Package 'BayesDesign'

July 21, 2025

| Type Package |
|--|
| Title Bayesian Single-Arm Design with Survival Endpoints |
| Version 0.1.1 |
| Description The proposed event-driven approach for Bayesian two-stage single-arm phase II trial design is a novel clinical trial design and can be regarded as an extension of the Simon's two-stage design with the time-to-event endpoint. This design is motivated by cancer clinical trials with immunotherapy and molecularly targeted therapy, in which time-to-event endpoint is often a desired endpoint. |
| License GPL-2 |
| Encoding UTF-8 |
| NeedsCompilation no |
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| optimal_OneStage | Obtain design settings for one-stage Bayesian Single-Arm Phase II Trial with Time-to-Event Endpoints |
|------------------|--|
| | |

Description

Obtain design parameters, type I error, power and operating characteristics of the Bayesian Single-Arm Phase II Trial Designs with Time-to-Event Endpoints (Wu et al. 2021). The exponential distribution is assumed for the survival time. The gamma prior is used here

Usage

Arguments

| • | • | |
|---|-------------|--|
| | alphacutoff | the desired type I error to be controlled |
| | powercutoff | the desired power to be achieved |
| | S0 | the survival probability at timepoint x |
| | x | the survival probability S0 at timepoint x |
| | ta | accrual duration |
| | tf | follow-up duration |
| | а | shape parameter of prior distribution. The default value is $a = 2$ |
| | delta | hazard ratio |
| | ntrial | the number of simulated trials |
| | complete | whether output the full or partial information. The default value is complete = "partial". If want to show full results, it would be complete = "complete" |
| | seed | the seed. The default value is seed = 8232 |
| | | |

Value

optimal_OneStage() depending on the argument "complete", it returns a vector of partial information/complete information which includes:

partial information: (1) m: number of events of the whole design (2) n: number of patients of the whole design (3) k: total observation time of the whole design (4) typeI: type I error of the whole design (5) power: power of the whole design (6) ES1: expected sample size under alternative hypothesis (7) ES0: expected sample size under null hypothesis

full information: (1) eta: cutoff point of "Go" at final stage of analysis (2) zeta: cutoff point of "no-Go" at final stage of analysis (3) m: number of events of the whole design (4) n: number of patients of the whole design (5) k: total observation time of the whole design (6) typeI: type I error of the whole design (7) power: power of the whole design (8) ES1: expected sample size under alternative hypothesis (9) ES0: expected sample size under null hypothesis

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Author(s)

Chia-Wei Hsu, Haitao Pan, Jianrong Wu

References

Jianrong Wu, Haitao Pan, Chia-Wei Hsu (2021). "Bayesian Single-Arm Phase II Trial Designs with Time-to-Event Endpoints." Pharmaceutical Statistics. Accepted

Examples

optimal_TwoStage

Obtain design settings for two-stage Bayesian Single-Arm Phase II Trial with Time-to-Event Endpoints

Description

Obtain design parameters, type I error, power and operating characteristics of the Bayesian Single-Arm Phase II Trial Designs with Time-to-Event Endpoints (Wu et al. 2021). The exponential distribution is assumed for the survival time. The gamma prior is used here

Usage

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Arguments

alphacutoff the desired type I error to be controlled powercutoff the desired power to be achieved

S0 the survival probability at timepoint x

x the survival probability S0 at timepoint x

ta accrual duration
tf follow-up duration

a shape parameter of prior distribution. The default value is a = 2

delta hazard ratio

frac a information fraction for interim analysis. The fefault value is frac = 0.5

ntrial the number of simulated trials

complete whether output the full or partial information. The default value is complete =

"partial". If want to show full results, it would be complete = "complete"

seed the seed. The default value is seed = 8232

Value

optimal() depending on the argument "complete", it returns a vector of partial information/complete information which includes:

partial information: (1) m1: number of events at stage 1 (2) n1: number of patients at stage 1 (3) k1: total observation time at stage 1 (4) m: number of events of the whole design (5) n: number of patients of the whole design (6) k: total observation time of the whole design (7) typeI: type I error of the whole design (8) power: power of the whole design (9) PET1: early stopping probabilites under alternative hypothesis (10) ES1: expected sample size under alternative hypothesis (11) PET0: early stopping probabilites under null hypothesis (12) ES0: expected sample size under null hypothesis

full information: (1) eta: cutoff point of "Go" at final stage of analysis (2) xi: cutoff point of "no-Go" at final stage of analysis (3) m1: number of events at stage 1 (4) n1: number of patients at stage 1 (5) k1: total observation time at stage 1 (6) m: number of events of the whole design (7) n: number of patients of the whole design (8) k: total observation time of the whole design (9) typeI: type I error of the whole design (10) power: power of the whole design (11) PET1: early stopping probabilites under alternative hypothesis (12) ES1: expected sample size under alternative hypothesis (13) PET0: early stopping probabilites under null hypothesis (14) ES0: expected sample size under null hypothesis

Author(s)

Chia-Wei Hsu, Haitao Pan, Jianrong Wu

References

Jianrong Wu, Haitao Pan, Chia-Wei Hsu (2021). "Bayesian Single-Arm Phase II Trial Designs with Time-to-Event Endpoints." Pharmaceutical Statistics. Accepted

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Examples

tot_time

Sum up transformed observation time for each patient

Description

Sum up transformed observation time for each patient to get U in order to determine the trial: (1) goes to second stage (2) stops for futility (3) declares the treatment is promising and warrants for further study in a large scale phase III trial (4) declares the treatment is unpromising and is not worth for further study.

Usage

```
tot_time(obs_time, S0, x)
```

Arguments

obs_time a vector. Each element represents an observation time of the patient
the survival probability at timepoint x

the survival probability S0 at timepoint x

Value

the function returns the total transformed observation time for all patients

Author(s)

Chia-Wei Hsu, Haitao Pan, Jianrong Wu

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References

Jianrong Wu, Haitao Pan, Chia-Wei Hsu (2021). "Bayesian Single-Arm Phase II Trial Designs with Time-to-Event Endpoints." Pharmaceutical Statistics. Accepted

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