Package 'BDP2'

July 31, 2018

Type Package

Title Bayesian Adaptive Designs for Phase II Trials with Binary
Endpoint

Version 0.1.3

Date 2018-07-31

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Description Tools and workflow to choose design parameters in Bayesian adaptive single-
arm phase II trial designs with binary endpoint (response, success) with possible stopping for ef-
ficacy and futility at interim analyses. Also contains routines to determine and visualize operat-
ing characteristics. See Kopp-Schneider et al. (2018) <doi:10.1002/bimj.201700209>.

License GPL-2

Depends rmarkdown, shiny, shinyBS

LazyData TRUE

Suggests knitr

VignetteBuilder knitr

NeedsCompilation no

Repository CRAN

Date/Publication 2018-07-31 12:30:06 UTC

R topics documented:

1

BDP2-package   *Bayesian Adaptive Designs for Phase II Trials with Binary Endpoint*

Description

Tools and workflow to choose design parameters in Bayesian adaptive single-arm phase II trial designs with binary endpoint (response, success) with possible stopping for efficacy and futility at interim analyses. Also contains routines to determine and visualize operating characteristics. See Kopp-Schneider et al. (2018) <doi:10.1002/bimj.201700209>.

Details

The DESCRIPTION file:

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Type:	Package
Title:	Bayesian Adaptive Designs for Phase II Trials with Binary Endpoint
Version:	0.1.3
Date:	2018-07-31
Author:	Annette Kopp-Schneider, Manuel Wiesenfarth, Ulrich Abel
Maintainer:	Manuel Wiesenfarth <m.wiesenfarth@dkfz.de>
Description:	Tools and workflow to choose design parameters in Bayesian adaptive single-arm phase II trial designs wit
License:	GPL-2
Depends:	rmarkdown, shiny, shinyBS
LazyData:	TRUE
Suggests:	knitr
VignetteBuilder:	knitr

Index of help topics:

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Tools and workflow to choose design parameters in Bayesian adaptive single-arm phase II trial designs with binary endpoint (response, success) with possible stopping for efficacy and futility at interim analyses. Also contains routines to determine and visualize operating characteristics.

Main functions: BDP2workflow, BDP2 and plotBDP2. BDP2workflow provides an interactive shiny app which also generates Word/pdf/html reports.

Author(s)

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References

Kopp-Schneider, A., Wiesenfarth, M., Witt, R., Edelmann, D., Witt, O. and Abel, U. (2018). Monitoring futility and efficacy in phase II trials with Bayesian posterior distributions - a calibration approach. *Biometrical Journal*, to appear.

Examples

```
## Not run:
# Starts Shiny app
  BDP2workflow()

## End(Not run)
```

BDP2   *Operating characteristics of a single-arm trial with a binary endpoint*

Description

Determines the operating characteristics of a single-arm trial with a binary endpoint (response, success) and interim efficacy and futility analyses. Declaration of efficacy and futility (including possibly early stopping) is based on the posterior probability that the true response rate is at least pE , pF respectively.

Usage

```
BDP2(n, interim.at, ptrue,
     eff.stop = FALSE,
     pF, cF, pE = NULL, cE = NULL,
     type="PostProb", alpha=0.05,
     shape1F, shape2F, shape1E = NULL, shape2E = NULL,
     simulate = FALSE, nsim = 10000)
```

Arguments

n	sample size at the final analysis
interim.at	vector of sample sizes at the interim analyses
ptrue	true (assumed) response rate used for analytical evaluations or simulating the trial
eff.stop	FALSE: No evaluation of efficacy. "call": no stop for efficacy; in this case the program merely calculates the probability that the efficacacy criterion is satisfied (possibly triggering a notification of the DMC and the start of the planning of a subsequent trial). "stop": the study ends if the efficacy criterion is reached at an interim analysis.
pF	response rate used for the futility criterion (may be identical to pE)
cF	critical level of posterior probabilities used for declaring futility
pE	response rate used for the efficacy criterion
cE	critical level of posterior probabilities used for declaring efficacy
type	"PostProb" for decisions based on posterior probabilities (default) or "PredictivePower" for decisions based on predictive power (currently only implemented for simulate==TRUE)
alpha	significance level for final test (only for simulate==TRUE & type=="PredictivePower")
shape1F	first parameter of the Beta prior for futility analysis
shape2F	second parameter of the Beta prior for futility analysis
shape1E	first parameter of the Beta prior for efficacy analysis
shape2E	second parameter of the Beta prior for efficacy analysis
simulate	FALSE for analytical evaluation and TRUE for simulation
nsim	number of simulation runs (only used if simulate==TRUE)

Details

Assumptions: Endpoint (response/no response) data available for all study patients. Beta-binomial model. Prior distribution = Beta(shape1, shape2).

