Package 'BE'

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Title Bioequivalence Study Data Analysis

Description Analyze bioequivalence study data with industrial strength. Sample size could be determined for various crossover designs, such as 2x2 design, 2x4 design, 4x4 design, Balaam design, Two-sequence dual design, and William design.

Reference: Chow SC, Liu JP. Design and Analysis of Bioavailability and Bioequivalence Studies. 3rd ed. (2009, ISBN:978-1-58488-668-6).

Depends R (>= 3.0.0), rtf

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R topics documented:

BE-package	2
be2x2	2
ci2cv	3
ci2mse	4
cv2mse	5
hodges	5
mse2cv	6
NCAResult4BE	7
plot2x2	7
pow2x2ci	8
pow2x2mse	9
powcv	0
powmse	.1
scaledBound	2
ss2x2ci	3
sscv	3
ssmse	5

2 be2x2

BE-package	Bioequivalence Study Data Analysis	
Index		19

Description

Analyze bioequivalence study data with industrial strength. Sample size could be determined for various crossover designs, such as 2x2 design, 2x4 design, 4x4 design, Balaam design, Two-sequence dual design, and William design. Basic assumption is that the variable is distributed as a log-normal distribution. This is SAS PROC GLM style. If you want PROC MIXED style, use nlme::lme.

Details

It performs bioequivalency tests for several variables of a 2x2 study in a data file.

Author(s)

Kyun-Seop Bae <k@acr.kr>

References

- 1. Chow SC, Liu JP. Design and Analysis of Bioavailability and Bioequivalence Studies. 3rd ed. (2009, ISBN:978-1-58488-668-6)
- 2. Hauschke D, Steinijans V, Pigeot I. Bioequivalence Studies in Drug Development. (2007, ISBN:978-0-470-09475-4)
- 3. Diletti E, Hauschke D, Steinijans VW. Sample size determination for bioequivalence assessment by means of confidence intervals. Int J Clinical Pharmacol Ther Tox. 1991;29(1):1-8

Examples

```
# write.csv(NCAResult4BE, "temp.csv", quote=FALSE, row.names=FALSE)
# be2x2("temp.csv", c("AUClast", "Cmax", "Tmax"))
```

be2x2

Bioequivalence test of a 2x2 study

Description

It performs conventional bioequivalence test for 2x2 study. Input is a file. Basic assumption is that the variable is distributed as a log-normal distribution. This is SAS PROC GLM style. If you want PROC MIXED style, use nlme::lme.

Usage

```
be2x2(Data, Columns = c("AUClast", "Cmax", "Tmax"), rtfName="")
```

ci2cv 3

Arguments

Data A data.frame or a csv file name. This should have at least the following

columns and variable column(s) to be tested. AUC and Cmax should be all

positive values.

GRP : Group or Sequence, 'RT' or 'TR'

PRD : Period, 1 or 2 SUBJ : Subject ID

TRT : Treatment or Drug, 'R' or 'T'

Column names of variables to be tested. This is usually c("AUClast", "Cmax",

"Tmax") or c("AUClast", "AUCinf", "Cmax", "Tmax")

rtfName Output filename of rich text format(rtf)

Details

It performs bioequivalency tests for several variables of a 2x2 study in a data file. If you specify output filename in rtfName, the output will be saved in the file.

Value

Returns text output of equivalence test result.

Author(s)

Kyun-Seop Bae <k@acr.kr>

See Also

```
test2x2, plot2x2
```

Examples

```
be2x2(NCAResult4BE, c("AUClast", "Cmax", "Tmax"))
```

ci2cv

Coefficient of variation (CV) from a confidence interval of previous 2x2 study

Description

It calculates coefficient of variation (CV) from a confidence interval of previous 2x2 study.

