## Chapter 143

# Superiority by a Margin Tests for the Ratio of Two Within-Subject Variances in a 2×2M Replicated Cross-Over Design

## Introduction

This procedure calculates power and sample size of *superiority by a margin* tests of within-subject variabilities from a 2×2M replicated cross-over design. This routine deals with the case in which the statistical hypotheses are expressed in terms of the ratio of the within-subject variances.

This design is used to compare two treatments which are administered to subjects in different orders. It has two treatment sequences. Here, M is the number of times a particular treatment is received by a subject. For example, if M = 2, the design is a 2×4 cross-over. The two sequences would often be

sequence 1: RTRT sequence 2: TRTR

It is assumed that either there is no carry-over from one measurement to the next, or there is an ample washout period between measurements.

## **Technical Details**

This procedure uses the formulation given in Chow, Shao, Wang, and Lokhnygina (2018).

Suppose  $x_{ijkl}$  is the response in the *i*th sequence (i = 1, 2), *j*th subject (j = 1, ..., Ni), *k*th treatment (k = T, C), and *l*th replicate (l = 1, ..., M). The mixed effect model analyzed in this procedure is

$$x_{ijkl} = \mu_k + \gamma_{ikl} + S_{ijk} + e_{ijkl}$$

where  $\mu_k$  is the kth treatment effect,  $\gamma_{ikl}$  is the fixed effect of the lth replicate on treatment k in the ith sequence,  $S_{ij1}$  and  $S_{ij2}$  are random effects of the ijth subject, and  $e_{ijkl}$  is the within-subject error term which is normally distributed with mean 0 and variance  $V_k = \sigma_{Wk}^2$ .

Unbiased estimators of these variances are found after applying an orthogonal transformation matrix *P* to the x's as follows

$$z_{ijk} = P'x_{ijk}$$

where P is an  $m \times m$  matrix such that P'P is diagonal and  $var(z_{ijkl}) = \sigma_{Wk}^2$ .

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For example, in a 2×4 cross-over design the z's become

$$z_{ijk1} = \frac{x_{ijk1} + x_{ijk2}}{2} = \bar{x}_{ijk}.$$

and

$$z_{ijk2} = \frac{x_{ijk1} + x_{ijk2}}{\sqrt{2}} = \bar{x}_{ijk}.$$

In this case, the within-subject variances are estimated as

$$\hat{V}_T = \frac{1}{(N_1 + N_2 - 2)(M - 1)} \sum_{i=1}^{2} \sum_{j=1}^{N_i} \sum_{l=1}^{M} (z_{ijTl} - \bar{z}_{i.Tl})^2$$

and

$$\hat{V}_C = \frac{1}{(N_1 + N_2 - 2)(M - 1)} \sum_{i=1}^{2} \sum_{j=1}^{N_i} \sum_{l=1}^{M} (z_{ijCl} - \bar{z}_{i.Cl})^2$$

## **Testing Superiority by a Margin**

The following hypotheses are usually used to test for non-inferiority

$$H_0: \frac{\sigma_{WT}^2}{\sigma_{WC}^2} \ge R0$$
 versus  $H_1: \frac{\sigma_{WT}^2}{\sigma_{WC}^2} < R0 < 1$ ,

where R0 is the superiority by a margin limit.

The corresponding test statistics are  $T=(\hat{V}_1/\hat{V}_2)/R0$ . Upon making the usual normality assumptions, T is distributed as an  $F_{d,d}$  random variable where

$$d = (N_1 + N_2 - 2)(M - 1).$$

#### Power

The power of this combination of tests is given by

Power = 
$$P\left(F < \frac{R0}{R1} F_{\alpha,d,d}\right)$$

where F is the common F distribution with the indicated degrees of freedom,  $\alpha$  is the significance level, and R1 is the value of the variance ratio stated by the alternative hypothesis. Lower quantiles of F are used in the equation.

A simple binary search algorithm can be applied to this power function to obtain an estimate of the necessary sample size.

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# **Example 1 - Finding Sample Size**

A company has developed a generic drug for treating rheumatism and wants to show that it is superior (has a smaller variance) to the standard drug. A  $2 \times 4$  replicated cross-over design will be used.

