# Package 'CRM2s'

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Type Package

Title C-Optimality Based Two-Stage Continual Reassessment Method (CRM)
Version 0.1.0
<b>Description</b> Implements CRM2s, a Two-Stage Continual Reassesment Method based in coptimality. Simulates and compares this dose-escalation method in phase I clinical trials with the traditional 3+3 design for two types of dose-toxicity models: power with one parameter and logistic with two parameters.
<pre>URL https://github.com/alvarocia/CRM2s</pre>
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crm2s-imports

Import functions from base packages

#### **Description**

This block imports commonly used base R functions into the package namespace.

```
{\it export\_simulation\_table\_manual} \\ {\it Export~Simulation~Summary~to~LaTeX}
```

## **Description**

Writes a LaTeX table with MTD and toxicity summaries using cat(), without external dependencies.

#### Usage

```
export_simulation_table_manual(
  df,
  file_path = "RESULTS/simulation_results.tex"
)
```

## **Arguments**

df A data frame with simulation summaries for one model (e.g., result\_list\$power)

as returned by simulate\_across\_n\_initial().

file\_path Path to the output .tex file.

#### **Examples**

```
result_list <- simulate_across_n_initial()
export_simulation_table_manual(result_list$power, "table_power.tex")
export_simulation_table_manual(result_list$logistic, "table_logistic.tex")</pre>
```

logistic\_3\_3

3+3 Design Simulation for Logistic model with 2 parameters

## Description

Simulates a single trial using a 3+3 dose-escalation method for estimating the Maximum Tolerated Dose (MTD).

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#### Usage

```
logistic_3_3(
   theta = c(-3, 2),
   theta_0 = c(-3.1, 1.8),
   n_initial = 3,
   p_tox_init = 0.05,
   delta_dosis = 0.092,
   seed = 1234,
   show_plot = FALSE
)
```

## Arguments

theta	True value of the vector of parameters for the dose-toxicity curve. Default is $c(-3,2)$ .
theta_0	Nominal value for the vector of parameters. Default is c(-3.1,1.8).
n_initial	Number of patients per dose level. Default is 3.
p_tox_init	Initial toxicity probability for computing starting dose. Default is 0.05.
delta_dosis	Step size for dose escalation. Default is 0.05.
seed	Random seed for reproducibility. Default is 1234.
show_plot	Logical. If TRUE, plot dose level vs. patient index. Default is FALSE.

#### Value

A list with:

```
n_toxicities Total number of toxicities observed.
```

mtd\_estimated Estimated MTD (last safe dose).

- **n\_patients** Total number of patients enrolled.
- x Dose levels assigned.
- y Observed toxicity outcomes (1 = toxic, 0 = non-toxic).

## **Examples**

```
res <- logistic_3_3()
print(res$mtd_estimated)</pre>
```

power\_3\_3

3+3 Design Simulation for power model

## Description

Simulates one trial using a 3+3 dose-escalation method for estimating the Maximum Tolerated Dose (MTD).

#### Usage

```
power_3_3(
   theta = 3,
   theta_0 = 2.7,
   n_initial = 3,
   p_tox_init = 0.02,
   delta_dosis = 0.05,
   seed = 1234,
   show_plot = FALSE
)
```

#### **Arguments**

theta True value for the dose-toxicity curve. Default is 3.

theta\_0 Nominal value for the dose-toxicity curve. Default is 2.7.

n\_initial Number of patients per dose level. Default is 3.

p\_tox\_init Initial toxicity probability for computing starting dose. Default is 0.02.

delta\_dosis Step size for dose escalation. Default is 0.05.

seed Random seed for reproducibility. Default is 1234.

show\_plot Logical. If TRUE, plot dose level vs. patient index. Default is FALSE.

#### Value

A list with:

```
n_toxicities Total number of toxicities observed.
mtd_estimated Estimated MTD (last safe dose).
n_patients Total number of patients enrolled.
x Dose levels assigned.
```

y Observed toxicity outcomes (1 = toxic, 0 = non-toxic).