Usage

```
ci2cv(n1, n2, LL, UL, Alpha = 0.1)
```

Arguments

n2 Subject count of group 2 LL Lower limit of the confidence interval of geometric mean ratio (Test/Reference) UL Upper limit of the confidence interval of geometric mean ratio (Test/Reference) Alpha Alpha level. This means (1 - alpha/2)*100 % confidence interval is given	n1	Subject count of group 1
UL Upper limit of the confidence interval of geometric mean ratio (Test/Reference)	n2	Subject count of group 2
	LL	Lower limit of the confidence interval of geometric mean ratio (Test/Reference)
Alpha Alpha level. This means (1 - alpha/2)*100 % confidence interval is given	UL	Upper limit of the confidence interval of geometric mean ratio (Test/Reference)
	Alpha	Alpha level. This means (1 - alpha/2)*100 % confidence interval is given

ci2mse

Details

It calculates coefficient of variation (CV) from a confidence interval of 2x2 bioequivalence study.

Value

Returns coefficient of variation (CV) in percent (%).

Author(s)

Kyun-Seop Bae <k@acr.kr>

Examples

```
ci2cv(12, 13, 0.85, 1.11)
```

ci2mse	Mean squared error (MSE) from a confidence interval of previous 2x2
	study

Description

It calculates mean squared error (MSE) from a confidence interval of previous 2x2 study.

Usage

```
ci2mse(n1, n2, LL, UL, Alpha = 0.1)
```

Arguments

n1	Subject count of group 1
n2	Subject count of group 2
LL	Lower limit of the confidence interval of geometric mean ratio (Test/Reference)
UL	Upper limit of the confidence interval of geometric mean ratio (Test/Reference)
Alpha	Alpha level. This means (1 - alpha/2)*100 % confidence interval is given

Details

It calculates coefficient of variation (CV) from a confidence interval of 2x2 bioequivalence study.

Value

Returns mean squared error (MSE).

Author(s)

Kyun-Seop Bae <k@acr.kr>

```
ci2mse(12, 13, 0.85, 1.11)
```

cv2mse 5

cv2mse

Mean squared error (MSE) from coefficient of variation (CV)

Description

It calculates mean squared error (MSE) from coefficient of variation (CV).

Usage

```
cv2mse(cv)
```

Arguments

C۷

Coefficient of variation (%) in the original scale

Details

Coefficient of variation (CV) is percent in original scale and mean squared error (MSE) is log scale.

Value

Returns mean squared error (MSE) in log scale).

Author(s)

Kyun-Seop Bae <k@acr.kr>

Examples

```
cv2mse(25)
```

hodges

Hodges-Lehmann estimation for a variable of a 2x2 study

Description

It performs Hodges-Lehmann estimation for 2x2 study. This is usually for Tmax variable.

Usage

```
hodges(bedata, Var)
```

Arguments

bedata Data table name. This should have at least the following columns and a variable

column to be tested.

GRP : Group or Sequence, 'RT' or 'TR'

PRD : Period, 1 or 2 SUBJ : Subject ID

TRT : Treatment or Drug, 'R' or 'T'

Var Variable to be estimated. This should be one of the column names in bedata

table. Usually 'Tmax'

6 mse2cv

Details

It nonparametrically tests Var variable equivalency from a 2x2 study. This is done for a variable which we cannot assume log-normal distribution.

Value

```
Wilcoxon Signed-Rank Test
A kind of nonparametric test
```

Hodges-Lehmann Estimate

90% confidence interval in the original scale and the percent scale

Author(s)

Kyun-Seop Bae <k@acr.kr>

Examples

```
hodges(NCAResult4BE, "Tmax")
```

mse2cv

Coefficient of variation (CV) from mean squared error (MSE)

Description

It calculates coefficient of variation (CV) from mean squared error (MSE).

Usage

mse2cv(mse)

Arguments

mse

Mean square error (MSE) in log scale

Details

Coefficient of variation (CV) is percent in the original scale and mean squared error (MSE) is the log scale.

Value

Returns coefficient of variation (CV) in percent (%).

Author(s)

Kyun-Seop Bae <k@acr.kr>

Examples

mse2cv(0.06062462)

NCAResult4BE 7

NCAResult4BE	An Example of Noncompartmental Analysis Result for Bioequivalence Test

Description

Contains a noncompartmental analysis result table from a concentration simulated bioequivalence study.