Decisions based on posterior probabilities:
The posterior distribution at interim analysis with n.int patients and k.int successes is Beta(k.int + shape1F, n.int + shape2F - k.int) and Beta(k.int + shape1E, n.int + shape2E - k.int), respectively. Efficacy is declared if the posterior probability P(true response rate > pE) is >= cE. Futility is declared if the posterior probability P(true success rate > pF) is < cF. cF, cE translate into futility/efficacy boundaries (maximum number of responses leading to early termination for futility/ minimum number of responses leading to declaring of, or early termination for, efficacy).

Decisions based on predictive power: Given the results of the interim analysis, the predictive power at the final analysis (n patients, critical number of successes k.crit) is P(X >= k.crit - k.int), where X follows a beta-binomial distribution with parameters n'= n - n.int, a = k.int + shape1, and b = n.int - k.int + shape2.
Efficacy is declared if the predictive power is >= cE (cE must be high, e.g. 0.70). Futility is declared if the predictive power is < cF (cF must be small, e.g. 0.10). cE, cF translate into futility/efficacy boundaries (maximum number of responses leading to early termination for futility/ minimum number of responses leading to declaring of, or early termination for, efficacy).

References

Kopp-Schneider, A., Wiesenfarth, M., Witt, R., Edelmann, D., Witt, O. and Abel, U. (2018). Monitoring futility and efficacy in phase II trials with Bayesian posterior distributions - a calibration approach. *Biometrical Journal*, to appear.

Examples

```
# Operating characteristics with calling for efficacy
BDP2(n=20, interim.at = c(3,9,13,18), ptrue = 0.3,
     eff.stop = "call",
     pF=0.3, cF=0.01, pE=0.12, cE = 0.9,
     type="PostProb",
     shape1F=0.3, shape2F=0.7, shape1E=0.12, shape2E=0.88)

# Operating characteristics with stopping for efficacy
BDP2(n=20, interim.at = c(3,9,13,18), ptrue = 0.3,
     eff.stop = "stop",
     pF=0.3, cF=0.01, pE=0.12, cE = 0.9,
     type="PostProb",
     shape1F=0.3, shape2F=0.7, shape1E=0.12, shape2E=0.88)
```

BDP2workflow *Shiny app for workflow*

Description

Starts a shiny app in the web browser. It provides a workflow to choose design parameters single-arm trial with a binary endpoint (response, success) and interim efficacy and futility analyses as well as routines to determine and visualize operating characteristics. Also Word/pdf/html reports can be generated.

Usage

```
BDP2workflow(display.mode = "normal")
```

Arguments

display.mode   display.mode passed to shiny::runApp()

References

Kopp-Schneider, A., Wiesenfarth, M., Witt, R., Edelmann, D., Witt, O. and Abel, U. (2018). Monitoring futility and efficacy in phase II trials with Bayesian posterior distributions - a calibration approach. *Biometrical Journal*, to appear.

Examples

```
## Not run:
# Starts Shiny app
  BDP2workflow()

## End(Not run)
```

BDP2_simulate	*Simulated operating characteristics of a single-arm trial with a binary*
	endpoint for different types of futility criteria

Description

NOTE: Usually function [BDP2](#) will be preferred.

Determines the operating characteristics of a single-arm trial with a binary endpoint (success - failure) and interim futility analyses. The user can choose among 10 futility criteria, which are based on predictive or conditional power (the latter either assuming H1 or the MLE), posterior or predictive probabilities, tail probabilites (under H0 or H1), constant success rates, or arbitrary user-defined futility bounds.

Assumptions: Endpoint (success/no success) data available for all study patients. In case of Bayesian analysis: Beta-binomial model. Prior distribution = Beta(shape1,shape2) (uniform if prior.mean=0.5,prior.sampleSize=2) In case of predictive or conditional power, tail probabilities, or rates: One-sided testing in the final analysis.

