, or if the patient is lost to follow-up.
- Follow-up: The period during which patients in a study are monitored for outcomes.
- Logrank Test: A statistical test used to compare survival distributions of two groups.
- **Proportional Hazards**: The assumption that the ratio of hazard rates (risk of event happening) between two groups is constant over time.

# Key Findings

- Unequal censoring and insufficient follow-up can lead to inaccurate conclusions about treatment effects in clinical studies.
- Simulation studies show that different censoring proportions can affect the performance of the logrank test.
- Insufficient follow-up may result in random censoring, which can bias the results.

## **Introduction**

The study examines how unequal censoring and insufficient follow-up in clinical trials affect the comparison of survival outcomes between treatment groups. Clinical trials often have a specified follow-up period, and patients who do not experience the event by the end of the study are censored. This can lead to unequal censoring, which impacts the validity of survival analysis.

## Main Content

#### Background

Clinical trials are designed to compare treatment effects on survival. Patients are followed for a set time, and those who don't experience the event are censored. Unequal censoring occurs when different numbers of patients are censored in each treatment group, which can skew results.

## Methods

- Simulation Studies:
  - Created scenarios with different censoring proportions and follow-up times.
  - Used statistical methods to evaluate the impact on survival analysis.

• Compared performance of the logrank test under these conditions.

### Results

- Unequal Censoring:
  - When censoring proportions were unequal, the logrank test's type I error rate (false positive) increased.
  - Greater differences in censoring proportions led to higher bias.

### • Insufficient Follow-up:

• Random censoring due to insufficient follow-up can misclassify failure times, leading to biased estimates.

### **Conclusion**

The study highlights the importance of considering censoring mechanisms in the design and analysis of clinical trials. Unequal censoring and insufficient follow-up can lead to invalid conclusions about treatment effects. Proper planning and analysis strategies are needed to mitigate these issues and ensure accurate survival comparisons.

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