### Chapter 648

# Meta-Analysis of Tests for the Risk Ratio of Two Proportions using a Random-Effects Model in a Cluster-Randomized Design

# Introduction

This procedure calculates power and number of studies for random-effect meta-analyses of clusterrandomized, two-group studies that are summarized by the risk ratios of two group proportions.

### Meta-Analysis

Meta-analysis refers to the systematic assessment of a group of two or more individual studies (experiments or clinical trials). The studies must be similar so that their results may be numerically compared. For example, in this procedure all studies must have as their outcome two group proportions whose risk ratio is used to compare their values.

By combining the results of several studies, their individual results may be summarized and assessed. The power of the combined test will be increased since it represents a larger sample.

#### Fixed-Effects versus Random-Effects Models

The *fixed-effects model*, which is assumed by another procedure, assumes that all studies have the same true effect size. That is, it assumes that all factors that might influence the effect size are the same so that the true effect size for all studies is the same. Therefore, only the *within study variability* needs to be considered since the effect size variation occurs because of random error.

The *random-effects model*, which is assumed by this procedure, assumes that the studies in the analysis are similar, but not to the extent that they all have the same true effect size. Instead, it is assumed that each study has its own true effect size and that the particular outcome of the study is a sample from a distribution with that proportion. The random-effects model must not only account for the *within study variability*, but it must also account for the *between study variability*.

# **Technical Details**

### The Meta-Analysis of the Risk Ratio of Two Proportions

The power formulas used by this procedure are an adaptation of those given in Harrer *et al.* (2022), Campbell and Walters (2014), Ahn, Heo, and Zhang (2015), Donner and Klar (2002), Donner, Piaggio, and Villar (2003), and Eldridge and Kerry (2012) page 204.

Cluster-randomized designs are those in which whole clusters of subjects (classes, hospitals, communities, etc.) are randomized into a treatment group or a control group. It is assumed that each study resulted in two group proportions, made up of  $K_i$  clusters of  $M_{ij}$  subjects each, and was analyzed using summary statistics created from the individual subject data. The sample sizes for each study were adjusted to compensate for the cluster-randomization using a *design effect* (DE) factor.

A series of *H* studies are assumed to each have an effect size of *RR* which is the risk ratio of two group proportions. The results of these individual studies may be combined into one overall statistical test that draws a more precise conclusion.

### **Test Statistic**

Suppose each study has two populations from which dichotomous (binary) responses are recorded. The probability (or risk) of obtaining the event of interest in population 1 (the treatment group) is  $p_1$  and in population 2 (the control group) is  $p_2$ . The corresponding failure proportions are given by  $q_1 = 1 - p_1$  and  $q_2 = 1 - p_2$ .

An assumption is made that the responses from each group follow a binomial distribution. This means that the event probability,  $p_i$ , is the same for all subjects within the group and that the response from one subject is independent of that of any other subject.

Random samples of *m* and *n* individuals are obtained from these two populations. The data from these samples can be displayed in a 2-by-2 contingency table as follows

Group	Success	Failure	Total
Treatment	а	С	т
Control	b	d	п
Total	S	f	Ν

The following alternative notation is sometimes used.

Group	Success	Failure	Total
Treatment	<i>x</i> <sub>11</sub>	<i>x</i> <sub>12</sub>	$n_1$
Control	<i>x</i> <sub>21</sub>	<i>x</i> <sub>22</sub>	$n_2$
Total	$m_1$	$m_2$	Ν

The binomial proportions,  $p_1$  and  $p_2$ , are estimated from these data using the formulae

$$\hat{p}_1 = \frac{a}{m} = \frac{x_{11}}{n_1}$$
 and  $\hat{p}_2 = \frac{b}{n} = \frac{x_{21}}{n_2}$ 

#### **Risk Ratio**

The risk ratio is computed as

$$RR = \frac{p_1}{p_2}$$

#### Meta-Analysis for the Risk Ratio of Proportions

Suppose the individual tables are summarized into one 2-by-2 table whose individual cells now contain the corresponding cell averages which are still named *a*, *b*, *c*, and *d*. These counts can be reparametrized using the two group proportions *P*1 and *P*2 and the two average-group sample sizes *N*1 and *N*2.

