

Package ‘CRM’

January 20, 2025

Title Continual Reassessment Method (CRM) for Phase I Clinical Trials

Version 1.2.4

Date 2018-9-25

Depends R (>= 2.10.0)

Author Qianxing Mo

Maintainer Qianxing Mo <qianxing.mo@moffitt.org>

Description Functions for phase I clinical trials using the continual reassessment method.

LazyData no

License GPL (>= 2)

NeedsCompilation yes

Repository CRAN

Date/Publication 2018-09-26 16:10:02 UTC

Contents

crm	1
crmsim	3
Index	6

crm *A function for the continued reassessment method (CRM) used in Phase I clinical trials*

Description

Function `crm` implement the continued reassessment method (CRM) for dose finding in Phase I clinical trials.

Usage

```
crm(target,prior,ptdata,model=1,a0=1,b=3)
```



```

y <- c(0,0,1,0,1,1,0,0,0,0,0,0,1,0,0,0,0,0,1,0,0,0,0,0,1,0,0,1,0,1,1)
pdata <- cbind(x,y)
for(i in 1:25){
  if(i == 1){
    cat(1,1,3,0,"\\n")
  }
  res <- crm(target,prior,pdata[1:i,],model=1,a0=1)
  if(i < 25){
    cat(i+1,res$a,res$MTD,pdata[i+1,2],"\\n")
  }else {
    cat(i+1,res$a,res$MTD,"\\n")
  }
}

#the proposed MTD is
res$MTD

```

crmsim

CRM Simulator

Description

crmsim, crmsiminc1 and crmsiminc2 implement the continued reassessment method (CRM) for dose finding in Phase I clinical trials. The operating characteristics of CRM are summarized through simulations.

crmsim allows users to select a variety of cohort sizes. A cohort of subjects are treated at the same dose.

The cohort size is fixed to 1 in crmsiminc1 and crmsiminc2. crmsiminc1 implements an algorithm that allows a clinical trial to proceed to the next subject's dose assignment before observing the last subject's toxicity data. crmsiminc2 allows a clinical trial to proceed to the next subject's dose assignment before observing the last two subject's toxicity data (see Iasonos et al. for details).

Usage

```

crmsim(target,prior,true,rate,cycle,cohort=1,nsubject=24,nsim=1000,
        model=1,a0=1,b=3,jump=FALSE,start.dose=1,seed=777)
crmsiminc1(target,prior,true,rate,cycle,nsubject=24,nsim=1000,
            model=1,a0=1,b=3,jump=FALSE,start.dose=1,seed=777)
crmsiminc2(target,prior,true,rate,cycle,nsubject=24,nsim=1000,
            model=1,a0=1,b=3,jump=FALSE,start.dose=1,seed=777)

```

Arguments

target	Target probability of toxicity. The value must be in (0, 1).
prior	Prior probabilities of toxicity for each dose. The values must be in (0,1) and in an ascending order. For example, prior=c(0.05,0.1,0.2,0.3,0.5,0.7),which corresponds to dose levels 1, 2, 3, 4, 5 and 6,respectively.