#### **Examples**

```
res <- power_3_3()
print(res$mtd_estimated)</pre>
```

```
run_simulation_logistic
```

Simulation Comparison of 3+3 and CRM2s Designs for a Logistic Model with Two Parameters

#### **Description**

Simulates and compares MTD estimates and toxicity outcomes between the 3+3 design and a two-stage CRM design over multiple replications.

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#### Usage

```
run_simulation_logistic(
 num_rep = 1000,
 seed = 1234,
 save_plot = FALSE,
 p0 = 0.4,
  theta = c(-3, 2),
  theta_0 = c(-3.1, 1.8),
 N = 24,
 n_initial = 1,
 q_0 = 0.05,
 q_2 = 0.4,
 q_1 = 0.9
 lim_sup_prob = 0.7,
 p_{tox_i} = 0.05
 delta_dosis_3_3 = 0.092
)
```

#### **Arguments**

num_rep	Number of replications to run. Default is 1000.
seed	Base random seed for reproducibility. Default is 1234.
save_plot	Logical. If TRUE, saves comparison plots as PDF. Default is FALSE.
p0	Target toxicity probability. Default is 0.4.
theta	True value of the vector of parameters for the dose-toxicity curve. Default is $c(-3,2)$ .
theta_0	Nominal value for the vector of parameters. Default is c(-3.1,1.8).
N	Total number of patients in the CRM design. Default is 24.
n_initial	Number of patients per dose level in the CRM design (not used in 3+3). Default is 1.
q_0	Initial toxicity probability for CRM design. Default is 0.05.
q_2	Fraction of patients in CRM stage 1. Default is 0.4.
q_1	Target probability of observing at least one toxicity in CRM stage 1. Default is $0.9$ .
lim_sup_prob	Maximum probability of toxicity the model allows. Default is 0.7.
<pre>p_tox_init_3_3 delta_dosis_3_3</pre>	Initial toxicity probability for the 3+3 model. Default is 0.05.
	Step size for dose escalation in the 3+3 model. Default is 0.092.

#### Value

```
A data.frame with one row per method ("3+3" and "CRM2s") and the following columns:

method Design used ("3+3" or "CRM2s")

mean_pat Mean of patients

median_pat Median of patients

mean_mtd Mean of the estimated MTDs
```

sd\_mtd Standard deviation of the estimated MTDs

```
median_mtd Median of the estimated MTDs
min_mtd Minimum of the estimated MTDs
q1_mtd First quartile (Q1) of the estimated MTDs
q3_mtd Third quartile (Q3) of the estimated MTDs
max_mtd Maximum of the estimated MTDs
siqr_mtd Semi-interquartile range of the estimated MTDs
mean_tox Mean number of toxicities
sd_tox Standard deviation of the number of toxicities
median_tox Median number of toxicities
min_tox Minimum number of toxicities
q1_tox First quartile (Q1) of the number of toxicities
q3_tox Third quartile (Q3) of the number of toxicities
max_tox Maximum number of toxicities
siqr_tox Semi-interquartile range of the number of toxicities
```

## **Examples**

```
df <- run_simulation_logistic(num_rep = 100)
head(df)</pre>
```

 $\verb"run_simulation_power" \textit{ Run Simulation Comparison}$ 

Run Simulation Comparison Between 3+3 and CRM2s for the power model

#### **Description**

Compares the MTD estimation and toxicity count between the power 3+3 method and the two-stage CRM using multiple replications.