These sample sizes are obtained from the cluster-randomized design parameters *K*1, *M*1, *K*2, and *M*2 using the formulas

$$Ni = \frac{Ki(Mi)}{DE_i}, i = 1,2$$

where

$$DE_1 = 1 + ((COV^2 + 1)M_1 - 1)\rho$$

$$DE_2 = 1 + ((COV^2 + 1)M_2 - 1)\rho$$

These are called the *effective* sample sizes. These adjusted sample sizes can then be used in the regular, twogroup formulas given next. The average cell counts are

$$a = (P1)(N1)$$
  

$$b = (P2)(N2)$$
  

$$c = (1 - P1)(N1)$$
  

$$d = (1 - P2)(N2)$$

Using these summary statistics, a summary effect size called the log risk ratio can be calculated using

$$\log(RR) = \log\left(\frac{P1}{P2}\right)$$

along with an estimate of the within-group variance

$$V_W = \frac{1}{a} + \frac{1}{c} - \frac{1}{a+b} - \frac{1}{c+d}$$

For planning purposes, the between-group variance is estimated as a multiple of the within-group variance as follows

$$V_B = RV_W$$

Hedges and Pigott (2001) recommend using 0.333 for a *small* value, 0.667 for a *medium* value, and 1.0 for a *large* value of *R*. Note that the fixed-effect results is achieved when *R* is set to 0.

#### I<sup>2</sup> - Heterogeneity Index

Some authors measure the amount of heterogeneity in the meta-analysis using the heterogeneity index *I*<sup>2</sup> (also called the *degree of heterogeneity* or *inconsistency*) instead of *R*. This value is related to *R* as follows. Note that often this value is presented as a percentage.

$$I^2 = \frac{V_B}{V_B + V_W}$$

This value is related to *R* using the relationship

$$R = \frac{I^2}{1 - I^2}$$

#### **Random-Effects Standard Error**

The final random-effects standard error used in the power calculation is

$$SE_{\log RR} = \sqrt{\frac{V_W + V_B}{H}}$$

The meta-analysis z-test is then calculated as

$$Z = \frac{\log(RR) - \log(RR_0)}{SE_{\log RR}}$$

We assume that *Z* follows the standard normal distribution and so it can be used to test the null hypothesis  $H_0$ :  $RR = RR_0$  at significance level of  $\alpha$ .

#### **Hypotheses**

Three statistical hypotheses commonly used are

 $H_0: RR \le RR_0 \quad \text{vs.} \quad H_1: RR > RR_0$  $H_0: RR \ge RR_0 \quad \text{vs.} \quad H_1: RR < RR_0$  $H_0: RR = RR_0 \quad \text{vs.} \quad H_1: RR \ne RR_0$ 

### **Power Calculation**

The power of a one-tailed test that  $RR = RR_0$  is given by

$$Power = 1 - \Phi(c_{\alpha} - \lambda)$$

where OR is set by the alternative hypothesis,  $c_{\alpha}$  is the  $1 - \alpha$  quantile of the standard normal distribution, and

$$\lambda = \frac{\log(RR) - \log(RR_0)}{SE_{\log RR}}$$

The power of a two-tailed test that  $RR = RR_0$  is given by

$$Power = 1 - \Phi(c_{\alpha/2} - \lambda) + \Phi(-c_{\alpha/2} - \lambda)$$

These formulas can be used in search algorithms to determine the necessary number of studies required to achieve a desired level of power.

Note that the values of P1.0 (P1 assuming H0) and P1.1 (P1 assuming H1) can be obtained from the risk ratios using

$$P1.0 = RR_0P2$$

and

$$P1.1 = RR_1(P2)$$

where  $RR_1$  is the risk ratio at which the power is calculated.

## **Example 1 – Finding the Number of Studies**

Researchers wish to determine a reasonable number of studies for a meta-analysis with the following characteristics: two-group, equal-allocation trials, two-sided significance level of 0.05, power of 0.9, P2 = 0.5,  $RR_1 = 1.1, 1.25, 1.5$ , and  $RR_0 = 1.0$ . The average number of clusters per study in both groups is 7. The average size of each cluster is 8. The coefficient of variation of cluster sizes is 0.65. The intracluster correlation is 0.05. I2 is used to specify the size of the between study variation. It is set at 0.5.

#### 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.

Design Tab	
Solve For	Number of Studies
Alternative Hypothesis	Two-Sided (H1: RR ≠ RR0)
Power	0.90
Alpha	0.05
K1 (Average Number of Clusters)	7
M1 (Average Cluster Size)	8
K2 (Average Number of Clusters)	K1
M2 (Average Cluster Size)	<b>M</b> 1
COV of Cluster Sizes	0.65
RR0 (Risk Ratio H0)	1.0
RR1 (Risk Ratio H1)	1.1 1.25 1.5
P2 (Group 2 Proportion)	0.5
ρ (Intracluster Correlation, ICC)	0.05
Calculate Between-Study Variance From	I <sup>2</sup> (Between-Study Var / Total-Study Var)
l <sup>2</sup> (Between-Study Var / Total-Study Var)	0.5