Usage

```
BDP2_simulate(n, vn.int, p, p0, p1,
              alpha = 0.05, crit, type = 5, nsim,
              shape1 = 1, shape2 = 1)
```

Arguments

n	sample size at the final analysis
vn.int	vector of sample sizes at the interim analyses (the vector may be equal to 1:(n-1) = continuous monitoring of futility)
p	true (assumed) success rate used for simulating the trial
p0	success rate corresponding to H0
p1	success rate corresponding to H1 (p1 > p0)
alpha	nominal probability of type 1 error used for the final test
crit	critical level(s) of predictive/conditional power, (posterior) probabilities, rates, or patient numbers used for early termination. crit translates into futility boundaries (maximum number of successes leading to early termination). If simple rates are used for monitoring futility (type=7), crit is the critical success rate. Rates or probabilites must be input as percentages (???). In case of type 10 analyses, crit must be a vector of numbers of successes indicating futility bounds at each (interim or final)analysis.

type	type of futility analysis: 1=predictive power; 2=conditional power under H1; 3=conditional power under the MLE; 4=posterior probability of a success rate > p0; 5=posterior probability of a success rate >= p1; 6=predictive probability of reaching a "positive" final study result, where "positive" is defined in terms of the posterior probability of a success rate > p0. (Here, the ciritcal level must be >= 1-alpha). 7=estimated success rate; 8=p-values; 9="p-values under H1"; 10=user-defined bounds (a vector of critical numbers of successes for each analysis)
nsim	number of simulation runs
shape1	shape parameter for prior distribution
shape2	shape parameter for prior distribution

Details

Some methodological details on the 10 types of futility criteria:

Type 1: Predictive power. Given the results of the interim analysis the predictive power at the final analysis (n patients, critical number of successes k.crit) is $P(X \geq k.crit - k.int)$, where X follows a beta-binomial distribution with parameters n'= n - n.int, a = k.int + shape1, and b= n.int - k.int + shape2.

Type 2,3: Conditional power. The conditional power at the interim analysis is $P(X \geq k.crit - k.int)$, where X follows a binomial distribution with parameters n'= n - n.int, and success probability either equal to p1 (futility analysis type 2) or to the estimated success rate (MLE) at the interim analysis (type 3.)

Type 4,5:Posterior probabilities. The posterior distribution at interim analysis with n.int patients and k.int successes is Beta(k.int + shape1, n + shape2 - k.int) Type 4: Futility is declared if the posterior probability P(true success rate > p0) is < crit. (Here, crit must be large,e.g. 70%). Type 5: Futility is declared if the posterior probability P(true success rate >= p1) is < crit. (Here, crit must be small, e.g. 10%).

Type 6: Predictive probability combined with posterior probability. Futility is declared if the posterior predictive probability that the study will be a success is < crit (e.g. 10%). Here, the success is defined by the total number of successes in the trial yielding a posterior probability of at least 1 - alpha (when evaluated in the final analysis) that the true success rate is > p0.

Type 7: Estimated success rates. Futility is declared if the success rate is smaller than a fixed benchmark crit. The final analysis is test-based.

Type 8,9: Tail probabilites under H0,H1. Type 8: The futility criterion uses an alpha level crit that is constant across all interim analysis. The final analysis is test-based. Futility is declared if the p-value (upper tail) is >= crit. Type 8 futility analyses should only be used if the number of patients at the first interim analysis is not too low (say, at least 5 to 10). The value of crit is not identicalt o the alpha level used in the final test. Generally, a fairly high value of crit will be appropriate (e.g. 70%). Type 9: Similar to type 8, but with lower-tail probabilites calculated under H1 ("p-values under H1"). I.e., futility is declared if, under H1, the probability of obtaining at most as many successes as the observed number is < crit ("observed number of successes "too low" to be compatible with H1 at one-sided significance level = crit). Generally, a small value of crit (e.g. 5% or 10%) should be chosen.

Type 10: User-defined boundaries. Here, the futility boundaries (maximum numbers of successes leading to early termination) are directly input by the user. crit ist the vector of these boundaries at

each (interim or final) analysis. The study is terminated if the number of successes is at analysis no. m is <= the crit[m].