## Usage

```
run_simulation_power(
 num_rep = 1000,
  seed = 1234,
  save_plot = FALSE,
  p0 = 0.4,
  theta_0 = 2.7,
  theta = 3,
 N = 24,
 n_{initial} = 1,
 q_0 = 0.05
 q_2 = 0.5,
 q_1 = 0.9
 p_{tox_init_3_3} = 0.02,
 delta_dosis_3_3 = 0.055,
  fixed_optimal_dose = 0.2032
)
```

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#### **Arguments**

num_rep	Number of replications to run. Default is 1000.	
seed	Base random seed for reproducibility. Default is 1234.	
save_plot	Logical. If TRUE, saves comparison plots as PDF. Default is FALSE.	
p0	Target toxicity probability. Default is 0.4.	
theta_0	Nominal value of theta used in CRM and 3+3 escalation models. Default is 2.7.	
theta	True theta used for generating the MTD reference. Default is 3.	
N	Total number of patients in the CRM design. Default is 24.	
n_initial	Number of patients per dose level in the CRM design (not used in 3+3). Default	
	is 1.	
q_0	Initial toxicity probability for CRM design. Default is 0.05.	
q_2	Fraction of patients in CRM stage 1. Default is 0.5.	
q_1	Target probability of observing at least one toxicity in CRM stage 1. Default is $0.9$ .	
<pre>p_tox_init_3_3 delta_dosis_3_3</pre>	Initial toxicity probability for the 3+3 model. Default is 0.02.	
	Step size for dose escalation in the 3+3 model. Default is 0.055.	
fixed_optimal_dose		
	Reference dose for optimal CRM design. Default is 0.2032.	

#### Value

```
A data. frame with one row per method ("3+3" and "CRM2s") and the following columns:
method Design used ("3+3" or "CRM2s")
mean_pat Mean of patients
median_pat Median of patients
mean_mtd Mean of the estimated MTDs
sd_mtd Standard deviation of the estimated MTDs
median_mtd Median of the estimated MTDs
min_mtd Minimum of the estimated MTDs
q1_mtd First quartile (Q1) of the estimated MTDs
q3_mtd Third quartile (Q3) of the estimated MTDs
max_mtd Maximum of the estimated MTDs
siqr_mtd Semi-interquartile range of the estimated MTDs
mean_tox Mean number of toxicities
sd_tox Standard deviation of the number of toxicities
median_tox Median number of toxicities
min tox Minimum number of toxicities
q1_tox First quartile (Q1) of the number of toxicities
q3_tox Third quartile (Q3) of the number of toxicities
max_tox Maximum number of toxicities
siqr_tox Semi-interquartile range of the number of toxicities
```

#### **Examples**

```
df <- run_simulation_power(num_rep = 100)
head(df)</pre>
```

```
simulate_across_n_initial
```

Simulation Study for Different n\_initial Values (Power and Logistic Models)

#### **Description**

Runs simulations using run\_simulation\_power() and run\_simulation\_logistic() with varying n\_initial values and summarizes key results.

#### Usage

```
simulate_across_n_initial(num_rep = 500, seed = 1234)
```

## **Arguments**

num\_rep Number of repetitions for each simulation. Default is 500.

seed Random seed. Default is 1234.

#### Value

A list of two data frames: \$power and \$logistic, each containing results for CRM2s at n\_initial = 1:4 and 3+3.

#### **Examples**

```
result_list <- simulate_across_n_initial()
head(result_list$power)</pre>
```

```
two_stage_crm_logistic
```

C-optimal based two-stage Continual Reassessment Method (CRM2s) Simulation for logistic model with 2 parameters

#### **Description**

Performs one simulation run of a two-stage CRM design for estimating the Maximum Tolerated Dose (MTD) in phase I trials.

## Usage

```
two_stage_crm_logistic(
  p0 = 0.4,
  theta = c(-3, 2),
  theta_0 = c(-3.1, 1.8),
  N = 24,
  n_initial = 3,
  q_0 = 0.05,
  q_2 = 0.4,
```

```
q_1 = 0.9,
lim_sup_prob = 0.7,
show_plot = FALSE,
seed = 1234
)
```

#### **Arguments**

p0	Target toxicity probability. Default is 0.4.
theta	True value for the vector of parameters for the dose-toxicity curve. Default is $c(-3,2)$ .
theta_0	Nominal value of the vector of parameters. Default is c(-3.1, 1.8).
N	Total number of patients (including both stages). Default is 24.
n_initial	Number of patients per dose in stage 1. Default is 3.
q_0	Toxicity probability at first dose. Default is 0.02.
q_2	Fraction of patients in stage 1. Default is 0.5.
q_1	Target probability of observing at least one toxicity during stage 1. Default is 0.9.
lim_sup_prob	Maximum acceptable probability of toxicity. Default is 0.7.
show_plot	Logical. If TRUE, plots the dose levels for patients in the trial. Default is FALSE.
seed	Random seed for reproducibility. Default is 1234.

