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Dose Finding in Clinical Trials | Dose Finding by the ...

The continual reassessment method (CRM) is a model-based design for phase I trials, which aims to find the maximum tolerated dose (MTD) of a new therapy. The CRM has been shown to be more accurate in targeting the MTD than traditional rule-based approaches such as the 3 + 3 design, which is used in most phase I trials. Furthermore, the CRM has been shown to assign more trial participants at or ...

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As clinicians begin to realize the important role of dose-finding in the drug development process, there is an increasing openness to "novel" methods proposed in the past two decades. In particular, the Continual Reassessment Method (CRM) and its variations have drawn much attention in the medical community, though it has yet to become a commonplace tool.

Dose Finding by the Continual Reassessment Method

Dose Finding by the Continual Reassessment Method (Chapman & Hall/CRC Biostatistics Series Book 41) eBook: Ying Kuen Cheung: Amazon.ca: Kindle Store

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The continual reassessment method (CRM) proposed by O'Quigley, Pepe and Fisher (1990) is an influential phase I clinical trial design for finding the maximum tolerated dose (MTD) of a new drug. The CRM assumes a single-parameter working dose-toxicity model and continuously updates the estimates of the toxicity probabilities of the considered doses to guide dose escalation.

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Dose Finding by the Continual Reassessment Method: Cheung ...

The book focuses on the design (not analysis) of phase I and phase II dose-finding trials using the continual reassessment method (CRM) and its variants. The method is introduced alongside a description of the R package `dfcrm`, aiming to provide the reader with the skills to implement the method in R. ...

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