Examples

```
BDP2_simulate(n=30, vn.int=c(10,20),
              p=0.2, p0=0.1, p1=0.3, crit=0.1, type=5, nsim=1000,shape1=1,shape2=1)
```

---

crit_general   *Calculates CritBoundaries for different types of futility criteria*

Description

Calculates CritBoundaries for different types of futility criteria

Usage

```
crit_general(n, p0, p1, vn.int, alpha, crit, type=5, shape1 = 1, shape2 = 1)
```

Arguments

n	sample size at the final analysis
p0	success rate corresponding to H0
p1	success rate corresponding to H1 (p1 > p0)
vn.int	vector of sample sizes at the interim analyses (the vector may be equal to 1:(n-1) = continuous monitoring of futility)
alpha	nominal probability of type 1 error used for the final test
crit	critical level(s) of predictive/conditional power, (posterior) probabilities, rates, or patient numbers used for early termination. crit translates into futility boundaries (maximum number of successes leading to early termination). If simple rates are used for monitoring futility (type=7), crit is the critical success rate. Rates or probabilites must be input as percentages. In case of type 10 analyses, crit must be a vector of numbers of successes indicating futility bounds at each (interim or final)analysis.
type	see [BDP2_simulate](#) for details. Type of futility analysis: 1=predictive power; 2=conditional power under H1; 3=conditional power under the MLE; 4=posterior probability of a success rate > p0; 5=posterior probability of a success rate >= p1; 6=predictive probability of reaching a "positive" final study result, where "positive" is defined in terms of the posterior probability of a success rate > p0. (Here, the ciritcal level must be >= 1- alpha). 7=estimated success rate; 8=p-values; 9="p-values under H1"; 10=user- defined bounds (a vector of critical numbers of successes for each analysis)
shape1	for prior distribution
shape2	for prior distribution

---

pFstop	*Operating characteristics of a single-arm trial with a binary endpoint with futility stopping*

Description

Determines the operating characteristics of a single-arm trial with a binary endpoint (success - failure) on the basis of analytical derivations. The design allows for futility stopping. Outputs are

- probability of futility stopping (P.futil) at each interim analysis
- cumulative stopping probability up to the interim (P.futil.cum).

Usage

```
pFstop(p, vn.int, v.crit)
```

Arguments

p	true response rate
vn.int	vector of sample sizes at the interim analyses
v.crit	vector of critical boundaries for futility stopping at the interim analyses (stop for futility if number of successes <= boundary). Choose boundary=-1 if no stopping is allowed.

Examples

```
pFstop(p=0.3,vn.int=c(3,9,13,18,20),v.crit=c(0,1,2,3,3))
```

---

pFstopEcall	*operating characteristics of a single-arm trial with a binary endpoint with futility stopping and calling efficacy at interim*

Description

Determines the operating characteristics of a single-arm trial with a binary endpoint (success - failure) on the basis of analytical derivations. The design allows for futility stopping and evaluates efficacy at interim. Outputs are

- probability of calling efficacy (P.effic) at each interim analysis
- cumulative probability of calling efficacy up to the interim (P.effic.cum).
- probability of futility stopping (P.futil) at each interim analysis
- cumulative stopping probability up to the interim (P.futil.cum).

Usage

```
pFstopEcall(p, vn.int, v.critE, v.critF)
```

Arguments

p	true response rate
vn.int	vector of sample sizes at the interim analyses
v.critE	vector of critical boundaries for calling efficacy at the interim analyses (call efficacy if number of successes >= boundary)
v.critF	vector of critical boundaries for futility stopping at the interim analyses (stop for futility if number of successes <= boundary). Choose boundary=-1 if no stopping for futility is allowed.

Examples

```
pFstopEcall(p=0.3,vn.int=c(3,9,13,18,20),v.critE=4:8, v.critF=c(0,1,2,3,3))
```

---

pFstopEstop	*Operating characteristics of a single-arm trial with a binary endpoint with stopping for futility and stopping for efficacy*

Description

Determines the operating characteristics of a single-arm trial with a binary endpoint (success - failure) on the basis of analytical derivations. The design allows for stopping for futility and stopping for efficacy. Outputs are

- probability of stopping for efficacy (P.effic) at each interim analysis
- cumulative probability of stopping for efficacy up to the interim (P.effic.cum).
- probability of futility stopping (P.futil) at each interim analysis
- cumulative stopping probability up to the interim (P.futil.cum).

