

Package ‘EWOC.Comb’

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Title Escalation with Overdose Control using 2 Drug Combinations

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Description

Implements Escalation With Overdose Control trial designs using two drug combinations described by this paper <[doi:10.1002/sim.6961](https://doi.org/10.1002/sim.6961)>(Tighiouart et al., 2016). It calculates the recommended dose for next cohorts and perform simulations to obtain operating characteristics.

License GPL (>= 2)

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 EWOC .Comb-package

Escalation with Overdose Control using 2 Drug Combinations

Description

Implements Escalation With Overdose Control trial designs using two drug combinations described by this paper <doi:10.1002/sim.6961>(Tighiouart et al., 2016). It calculates the recommended dose for next cohorts and perform simulations to obtain operating characteristics.

Author(s)

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References

Tighiouart M, Li Q and Rogatko A. A Bayesian adaptive design for estimating the maximum tolerated dose curve using drug combinations in cancer phase I clinical trials. *Statistics in Medicine*. 2017, 36: 280-290.

 ewoc2

Escalation With Overdose Control for two drugs combination

Description

Finding the doses of next cohort for a phase I clinical trial based on Escalation with Overdose Control (EWOC) design considering the classic parametrization for binary response and two agents.

Usage

```
ewoc2(dose.a, dose.b, resp, theta, alpha, Min.Dose.A, Max.Dose.A, Min.Dose.B, Max.Dose.B,
a01, b01, a10, b10, a00, b00, a, b, delta1x, delta1y, burn, mm, delta1)
```

```
## Default S3 method:
```

```
ewoc2(dose.a, dose.b, resp, theta, alpha, Min.Dose.A, Max.Dose.A, Min.Dose.B, Max.Dose.B,
a01, b01, a10, b10, a00, b00, a, b, delta1x, delta1y, burn=4000, mm=2000, delta1=0.05)
```

Arguments

| | |
|--------|--|
| dose.a | a numeric vector of allowable doses for drug A |
| dose.b | a numeric vector of allowable doses for drug B |
| resp | a numeric vector of allowable responses, 0 or 1 |
| theta | a numeric value defining the proportion of expected patients to experience a medically unacceptable, dose-limiting toxicity (DLT) if administered the MTD. |

| | |
|------------|--|
| alpha | a numerical value defining the probability that dose selected by EWOC is higher than the MTD. |
| Min.Dose.A | a numeric value defining the lower bound of the support of the MTD for drug A |
| Max.Dose.A | a numeric value defining the upper bound of the support of the MTD for drug A |
| Min.Dose.B | a numeric value defining the lower bound of the support of the MTD for drug B |
| Max.Dose.B | a numeric value defining the upper bound of the support of the MTD for drug B |
| a01 | a numeric value for beta prior distribution associated with parameter rho01 |
| b01 | a numeric value for beta prior distribution associated with parameter rho01 |
| a10 | a numeric value for beta prior distribution associated with parameter rho10 |
| b10 | a numeric value for beta prior distribution associated with parameter rho10 |
| a00 | a numeric value for beta prior distribution associated with parameter rho00 |
| b00 | a numeric value for beta prior distribution associated with parameter rho00 |
| a | a numeric value for gamma prior distribution associated with parameter eta |
| b | a numeric value for the gamma prior distribution associated with the parameter eta |
| delta1x | Maximum dose escalation at each step for drug A, the default is $0.2*(Max.Dose.A - Min.Dose.A)$ if not assigned) |
| delta1y | Maximum dose escalation at each step for drug B, the default is $0.2*(Max.Dose.B - Min.Dose.B)$ if not assigned) |
| burn | Number of iterations for adaption, see n.adapt in jags.model for detail |
| mm | Number of iterations to monitor, see n.iter in code.samples for detail |
| delta1 | Threshold for toxicity |

Value

| | |
|------------|---|
| data | a data frame containing the current doses and responses set |
| parameters | list of input parameters |
| priors | list of prior parameters |
| nextdose.x | the next recommended doses for drug A |
| nextdose.y | the next recommended doses for drug B |

References

Tighiouart M, Li Q and Rogatko A. A Bayesian adaptive design for estimating the maximum tolerated dose curve using drug combinations in cancer phase I clinical trials. *Statistics in Medicine*. 2017, 36: 280-290.