Usage

NCAResult4BE

Format

A data frame with 48 observations on the following 10 variables.

```
SUBJ Subject ID
```

GRP Group or Sequence character code: 'RT' or 'TR"

PRD Period numeric value: 1 or 2

TRT Treatment or Drug code: 'R' or 'T'

AUClast AUClast positive numeric value

Cmax Positive numeric value

Tmax Tmax positive numeric value

Details

This contains a simulated data for 2x2 bioequivalence study data analysis. Noncompartmental analysis results are from the NonCompart package.

plot2x2

Plot bioequivalence variable of a 2x2 study

Description

It plots two 2x2 plots for a variable.

Usage

```
plot2x2(bedata, Var)
```

Arguments

bedata Data table name. This should have at least the following columns and a variable

column to be plotted.

GRP: Group or Sequence, 'RT' or 'TR'

PRD : Period, 1 or 2 SUBJ : Subject ID

TRT : Treatment or Drug, 'R' or 'T'

Var Variable to be plotted. This should be one of the column names in bedata table.

pow2x2ci

Details

It plots Var column values according to GRP, PRD, TRT.

Value

It just draws two 2x2 plots for equivalence exploration.

Author(s)

Kyun-Seop Bae <k@acr.kr>

Examples

```
plot2x2(NCAResult4BE, "AUClast")
plot2x2(NCAResult4BE, "Cmax")
plot2x2(NCAResult4BE, "Tmax")
```

pow2x2ci

Power using a confidence interval of previous 2x2 study

Description

It calculates power for the bioequivalence test on ratio using a confidence interval of previous 2x2 study.

Usage

```
pow2x2ci(n1, n2, LL, UL, Alpha = 0.1)
```

Arguments

n1	Subject count of group 1
n2	Subject count of group 2
LL	Lower limit of the confidence interval of geometric mean ratio (Test/Reference)
UL	Upper limit of the confidence interval of geometric mean ratio (Test/Reference)
Alpha	Alpha level. This means (1 - alpha/2)*100 % confidence interval is given

Details

It calculates power of sample size (n per group) with CV.

Value

```
Returns power [0, 1)
```

Author(s)

Kyun-Seop Bae <k@acr.kr>

```
pow2x2ci(12, 13, 0.85, 1.11)
```

pow2x2mse 9

pow2x2mse	Power using mean squared error (MSE) of previous 2x2 study

Description

It calculates power for the bioequivalence test on ratio using mean squared error (MSE of previous 2x2 study.

Usage

```
pow2x2mse(n1, n2, mse, True.R = 1, Alpha = 0.1, ThetaL = 0.8, ThetaU = 1.25)
```

Arguments

n1	Subject count of group 1
n2	Subject count of group 2
mse	Mean squared error
True.R	True ratio of test/reference
Alpha	Alpha level. This means (1 - alpha/2)*100 % confidence interval is given
ThetaL	Lower limit of equivalence criteria
ThetaU	Upper limit of equivalence criteria

Details

It calculates power of sample size (n per group) with CV.

Value

```
Returns power [0, 1)
```

Author(s)

```
Kyun-Seop Bae <k@acr.kr>
```

```
pow2x2mse(12, 13, 0.0756530)
```

10 powcy

powcv

Power using coefficient of variation (CV)

Description

It calculates power for the bioequivalence test on ratio using coefficient of variation (CV).

Usage

```
powcv(n, CV, DesignNo = 1, True.R = 1, Alpha = 0.1, ThetaL = 0.8, ThetaU = 1.25)
```

Arguments

n Sample size, n per group
CV Coefficient of Variation (%)
DesignNo Crossover design number.

