BOIN Design Simulation Set-up

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Objective

Replication of Bayesian Optimal Interval Design: A Simple and Well-Performing Design for Phase I Oncology Trials by Yuan et al.

Simulation setting

- 5 Dose levels;
- Max sample size: 30 patients;
- Number of Cohorts: 10;
- Dose Limiting Toxicity (DLT) rate: 15%, 20%, 25%, 30%;
- For each DLT, 16 toxicity scenarios (location of MTD and gaps around MTD);
- Under each scenario (see ToxicityScenarios_DLTrates.xlsx), do 10,000 trails.

- Traditional 3+3

- Dose level 1: A B C (Cohort 1)
 0 DLT (0 out of 3): D E F at Dose level 2;
 - 1 DLT (1 out of 3): *D E F* at Dose level 1; * 0 DLT (1 out of 6): *G H I* at Dose level 2.
- IF >1 DLT out of 3 or 6 THEN Current dose level $(Dose_i)$ > MTD:
 - IF less than 6 patients have already been at $Dose_{i-1}$, THEN add a cohort of 3 at $Dose_{i-1}$;
 - IF 6 patients have already been at $Dose_{i-1}$, THEN $Dose_{i-1} = MTD$;
 - IF $Dose_{i-1} = Dose_1$, THEN the trial is terminated and the MTD is not found.
- Remaining patients are considered treated at the selected MTD.

- Local BOIN

Boundaries

- Global BOIN

Boundaries

Data generation

how are the trial data (not the dose scenario) in each simulation generated?

BOIN_SuyuLiu&YingYuan.pdf Fig.3 replicate in Simulation.Rmd

Package

BOIN

BOIN2.4_manual.pdf BOIN2.4_tutorial.pdf

Escalation

Info Website:

https://cran.r-project.org/web/packages/escalation/vignettes/A700-Simulation.html

STAN

Performance metrics

- Percentage of correct selection (PCS) of the MTD

Percentage of correct selection (PCS) of the true MTD in 10000 simulation trials.

- Average number of patients allocated to the MTD
- Risk of overdosing
- Risk of underdosing