Examples

```
test = ewoc2(dose.a=c(0,0),dose.b=c(0,0),resp=c(0,0),theta=0.33,alpha=0.25,
Min.Dose.A=0, Max.Dose.A=1, Min.Dose.B=0, Max.Dose.B=1,a01=1,b01=1,a10=1,b10=1,
a00=1,b00=1,a=0.8,b=0.0384)
print(test)
```

ewoc2simu

*Generic EWOC2 simulation***Description**

Generic function for simulating EWOC trials for 2 drugs combination

Usage

```
ewoc2simu(ntrials, nsamples, type, trho00, trho01, trho10, teta, nx, ny, tp,
Min.Dose.A, Max.Dose.A, Min.Dose.B, Max.Dose.B, alpha, theta, vai, a01,
b01, a10, b10, a00, b00, a, b, delta1x, delta1y, burn, mm, delta1, seed)

## Default S3 method:
ewoc2simu(ntrials, nsamples, type, trho00, trho01, trho10, teta, nx, ny, tp,
Min.Dose.A, Max.Dose.A, Min.Dose.B, Max.Dose.B, alpha, theta, vai, a01,
b01, a10, b10, a00, b00, a, b, delta1x, delta1y, burn=4000, mm=2000, delta1=0.05, seed)
```

Arguments

| | |
|------------|--|
| ntrials | a number indicating the number of trials to be simulated |
| nsamples | a number indicating the number of patients enrolled for each clinical trial |
| type | a character indicating the type of design, could be 'continuous' or 'discrete' or their initials |
| trho00 | a numeric value indicating the true value of the parameter rho00, the probability of DLT when the levels of drugs A and B are both 0 |
| trho01 | a numeric value indicating the true value of the parameter rho01, the probability of DLT when the levels of drugs A and B are 0 and 1, respectively |
| trho10 | a numeric value indicating the true value of the parameter rho10, the probability of DLT when the levels of drugs A and B are 1 and 0, respectively |
| teta | a numeric value indicating the true value of the eta, the interaction parameter |
| nx | a numeric value indicating the number of dose levels for drug A. It's only necessary if type = 'discrete' |
| ny | a numeric value indicating the number of dose levels for drug B. It's only necessary if type = 'discrete' |
| tp | a numerical vector indicating the true probabilities of DLT at each dose combinations, the order is by Drug B first, only necessary if type = 'discrete' |
| Min.Dose.A | a numeric value defining the lower bound of the support of the MTD for drug A |
| Max.Dose.A | a numeric value defining the upper bound of the support of the MTD for drug A |
| Min.Dose.B | a numeric value defining the lower bound of the support of the MTD for drug B |
| Max.Dose.B | a numeric value defining the upper bound of the support of the MTD for drug B |
| alpha | a numerical value defining the probability that dose selected by EWOC is higher than the MTD. |

| | |
|---------|--|
| theta | a numeric value defining the proportion of expected patients to experience a medically unacceptable, dose-limiting toxicity (DLT) if administered the MTD. |
| vai | a numeric value indicating variable alpha increment for each new cohort |
| a01 | a numeric value for beta prior distribution associated with parameter rho01 |
| b01 | a numeric value for beta prior distribution associated with parameter rho01 |
| a10 | a numeric value for beta prior distribution associated with parameter rho10 |
| b10 | a numeric value for beta prior distribution associated with parameter rho10 |
| a00 | a numeric value for beta prior distribution associated with parameter rho00 |
| b00 | a numeric value for beta prior distribution associated with parameter rho00 |
| a | a numeric value for gamma prior distribution associated with parameter eta |
| b | a numeric value for gamma prior distribution associated with parameter eta |
| delta1x | Maximum dose escalation at each step for drug A, the default is $0.2 * (\text{Max.Dose.A} - \text{Min.Dose.A})$ if not assigned) |
| delta1y | Maximum dose escalation at each step for drug B, the default is $0.2 * (\text{Max.Dose.B} - \text{Min.Dose.B})$ if not assigned) |
| burn | Number of iterations for adaption, see n.adapt in jags.model for detail |
| mm | Number of iterations to monitor, see n.iter in code.samples for detail |
| delta1 | Threshold for toxicity |
| seed | a numeric value used in random number generation |

Value

| | |
|------------|--|
| type | same as input parameter type |
| parameters | list of input parameters |
| priors | list of prior parameters |
| Dose.A | a matrix ntrials x nsamples containing the doses of drug A assigned for each patient in a trial and each trial in the simulation |
| Dose.B | a matrix ntrials x nsamples containing the doses of drug B assigned for each patient in a trial and each trial in the simulation |
| Resp | a matrix ntrials x nsamples containing ones and zeros indicating the occurrence of DLT (1) and the absence of DLT (0) for each patient in the trial and each trial in the simulation |
| rho00 | a numeric vector ntrials x 1 containing the estimated rho00 parameter for each trial in the simulation |
| rho01 | a numeric vector ntrials x 1 containing the estimated rho01 parameter for each trial in the simulation |
| rho10 | a numeric vector ntrials x 1 containing the estimated rho10 parameter for each trial in the simulation |
| eta | a numeric vector ntrials x 1 containing the estimated eta parameter for each trial in the simulation |

| | |
|----------|--|
| postlow | a matrix ntrials x nsamples/2 containing posterior probability of DLT at lower doses (both 0 for drug A and B) at each step in a trial and each trial in the simulation |
| postdlts | a matrix (nx x ny x ntrials) x 4 containing posterior probability of DLT at each dose combination sets in each trial in the simulation. This is used to test whether or not a discrete set of MTDs was selected from a continuous MTD curve is kept or dropped. It's available only when type = 'discrete' |

References

Tighiouart M, Li Q and Rogatko A. A Bayesian adaptive design for estimating the maximum tolerated dose curve using drug combinations in cancer phase I clinical trials. *Statistics in Medicine*. 2017, 36: 280-290.