Design Number (treatment x sequence x period)

1 2x2x2 : RT TR

2 2x4x2 (Balaam Design) : TT RR RT TR

3 2x2x3 (Two-sequence Dual Design): TRR RTT

4 2x2x4 : TRRT RTTR

5 2x4x4 : TTRR RRTT TRRT RTTR

6 3x6x3 (William Design for 3 treatments) + carry-over effect

: RBA ARB BAR ABR BRA RAB

7~3x6x3 (William Design for 3 treatments) - carry-over effect

: RBA ARB BAR ABR BRA RAB

8 4x4x4 (William Design for 4 treatments) + carry-over effect

: RCAB ARBC BACR CBRA

9 4x4x4 (William Design for 4 treatments) - carry-over effect

: RCAB ARBC BACR CBRA

True .R True ratio of test/reference

Alpha error level

ThetaL Lower limit of equivalence criteria
ThetaU Upper limit of equivalence criteria

Details

It calculates power of sample size (n per group) with CV.

Value

Returns power [0, 1)

Author(s)

Kyun-Seop Bae <k@acr.kr>

```
powcv(12, 25)
```

powmse 11

powmse

Power using mean squared error (MSE)

Description

It calculates power for the bioequivalence test on ratio using mean squared error (MSE).

Usage

```
powmse(n, mse, DesignNo = 1, True.R = 1, Alpha = 0.1, ThetaL = 0.8, ThetaU = 1.25)
```

Arguments

n Sample size, n per group
mse Mean squared error
DesignNo Crossover design number.

Design Number (treatment x sequence x period)

1 2x2x2 : RT TR

2 2x4x2 (Balaam Design) : TT RR RT TR

3 2x2x3 (Two-sequence Dual Design): TRR RTT

4 2x2x4 : TRRT RTTR

5 2x4x4 : TTRR RRTT TRRT RTTR

6 3x6x3 (William Design for 3 treatments) + carry-over effect

: RBA ARB BAR ABR BRA RAB

7~3x6x3 (William Design for 3 treatments) - carry-over effect

: RBA ARB BAR ABR BRA RAB

8 4x4x4 (William Design for 4 treatments) + carry-over effect

: RCAB ARBC BACR CBRA

9 4x4x4 (William Design for 4 treatments) - carry-over effect

: RCAB ARBC BACR CBRA

True .R True ratio of test/reference

Alpha error level

ThetaL Lower limit of equivalence criteria
ThetaU Upper limit of equivalence criteria

Details

It calculates power of sample size (n per group) with mse.

Value

Returns power [0, 1)

Author(s)

Kyun-Seop Bae <k@acr.kr>

```
powmse(12, 0.06)
```

12 scaledBound

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Widened Bound for Scaled Average Bioequivalence

Description

It calculates widened bound for scaled average bioequivalence.

Usage

```
scaledBound(CV = 40, k = 0.76, digits = 4)
```

Arguments

CV	coefficient of variation in percent
k	0.76 is for EMA and Korea MFDS. US FDA uses 0.893. When CV is 30%, bound becomes (0.8, 1.25). Most regulartory body does not use a more accurate value.
digits	Korea MFDS use 4 digits only, while other regulatory bodies use more decimal values.

Details

CV must be larger than 30%. If CV is larger than 50

Value

widened bound for scaled average bioequivalence

Author(s)

Kyun-Seop Bae <k@acr.kr>

```
scaledBound(30)
scaledBound(35)
scaledBound(40)
scaledBound(45)
scaledBound(50)
```

ss2x2ci 13

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SS2X2C	1

Sample size using a confidence interval of previous 2x2 study

Description

It calculates sample size for the bioequivalence test on ratio using a confidence interval of previous 2x2 study.

Usage

```
ss2x2ci(n1, n2, LL, UL, Alpha = 0.1)
```

Arguments

n1	Subject count of group 1
n2	Subject count of group 2
LL	Lower limit of the confidence interval of geometric mean ratio (Test/Reference)
UL	Upper limit of the confidence interval of geometric mean ratio (Test/Reference)
Alpha	Alpha level. This means (1 - alpha/2)*100 % confidence interval is given

Details

It calculates sample size (n per group) with CV, Alpha, and Beta for bioequivalence test.

Value

Returns sample size (n per group) for bioequivalence test with ratio criteria.

Author(s)

```
Kyun-Seop Bae <k@acr.kr>
```

Examples

```
ss2x2ci(12, 13, 0.85, 1.11)
```

SSCV

Sample size using coefficient of variation (CV)

Description

It calculates sample size for the bioequivalence test on ratio using coefficient of variation (CV).