true	True probabilities of toxicity. The values must be in (0,1) and in an ascending order. e.g. (0.1,0.2,0.3,0.4,0.5,0.8)
rate	Recruitment/accrual rate of subjects in a 30 window. For example,if 1 subject can be recruited per 30 days, then set rate = 1/30 = 0.033; if 2 patients per 30 day then rate = 2/30 = 0.0667.
cycle	The length of treatment cycle in days.
cohort	Cohort size of subjects entering into the trials. Default is 1. The value for cohort must be less than or equal to the value for nsubject.
nsubject	Total number of subjects in one simulation(or trial). Default is 24. nsubject should be equal to n*cohort,where n is positive integer.
nsim	Total number of simulations. Default is 1000.
model	Dose-toxicity model. The value must be 1 (hyperbolic tangent model) or 2 (one-parameter logistic model). Default is 1. Hyperbolic tangent model: $p(y=1 x,a) = ((\tanh(x)+1)/2)^a$. One-parameter logistic model: $p(y=1 x,a,b) = \exp(b+ax)/(1+\exp(b+ax))$. For both models, y=1 indicates toxicity is observed; a is the parameter that can be updated based on the outcome of the trail; b is a fixed parameter. The prior for a is $\exp(-a)$.
a0	Initial value for parameter a. Default is 1.0.
b	A fixed parameter for the one-parameter logistic model. Default is 3.0.
jump	jump=FALSE means NOT allowing that the proposed dose by the CRM program has an increase of more than one level than the previous level; jump=TRUE means allowing more-than-one-level increase of the proposed dose by the CRM program.
start.dose	Initial dose for each trial. Default is 1.
seed	Seed for the random number generator. Default is 777.

Value

SimResult is a matrix that summarizes the operating characteristics of CRM. The column names are the dose levels. The row names are the operating characteristics.

% Selection	the percentage of selection of each dose as MTD
% Subjects Treated	the percentage of subjects treated at each dose
# Subjects Treated	the average number of subjects treated at each dose
Average Toxicities	the average toxicities per trial at each dose
True Probabilities	the true probability of toxicity of each dose

TrialDuration is a table that summarizes the time needed for the trial based on the simulation.

Author(s)

Qianxing Mo; <qianxing.mo@moffitt.org>

References

John O'Quigley, Margaret Pepe, and Lloyd Fisher. (1990). Continual Reassessment Method: A Practical Design for Phase I Clinical Trials in Cancer. *Biometrics*, 46, pp.33-48

Alexia Iasonos, Andrew S Wilton, Elyn R Riedel, Venkatraman E Seshan and David R Spriggs. A comprehensive comparison of the continual reassessment method to the standard 3+3 dose escalation scheme in Phase I dose-finding studies. *Clinical Trials*, 2008, nil:1-12

See Also

[crm](#)

Examples

```
prior1 <- c(0.05,0.1,0.2,0.3,0.5,0.7)
true1 <- c(0.1,0.15,0.2,0.4,0.5,0.8)

### simulations using model 1 (hyperbolic tangent model)

### uncomment the following code to run ###
#crmsim(target=0.2,prior=prior1,true=true1,rate=0.1,cycle=21,cohort=1,nsubject=24,nsim=1000,
#       model=1,a0=1,b=3,jump=FALSE,start.dose=1,seed=777)

#crmsiminc1(target=0.2,prior=prior1,true=true1,rate=0.1,cycle=21,nsubject=24,nsim=1000,
#           model=1,a0=1,b=3,jump=FALSE,start.dose=1,seed=777)

#crmsiminc2(target=0.2,prior=prior1,true=true1,rate=0.1,cycle=21,nsubject=24,nsim=1000,
#           model=1,a0=1,b=3,jump=FALSE,start.dose=1,seed=777)

# simulations using model 2 (one-parameter logistic model)
#crmsim(target=0.2,prior=prior1,true=true1,rate=0.1,cycle=21,cohort=1,nsubject=24,nsim=1000,
#       model=2,a0=1,b=3,jump=FALSE,start.dose=1,seed=777)

#crmsiminc1(target=0.2,prior=prior1,true=true1,rate=0.1,cycle=21,nsubject=24,nsim=1000,
#           model=2,a0=1,b=3,jump=FALSE,start.dose=1,seed=777)

#crmsiminc2(target=0.2,prior=prior1,true=true1,rate=0.1,cycle=21,nsubject=24,nsim=1000,
#           model=2,a0=1,b=3,jump=FALSE,start.dose=1,seed=777)
```

Index

* design

crm, 1

crmsim, 3

crm, 1, 5

crmsim, 2, 3

crmsiminc1, 2

crmsiminc1 (crmsim), 3

crmsiminc2, 2

crmsiminc2 (crmsim), 3