Examples

```
# continous
test1 = ewoc2simu(ntrials=10, nsamples=40, type="c", rho00=0.01, rho01=0.2, rho10=0.9, eta=20,
Min.Dose.A=0, Max.Dose.A=1, Min.Dose.B=0, Max.Dose.B=1, alpha=0.25, theta=0.20, a01=1, b01=1,
a10=1, b10=1, a00=1, b00=1, a=0.8, b=0.0384)

print(test1)
plot(test1, type="MTD")
plot(test1, type="bias")
plot(test1, type="percent")

# discrete
tp = c(0.03, 0.05, 0.08, 0.05, 0.08, 0.13, 0.08, 0.13, 0.2, 0.13, 0.2, 0.29, 0.2, 0.29, 0.4, 0.29, 0.4, 0.53)
test2 = ewoc2simu(ntrials=10, nsamples=40, type="d", nx=6, ny=3, tp=tp,
Min.Dose.A=0, Max.Dose.A=1, Min.Dose.B=0, Max.Dose.B=1, alpha=0.25, theta=0.20,
a01=1, b01=1, a10=1, b10=1, a00=1, b00=1, a=0.8, b=0.0384)

print(test2)
plot(test2, type="MTD")
plot(test2, type="percent")
```

mtdcurve

Generating MTD curve based on logistic model for two drugs

Description

Generating MTD curve based on logistic model for two drugs

Usage

```
mtdcurve(rho00, rho01, rho10, eta, theta)
```

Arguments

| | |
|-------|---|
| rho00 | a numeric value indicating the true value of the parameter rho00, the probability of DLT when the levels of drugs A and B are both 0 |
| rho01 | a numeric value indicating the true value of the parameter rho01, the probability of DLT when the levels of drugs A and B are 0 and 1, respectively |
| rho10 | a numeric value indicating the true value of the parameter rho10, the probability of DLT when the levels of drugs A and B are 1 and 0, respectively |
| eta | a numeric value indicating the true value of the eta, the interaction parameter |
| theta | a numerical value defining the proportion of expected patients to experience a medically unacceptable, dose-limiting toxicity (DLT) if administered the MTD |

Value

a plot showing the MTD curve based on the logistic model

Examples

```
mtdcurve(rho00=0.01, rho01=0.2, rho10=0.9, eta=20, theta=0.2)
```

pdl_t

Generating probability of DLT based on the EWOC2 model

Description

Generating probability of DLT based on the EWOC 2 drugs combination model

Usage

```
pdl_t(rho00, rho01, rho10, eta, theta, x, y)
```

Arguments

| | |
|-------|---|
| rho00 | a numeric value indicating the true value of the parameter rho00, the probability of DLT when the levels of drugs A and B are both 0 |
| rho01 | a numeric value indicating the true value of the parameter rho01, the probability of DLT when the levels of drugs A and B are 0 and 1, respectively |
| rho10 | a numeric value indicating the true value of the parameter rho10, the probability of DLT when the levels of drugs A and B are 1 and 0, respectively |
| eta | a numeric value indicating the true value of the eta, the interaction parameter |
| theta | a numerical value defining the proportion of expected patients to experience a medically unacceptable, dose-limiting toxicity (DLT) if administered the MTD |
| x | a numeric value of dose level for drug A |
| y | a numeric value of dose level for drug B |

Value

a numeric value indicating the probability of DLT with doses from input based on the logistic model

Examples

```
pdlt(rho00=0.01, rho01=0.2, rho10=0.9, eta=20, theta=0.2, x=0.2, y=0.3)
```

```
plot.ewoc2simu
```

```
EWOC for 2 drugs combination trial design characteristics
```

Description

Function to plot the trial design characteristics from EWOC 2 drugs combination simulation results

Usage

```
## S3 method for class 'ewoc2simu'
plot(x, type = "MTD", conf.reg=0.9, plot.figure="Y",...)
```

Arguments

| | |
|-------------|---|
| x | an object of class "ewoc2simu", usually a result of a call to ewoc2simu |
| type | a character indicating the type of plots a user requests, could be "MTD", "bias", or "percent". For discrete simulations, "bias" is not available |
| conf.reg | confidence level that controls the region of the doses from the last trial in the MTD plot |
| plot.figure | a character indicating whether user wants the plot, 'Y' would be yes, otherwise would be no. It's mainly for internal uses |
| ... | arguments passed to or from methods |

Value

No return value, called for side effects.