Usage

```
sscv(CV, DesignNo = 1, True.R = 1, Alpha = 0.1, Beta = 0.2,
    ThetaL = 0.8, ThetaU = 1.25, nMax = 999999)
```

14 sscv

Arguments

CV Coefficient of Variation (%)

DesignNo Crossover design number.

Design Number (treatment x sequence x period)

1 2x2x2 : RT TR

2 2x4x2 (Balaam Design) : TT RR RT TR

3 2x2x3 (Two-sequence Dual Design): TRR RTT

4 2x2x4 : TRRT RTTR

5 2x4x4 : TTRR RRTT TRRT RTTR

6 3x6x3 (William Design for 3 treatments) + carry-over effect

: RBA ARB BAR ABR BRA RAB

7 3x6x3 (William Design for 3 treatments) - carry-over effect

: RBA ARB BAR ABR BRA RAB

8 4x4x4 (William Design for 4 treatments) + carry-over effect

: RCAB ARBC BACR CBRA

9 4x4x4 (William Design for 4 treatments) - carry-over effect

: RCAB ARBC BACR CBRA

True . R True ratio of test/reference

Alpha error level

Beta error level

ThetaL Lower limit of equivalence criteria

ThetaU Upper limit of equivalence criteria

nMax Maximum subject number (sample size) per group

Details

It calculates sample size (n per group) with CV, Alpha, and Beta for bioequivalence test.

Value

Returns sample size (n per group) for bioequivalence test with ratio criteria.

Author(s)

Kyun-Seop Bae <k@acr.kr>

Examples

sscv(25)

ssmse 15

ssmse

Sample size using mean squared error (MSE)

Description

It calculates sample size for the bioequivalence test on ratio using mean squared error (MSE).

Usage

```
ssmse(mse, DesignNo = 1, True.R = 1, Alpha = 0.1, Beta = 0.2, ThetaL = 0.8, ThetaU = 1.25, nMax = 9999999)
```

Arguments

iguments			
	mse	Mean squared error	
	DesignNo	Crossover design number.	
		Design Number (treatment x sequence x period) 1 2x2x2 : RT TR 2 2x4x2 (Balaam Design) : TT RR RT TR 3 2x2x3 (Two-sequence Dual Design): TRR RTT 4 2x2x4 : TRRT RTTR 5 2x4x4 : TTRR RRTT TRRT RTTR 6 3x6x3 (William Design for 3 treatments) + carry-over effect : RBA ARB BAR ABR BRA RAB 7 3x6x3 (William Design for 3 treatments) - carry-over effect : RBA ARB BAR ABR BRA RAB 8 4x4x4 (William Design for 4 treatments) + carry-over effect : RCAB ARBC BACR CBRA 9 4x4x4 (William Design for 4 treatments) - carry-over effect	
		: RCAB ARBC BACR CBRA	
	True.R	True ratio of test/reference	
	Alpha	Alpha error level	
	Beta	Beta error level	
	ThetaL	Lower limit of equivalence criteria	
	ThetaU	Upper limit of equivalence criteria	

Details

nMax

It calculates sample size (n per group) with mse, Alpha, and Beta for bioequivalence test.

Maximum subject number (sample size) per group

Value

Returns sample size (n per group) for bioequivalence test with ratio criteria.

Author(s)

Kyun-Seop Bae <k@acr.kr>

16 ssscv

Examples

ssmse(0.06)

SSSCV

Sample Size for Scaled Average BE using coefficient of variation (CV)

Description

It calculates sample size for the scaled average bioequivalence test on ratio using coefficient of variation (CV).