## Value

A list with:

**n\_toxicities** Total number of toxicities observed.

**mtd\_estimated** Estimated Maximum Tolerated Dose (MTD). If no toxicity is observed in the first stage, the MTD is set to the largest dose level used.

**mle\_theta** Vector of estimated parameters  $\theta = (\alpha, \beta)$  for the logistic dose-toxicity model. Set to NA if no toxicity is detected in stage 1.

- x Vector of dose levels administered.
- y Vector of toxicity outcomes (1 = toxic, 0 = non-toxic).

### Note

If no toxicity is observed during the first stage of the trial (i.e., sum(y) == 0), the simulation is terminated. A warning is issued, and the MTD is conservatively estimated as the highest dose level reached. The value of mle\_theta is set to NA in this case.

## **Examples**

```
result <- two_stage_crm_logistic(show_plot = TRUE)
print(result$mtd_estimated)</pre>
```

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 $\begin{array}{ll} {\sf two\_stage\_crm\_power} & {\it C-optimal\ based\ two-stage\ Continual\ Reassessment\ Method\ (CRM2s)} \\ {\it Simulation\ for\ power\ model} \end{array}$ 

## Description

Performs one simulation run of a two-stage CRM design for estimating the Maximum Tolerated Dose (MTD) in phase I trials.

## Usage

```
two_stage_crm_power(
  p0 = 0.4,
  theta = 3,
  theta_0 = 2.7,
  N = 24,
  n_initial = 3,
  q_0 = 0.02,
  q_2 = 0.5,
  q_1 = 0.9,
  fixed_optimal_dose = 0.2032,
  show_plot = FALSE,
  seed = 1234
)
```

## **Arguments**

p0	Target toxicity probability. Default is 0.4.	
theta	True value for the dose-toxicity curve. Default is 3.	
theta_0	Nominal value of the dose-toxicity curve parameter (initial guess). Default is 2.7.	
N	Total number of patients (including both stages). Default is 24.	
n_initial	Number of patients per dose in stage 1. Default is 3.	
q_0	Toxicity probability at first dose. Default is 0.02.	
q_2	Fraction of patients in stage 1. Default is 0.5.	
q_1	Target probability of observing at least one toxicity during stage 1. Default is $0.9$ .	
fixed_optimal_dose		
	Reference dose for optimal design estimation. Default is 0.2032.	
show_plot	Logical. If TRUE, plots the dose levels for patients in the trial. Default is FALSE.	
seed	Random seed for reproducibility. Default is 1234.	

## Value

A list with:

**n\_toxicities** Total number of toxicities observed.

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**mtd\_estimated** Estimated Maximum Tolerated Dose (MTD). If no toxicity is observed in the first stage, the MTD is set to the largest dose level used.

**mle\_theta** Estimated value of the dose-toxicity parameter  $\theta$ . Set to NA if no toxicity is detected in stage 1.

- x Vector of dose levels administered.
- y Vector of toxicity outcomes (1 = toxic, 0 = non-toxic).

#### Note

If no toxicity is observed during the first stage of the trial (i.e., sum(y) == 0), the simulation is terminated. A warning is issued, and the MTD is conservatively estimated as the highest dose level reached. The value of mle\_theta is set to NA in this case.

## **Examples**

```
result <- two_stage_crm_power(show_plot = TRUE)
print(result$mtd_estimated)</pre>
```

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