Company researchers set the superiority limit to 0.75, the significance level to 0.05, the power to 0.90, M to 2, and the actual variance ratio values between 0.2 and 0.6. They want to investigate the range of required sample size values assuming that the two group sample sizes are equal.

## Setup

If the procedure window is not already open, use the PASS Home window to open it. The parameters for this example are listed below and are stored in the **Example 1** settings file. To load these settings to the procedure window, click **Open Example Settings File** in the Help Center or File menu.

Solve For	Sample Size
Power	0.90
Alpha	0.05
Sequence Allocation	Equal (N1 = N2)
M (Number of Replicates)	2
R0 (Superiority Variance Ratio)	0.75
R1 (Actual Variance Ratio)	0.2 0.3 0.4 0.5 0.6

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## **Output**

Click the Calculate button to perform the calculations and generate the following output.

#### **Numeric Reports**

#### **Numeric Results**

Solve For: Sample Size Variance Ratio: σ²wτ / σ²wc

Hypotheses: H0:  $\sigma^2 wr / \sigma^2 wc \ge R0$  vs. H1:  $\sigma^2 wr / \sigma^2 wc < R0$ 

Pow	ver.		Sequence ample Siz		Number of	Variance	Ratio	
Target	Actual	N1	N2	N	Replicates M	Superiority R0	Actual R1	Alpha
0.9	0.9181	12	12	24	2	0.75	0.2	0.05
0.9	0.9021	22	22	44	2	0.75	0.3	0.05
0.9	0.9013	45	45	90	2	0.75	0.4	0.05
0.9	0.9009	106	106	212	2	0.75	0.5	0.05
0.9	0.9004	346	346	692	2	0.75	0.6	0.05

**Target Power** The desired power value entered in the procedure. Power is the probability of rejecting a false null hypothesis. **Actual Power** The actual power achieved. Because N1 and N2 are discrete, this value is usually slightly larger than the target power. N<sub>1</sub> The number of subjects in sequence 1. N2 The number of subjects in sequence 2. Ν The total number of subjects. N = N1 + N2. Μ The number of replicates. That is, it is the number of times a treatment measurement is repeated on a subject. R0 The superiority limit for the within-subject variance ratio.

R1 The value of the within-subject variance ratio at which the power is calculated.

Alpha The probability of rejecting a true null hypothesis.

#### **Summary Statements**

A 2x2M replicated cross-over design will be used to test whether the treatment within-subject variance ( $\sigma^2$ wT) is superior to the control within-subject variance ( $\sigma^2$ wc) by a margin, by testing whether the within-subject variance ratio ( $\sigma^2$ wT /  $\sigma^2$ wc) is less than 0.75 (H0:  $\sigma^2$ wT /  $\sigma^2$ wc  $\geq$  0.75 versus H1:  $\sigma^2$ wT /  $\sigma^2$ wc < 0.75). Each subject will alternate treatments (T and C), with an assumed wash-out period between measurements to avoid carry-over. With 2 replicate pairs, each subject will be measured 4 times. For those in the Sequence 1 group, the first treatment will be C, and the sequence is [C T C T]. For those in the Sequence 2 group, the first treatment will be T, and the sequence is [T C T C]. The comparison will be made using a one-sided, variance-ratio F-test (with the treatment within-subject variance in the numerator), with a Type I error rate ( $\sigma^2$ ) of 0.05. To detect a within-subject variance ratio ( $\sigma^2$ vT /  $\sigma^2$ vc) of 0.2 with 90% power, the number of subjects needed will be 12 in Group/Sequence 1, and 12 in Group/Sequence 2.

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#### **Dropout-Inflated Sample Size**

	s	ample S	ze	E	pout-Infl Inrollme ample Si	nt	ı	Expecte Number Dropou	of
Dropout Rate	N1	N2	N	N1'	N2'	N'	D1	D2	D
20%	12	12	24	15	15	30	3	3	6
20%	22	22	44	28	28	56	6	6	12
20%	45	45	90	57	57	114	12	12	24
20%	106	106	212	133	133	266	27	27	54
20%	346	346	692	433	433	866	87	87	174
Dropout Rate	The percentag			) that are exp					
N1, N2, and N	The evaluable	sample si	zes at which		mputed. If	N1 and N2 s	subjects are	evaluate	ed out of
N1', N2', and N'	inflating N1	sed on the and N2 us ded up. (S	assumed di ing the formi ee Julious, S	ropout rate. <i>A</i> ulas N1' = N1 S.A. (2010) p	After solvir	ng for N1 and ) and N2' = N	l N2, N1' ar l2 / (1 - DR)	nd N2' are ), with N1	calcula and N2
D1, D2, and D	The expected	number of	dropouts. D	1 = N1' - N1	D2 = N2'	- N2, and D	= D1 + D2.		