Usage

```
ssscv(CV, DesignNo = 1, True.R = 1, Alpha = 0.1, Beta = 0.2, Region = "EU", nMax = 999999)
```

Arguments

- Samenes	
CV	Coefficient of Variation (%)
DesignNo	Crossover design number.
	Design Number (treatment x sequence x period) 1 2x2x2 : RT TR 2 2x4x2 (Balaam Design) : TT RR RT TR 3 2x2x3 (Two-sequence Dual Design): TRR RTT 4 2x2x4 : TRRT RTTR
	5 2x4x4 : TTRR RRTT TRRT RTTR 6 3x6x3 (William Design for 3 treatments) + carry-over effect : RBA ARB BAR ABR BRA RAB
	7 3x6x3 (William Design for 3 treatments) - carry-over effect : RBA ARB BAR ABR BRA RAB
	8 4x4x4 (William Design for 4 treatments) + carry-over effect : RCAB ARBC BACR CBRA
	9 4x4x4 (William Design for 4 treatments) - carry-over effect : RCAB ARBC BACR CBRA
True.R	True ratio of test/reference
Alpha	Alpha error level
Beta	Beta error level

Details

Region

nMax

It calculates sample size (n per group) with CV, Alpha, and Beta for scaled average bioequivalence test. US FDA uses this widened bound for both AUClast and Cmax, while EU EMA and Korea MFDA use this for Cmax only.

US or FDA for US FDA, KR or MFDS for Korea MFDS, EU or EMA for other

Value

Returns sample size (n per group) for scaled average bioequivalence test with ratio criteria.

Maximum subject number (sample size) per group

regions or countries

test2x2

Author(s)

Kyun-Seop Bae <k@acr.kr>

Examples

```
ssscv(42.2, DesignNo=4, True.R=0.9) # 14 per group, EU EMA. This applies only for Cmax
ssscv(42.2, DesignNo=4, True.R=0.9, Region="US") # 9 per group, US FDA
ssscv(42.2, DesignNo=4, True.R=0.9, Region="KR") # 14 per group, Korea MFDS. This applies only for Cmax
```

test2x2

Bioequivalence test for a variable of a 2x2 study

Description

It performs conventional bioequivalence test for 2x2 study. Basic assumption is that the variable is distributed as a log-normal distribution. This is SAS PROC GLM style. If you want PROC MIXED style use nlme::lme.

Usage

```
test2x2(bedata, Var)
```

Arguments

bedata Data table name. This should have at least the following columns and a variable

column to be tested. Var column values should be all positive values.

GRP: Group or Sequence, 'RT' or 'TR'

PRD : Period, 1 or 2 SUBJ : Subject ID

TRT : Treatment or Drug, 'R' or 'T'

Var Variable to be tested. This should be one of the column names in bedata table.

Usually 'AUClast' or 'Cmax'

Details

It tests Var variable equivalency from a 2x2 study. Current regulatory requirement is that the 90% confidence interval of geometric mean ratio (Test/Reference) should be within [0.8, 1.25].

Value

Analysis of Variance (log scale)

Analysis of Variance in log scale

Between and Within Subject Variability

Variance in log scale and coefficient of variance in original scale

Least Square Means

Geometric means

90% Confidence Interval

90% confidence interval of geometric mean ratio (T/R)

Sample Size Sample size for the replication of this study

18 test2x2

Author(s)

Kyun-Seop Bae <k@acr.kr>

Examples

test2x2(NCAResult4BE, "AUClast")
test2x2(NCAResult4BE, "Cmax")

Index

```
* Plot
    plot2x2, 7
* Power
    pow2x2ci, 8
    pow2x2mse, 9
    powcv, 10
    powmse, 11
* \ Report
    be2x2, 2
* Sample Size
    ss2x2ci, 13
    sscv, 13
    ssmse, 15
    ssscv, 16
* Statistical Test
    hodges, 5
    test2x2, 17
*\ datasets
    NCAResult4BE, 7
BE (BE-package), 2
BE-package, 2
be2x2, 2
ci2cv, 3
ci2mse, 4
cv2mse, 5
hodges, 5
mse2cv, 6
NCAResult4BE, 7
plot2x2, 3, 7
pow2x2ci, 8
pow2x2mse, 9
powcv, 10
powmse, 11
scaledBound, 12
ss2x2ci, 13
sscv, 13
ssmse, 15
ssscv, 16
test2x2, 3, 17
```