Usage

```
pFstopEstop(p, vn.int, v.critE, v.critF)
```

Arguments

p	true response rate
vn.int	vector of sample sizes at the interim analyses
v.critE	vector of critical boundaries for calling efficacy at the interim analyses (stop for efficacy if number of successes >= boundary)
v.critF	vector of critical boundaries for futility stopping at the interim analyses (stop for futility if number of successes <= boundary). Choose boundary=-1 if no stopping for futility is allowed.

Examples

```
pFstopEstop(p=0.3,vn.int=c(3,9,13,18,20),v.critE=4:8, v.critF=c(0,1,2,3,3))
```

---

plot.cE_vs_pEcall   *Plot objects returned by plotBDP2()*

Description

Plot objects returned by [plotBDP2](#) with arguments x="cE" and either y="PEcall" or y="PEstop" which can computationally be relatively expensive.

Usage

```
## S3 method for class 'cE_vs_pEcall'
plot(x, ...)
## S3 method for class 'cE_vs_pEstop'
plot(x, ...)
```

Arguments

x   Object returned by [plotBDP2](#) with arguments x="cE" and either y="PEcall" or
            y="PEstop"

...   arguments passed to [plot.default](#)

Examples

```
shape1F=0.3
shape2F=0.7
shape1E=0.12
shape2E=0.88
res=plotBDP2(x="cE",y="PEcall",n=30,interim.at=15,
             pF=0.3,cF=0.01,pE=0.12,cE=seq(.5,1,by=.01),p0=0.3,p1=0.12,
             shape1F=shape1F,shape2F=shape2F,shape1E=shape1E,shape2E=shape2E,
             col=c("green","red"),cex.sub=.8)
plot(res)
```

---

plotBDP2   *Plots*

Description

Output of desired plots

Usage

```
plotBDP2(x = c("n", "k", "ptrue", "cE", "cF"),
         y = c("Prob0Successes", "PostProb0or1Successes", "bFbE",
               "PEcall_p0_p1", "PEstop_p0_p1",
               "PFstopEcall", "PFstopEstop",
               "PEcall", "PEstop", "PFstop", "PFstopEstop",
               "ExpectedNumber",
               "PredictivePower"),
         n, interim.at, ptrue,
         pF, cF, pE, cE, p0, p1, Estop=FALSE,
         shape1F, shape2F, shape1E = NULL, shape2E = NULL,
         col = c("green", "red"), cex.legend=1, add = FALSE,
         show=TRUE, progress = FALSE, ...)
```

Arguments

x
: character string specifying what is given by the x axis

  - "n" for number of patients at final or at first interim, depending on plot
  - "ptrue" for the true response rate
  - "k" number of successes at interim (only for y=="PredictivePower")
  - "cE" for the critical level of posterior probabilities used for declaring efficacy
  - "cF" for the critical level of posterior probabilities used for declaring futility

y
: character string specifying what is given by the y axis

  - "Prob0Successes" for plot of probability of 0 successes out of n at first interim, i.e. $(1-p)^n$. Two curves are generated: one for p0 and one for p1. Choose x="n" and set n to a vector of number patients at first interim.
  - "PostProb0or1Successes" generates two curves of posterior probability that reponse rate exceeds cF. One given 0 successes and one given 1 success observed in n, i.e. P(p>pF| 0 successes out of n) and P(p>pF| 1 success out of n). Choose x="n" and set n to a vector of number patients at first interim.
  - "bFbE" for plot of boundaries for futility bF and for efficacy bE in terms of number of successes per n number of patients. Futility stop if number of successes <= bF, call efficacy if number of successes >= bE. Choose x="n" and set n to the maximal number of patients (not a vector).
  - "PEcall_p0_p1" for plot of probability to call efficacy at final (i.e. P(p>pE|Data)>=cE), evaluated for data generated with control response rate p0 (corresponds to type I error) and for data generated with target response rate p1 (corresponds to power) for varying number of patients at final. Choose x="n" and set n to a vector of number patients at final.
  - "PEstop_p0_p1" as above but with stopping for efficacy instead of calling efficacy
  - "PFstopEcall" for plotting operating characteristics for a given response rate as a function of number of patients at final, n. Shows one curve (default: in red) for the probability of stopping for futility up to final analysis and one curve (default: in green) for the probability of calling efficacy at final

analysis. For the setting ptrue=p0, the curves show the probability of true stopping (default: in red) and type I error (default: in green). For the setting ptrue=p1, the curves show the probability of false stopping (default: in red) and power (default: in green). Choose x="n" and set n to a vector of number patients at final.