#### **Dropout Summary Statements**

Anticipating a 20% dropout rate, 15 subjects should be enrolled in Group 1, and 15 in Group 2, to obtain final group sample sizes of 12 and 12, respectively.

#### References

Chow, S.C., Shao, J., Wang, H., and Lokhnygina, Y. 2018. Sample Size Calculations in Clinical Research, Third Edition. Taylor & Francis/CRC. Boca Raton, Florida.

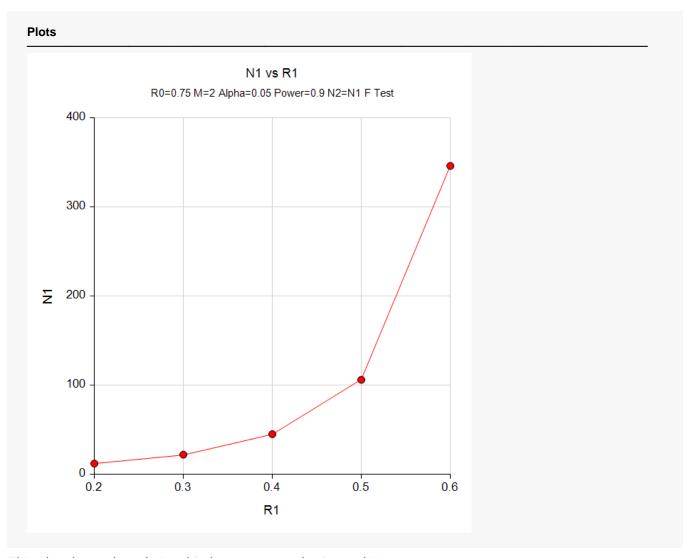
Chow, S.C. 2014. Biosimilars Design and Analysis of Follow-on Biologics, Third Edition. Taylor & Francis/CRC. Boca Raton, Florida.

Chow, S.C., and Liu, J.P. 2014. Design and Analysis of Clinical Trials, Third Edition. John Wiley & Sons. Hoboken, New Jersey.

This report gives the sample sizes for the indicated scenarios.

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#### **Plots Section**



This plot shows the relationship between sample size and R1.

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# **Example 2 - Validation using Hand Calculations**

We could not find an example in the literature, so we will present hand calculations to validate this procedure.

Set N1 to 25, the upper limit of equivalence to 0.75, the significance level to 0.05, M to 2, and the actual variance ratio value 0.5. Compute the power.

The calculations proceed as follows.

Power = 
$$P\left(F < \frac{R0}{R1} F_{\alpha,d,d}\right)$$
  
=  $P\left(F < 0.75/0.5 \left(F_{0.05,48,48}\right)\right)$   
=  $P(F < 1.5(0.61905310))$   
=  $P(F < 0.92857965)$   
=  $0.39923624$ 

Hence, the power is 0.3992 to four decimal places.

## Setup

If the procedure window is not already open, use the PASS Home window to open it. The parameters for this example are listed below and are stored in the **Example 2** settings file. To load these settings to the procedure window, click **Open Example Settings File** in the Help Center or File menu.

Solve For	Power
Alpha	0.05
Sequence Allocation	Equal (N1 = N2)
Sample Size Per Sequence	25
M (Number of Replicates)	2
R0 (Superiority Variance Ratio)	0.75
R1 (Actual Variance Ratio)	0.5

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## **Output**

Click the Calculate button to perform the calculations and generate the following output.

Solve For: Power Variance Ratio: $\sigma^2$ wt / $\sigma^2$ wc Hypotheses: H0: $\sigma^2$ wt / $\sigma^2$				≥ R0 vs. H1: c	σ²wτ / σ²wc < R0		
g		Sequence sample Size		Number of	Variance		
Power	N1	N2	N	Replicates M	Superiority R0	Actual R1	Alpha
		25	50	2	0.75	0.5	0.05

The power matches the hand calculations.