References

Tighiouart M, Li Q and Rogatko A. A Bayesian adaptive design for estimating the maximum tolerated dose curve using drug combinations in cancer phase I clinical trials. *Statistics in Medicine*. 2017, 36: 280-290.

Examples

```

# continous
test1 = ewoc2simu(ntrials=10, nsamples=40, type="c", trho00=0.01, trho01=0.2, trho10=0.9, teta=20,
Min.Dose.A=0, Max.Dose.A=1, Min.Dose.B=0, Max.Dose.B=1, alpha=0.25, theta=0.20, a01=1, b01=1,
a10=1, b10=1, a00=1, b00=1, a=0.8, b=0.0384)

print(test1)
plot(test1, type="MTD")
plot(test1, type="bias")
plot(test1, type="percent")

# discrete
tp = c(0.03, 0.05, 0.08, 0.05, 0.08, 0.13, 0.08, 0.13, 0.2, 0.13, 0.2, 0.29, 0.2, 0.29, 0.4, 0.29, 0.4, 0.53)
test2 = ewoc2simu(ntrials=10, nsamples=40, type="d", nx=6, ny=3, tp=tp,
Min.Dose.A=0, Max.Dose.A=1, Min.Dose.B=0, Max.Dose.B=1, alpha=0.25, theta=0.20,
a01=1, b01=1, a10=1, b10=1, a00=1, b00=1, a=0.8, b=0.0384)

print(test2)
plot(test2, type="MTD")
plot(test2, type="percent")

```

print.ewoc2

Summarizing EWOC2 next doses results

Description

Summarizing EWOC2 next doses result

Usage

```

## S3 method for class 'ewoc2'
print(x, ...)

```

Arguments

x an object of class "ewoc2", usually, a result of a call to ewoc2
... arguments passed to or from methods

Value

a data.frame of 2 x 4 with columns for cohort, patients, recommended dose of drug A and recommended dose of drug B for next cohort or 2 patients

References

Tighiouart M, Li Q and Rogatko A. A Bayesian adaptive design for estimating the maximum tolerated dose curve using drug combinations in cancer phase I clinical trials. *Statistics in Medicine*. 2017, 36: 280-290.

Examples

```
test = ewoc2(dose.a=c(0,0),dose.b=c(0,0),resp=c(0,0),theta=0.33,alpha=0.25, Min.Dose.A=0,
Max.Dose.A=1, Min.Dose.B=0, Max.Dose.B=1,a01=1,b01=1,a10=1,b10=1,a00=1,b00=1,a=0.8,b=0.0384)
print(test)
```

```
print.ewoc2simu
```

Summarizing EWOC2 simulation results

Description

Summarizing EWOC2 simulation results

Usage

```
## S3 method for class 'ewoc2simu'
print(x, ...)
```

Arguments

x an object of class "ewoc2simu", usually, a result of a call to ewoc2simu
... arguments passed to or from methods

Value

a data.frame of 7 x 1 with row represent Accuracy square discrepancy (sq), Accuracy absolute discrepancy (abs), Accuracy overdose (od), percent Selection, Average percent DLT, percent Trials with DLT rate > theta+0.05, percent Trials with LDT rate > theta+0.1

References

Tighiouart M, Li Q and Rogatko A. A Bayesian adaptive design for estimating the maximum tolerated dose curve using drug combinations in cancer phase I clinical trials. *Statistics in Medicine*. 2017, 36: 280-290.

Examples

```
# continous
test1 = ewoc2simu(ntrials=10, nsamples=40, type="c", trho00=0.01, trho01=0.2, trho10=0.9, teta=20,
Min.Dose.A=0, Max.Dose.A=1, Min.Dose.B=0, Max.Dose.B=1, alpha=0.25, theta=0.20, a01=1, b01=1,
a10=1, b10=1, a00=1, b00=1, a=0.8, b=0.0384)

print(test1)
plot(test1, type="MTD")
plot(test1, type="bias")
plot(test1, type="percent")
```

```
# discrete
tp = c(0.03,0.05,0.08,0.05,0.08,0.13,0.08,0.13,0.2,0.13,0.2,0.29,0.2,0.29,0.4,0.29,0.4,0.53)
test2 = ewoc2simu(ntrials=10, nsamples=40, type="d", nx=6, ny=3, tp=tp,
Min.Dose.A=0, Max.Dose.A=1, Min.Dose.B=0, Max.Dose.B=1, alpha=0.25, theta=0.20,
a01=1,b01=1,a10=1,b10=1,a00=1,b00=1,a=0.8,b=0.0384)

print(test2)
plot(test2, type="MTD")
plot(test2, type="percent")
```

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