- "PFstopEstop" as above but with stopping for efficacy instead of calling efficacy
- "PEcall" for plot of probability to call efficacy at final (i.e. P(p>pE|Data)>=cE).
  - For x="ptrue" then this is the power function.
  - For x="cE" this gives plots of probability to call efficacy at final, evaluated for data generated with control response rate p0 (corresponds to type I error) and for data generated with target response rate p1 (corresponds to power).
- "PEstop"
  - For x="ptrue" then this is the power function.
  - For x="cE" this gives plots of probability to call efficacy at final, evaluated for data generated with control response rate p0 (corresponds to type I error) and for data generated with target response rate p1 (corresponds to power).
- "PFstop" for plot of cumulative probability to stop for futility up to final.
  - For x="ptrue" this gives the futility stopping probability as function of ptrue.
  - For x="cF" this gives 2 curves, evaluated for data generated with control response rate p0 (corresponds to true stopping probability, default: in green) and for data generated with target response rate p1 (corresponds to false stopping probability, default: in red).
- "ExpectedNumber" for expected number of patients in the trial. Choose x= "ptrue". Takes stopping for efficacy into account if Estop==TRUE
- "PredictivePower" for predictive power (only for x=="k")

n	sample size at the final analysis, vector if x=="n"
interim.at	vector of sample sizes at the interim analyses
ptrue	true (assumed) response rate used for simulating the trial, vector if x=="ptrue"
pF	response rate used for the futility criterion P(p>pF|Data) < cF (may be identical to pE)
cF	critical level of posterior probabilities used for declaring futility, vector if x=="cF"
pE	response rate used for the efficacy criterion P(p>pE|Data) >= cE
cE	critical level of posterior probabilities used for declaring efficacy, vector if x=="cE"
p0	response rate corresponding to H0
p1	response rate corresponding to H1 (p1 > p0)
shape1F	first parameter of the Beta prior for futility analysis
shape2F	second parameter of the Beta prior for futility analysis
shape1E	first parameter of the Beta prior for efficacy analysis

shape2E	second parameter of the Beta prior for efficacy analysis
Estop	Stop for efficacy? Defaults to FALSE. Only relevant if y=="ExpectedNumber".
col	line color, for some plots vector of length 2.
add	add line to existing plot. Only supported if x=="ptrue"
show	show plot (otherwise computed objects are invisibly returned)
progress	only used by shiny app
cex.legend	size of legend text relative to cex
...	additional arguments passed to plot.default()

References

Kopp-Schneider, A., Wiesenfarth, M., Witt, R., Edelmann, D., Witt, O. and Abel, U. (2018). Monitoring futility and efficacy in phase II trials with Bayesian posterior distributions - a calibration approach. *Biometrical Journal*, to appear.

Examples

```
# See vignette for more details and examples
pF=0.3
pE=0.12
shape1F=0.3
shape2F=0.7
shape1E=0.12
shape2E=0.88
cF=0.01
cE=0.9
nvec=c(18:40)
interim.at=c(10,20,30)

# Type I error and probability of true stopping for the uninteresting response rate.
ptrue=0.12
plotBDP2(x="n", y="PFstopEcall",
        n =nvec, interim.at = interim.at,
        pF=pF,cF=cF,pE=pE,cE=cE,ptrue=ptrue,
        shape1F=shape1F,shape2F=shape2F,shape1E=shape1E,shape2E=shape2E)


# Power and probability of false stopping for the target response rate.
ptrue=0.3
plotBDP2(x="n", y="PFstopEcall",
        n =nvec, interim.at = interim.at,
        pF=pF,cF=cF,pE=pE,cE=cE,ptrue=ptrue,
        shape1F=shape1F,shape2F=shape2F,shape1E=shape1E,shape2E=shape2E)
```

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