### Output

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

#### **Numeric Reports**

Solve For: Hypotheses: Calculate Between-Study Variance From:			H0: RR =	Number of Studies H0: RR = RR0 vs. H1: RR ≠ RR0 I² (Between-Study Variance / Total-Study Variance)										
N	Number of	Number of Clusters		Cluster Size		Sample Size R		Risk	Risk Ratio		ortion	Between vs. Total		
	Studies H	Average K	Total HK	Average M	cov	Average N	Total HN	RR H1 RR1	RR H0 RR0	P H1 P.1	P H0 P.0	Variance Ratio I <sup>2</sup>	ICC P	Alpha
0.90062 [Group 1] [Group 2]		14 7 7	1792 896 896	8 8	0.65 0.65	112 56 56	14336 7168 7168	1.10	1	0.550 0.500	0.5 0.5	0.5 0.5	0.05 0.05	0.05
0.90452 [Group 1] [Group 2]	26	14 7 7	364 182 182	8 8	0.65 0.65	112 56 56	2912 1456 1456	1.25	1	0.625 0.500	0.5 0.5	0.5 0.5	0.05 0.05	0.05
0.92090 [Group 1] [Group 2]		14 7 7	168 84 84	8 8	0.65 0.65	112 56 56	1344 672 672	1.50	1	0.750 0.500	0.5 0.5	0.5 0.5	0.05 0.05	0.05
Power [Group] H K HK	ratio. The group The firs The numb The avera count in The total	p to which t line show ber of stuc age number n each gro number of	the res vs the s lies incl er of clu up is sh	ults on thi ettings tha uded in th usters per nown below	s line t at are u e meta study. w on th eta-an	pelong. Gro used for bo analysis. These clus in next two alysis. The	oup 1 is oth grou sters are lines. ese clus	the trea ps. e split be ters are	tment gr etween g	roup an	d group and 2.	ulated at the c 2 is the c . The avera and 2. Th	ontrol age clu	group. uster

M The average cluster size (i.e., number of subjects per cluster) in groups 1 and 2.

COV The coefficient of variation of the cluster sizes of the clusters within the group. This value is used for both groups.

N The average number of subjects per study. The total sample size is shown as the top row followed by N1 and N2.

- HN The total number of subjects represented in the meta-analysis. HN = H x N. The subtotals for groups 1 and 2 are shown below.
- RR1 The value of the risk ratio assumed by the alternative hypothesis, H1. This is the risk ratio at which the power is calculated.
- RR0 The value of the risk ratio assumed by the null hypothesis, H0.
- P.1 The value of the response proportion of groups 1 and 2 assumed by the alternative hypothesis, H1.

P.0 The value of the response proportion of groups 1 and 2 assumed by the null hypothesis, H0.

I<sup>2</sup> The value of the heterogeneity index used to specify the Between Study variance. The formula is (Between Study Variance) = I<sup>2</sup> / (1 - I<sup>2</sup>) x (Within Study Variance). The following values are often used: Small = 0.25, Medium = 0.5, Large = 0.75.

ρ The intracluster correlation (ICC) between any two observations from within a cluster. This statistic has two common interpretations: (1) the proportion of variation related to the between-cluster variation or (2) the correlation between two subjects from the same cluster.

Alpha The probability of rejecting a true null hypothesis.

#### **Summary Statements**

A collection of two-group cluster-randomized-design studies will be used to test whether the Group 1 proportion is different from the Group 2 proportion (H0: RR = 1 versus H1: RR  $\neq$  1, RR = P1 / P2). The meta-analysis comparison will be made using a two-sided random-effects-model Z-test, with a Type I error rate ( $\alpha$ ) of 0.05. In Group 1, the studies are assumed to have an average of 7 clusters with an average of 8 subjects per cluster (totaling 56 subjects), and in Group 2, an average of 7 clusters with an average of 8 subjects per cluster (totaling 56 subjects). The coefficient of variation of the cluster sizes within a group is assumed to be 0.65. The intracluster correlation coefficient of responses within a cluster is assumed to be 0.05. The response proportion of Group 2 (the control group) is assumed to be 0.5. The between-to-total variance ratio (I<sup>2</sup>), used to calculate the between-study variance, is assumed to be 0.5. To detect a risk ratio (RR) of 1.1 with 90% power, the number of needed studies will be 128.

#### References

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Hedges, L.V. and Pigott, T.D. 2001. 'The Power of Statistical Tests in Meta-Analysis', Psychological Methods, Volume 6, No. 3, pages 203-217.

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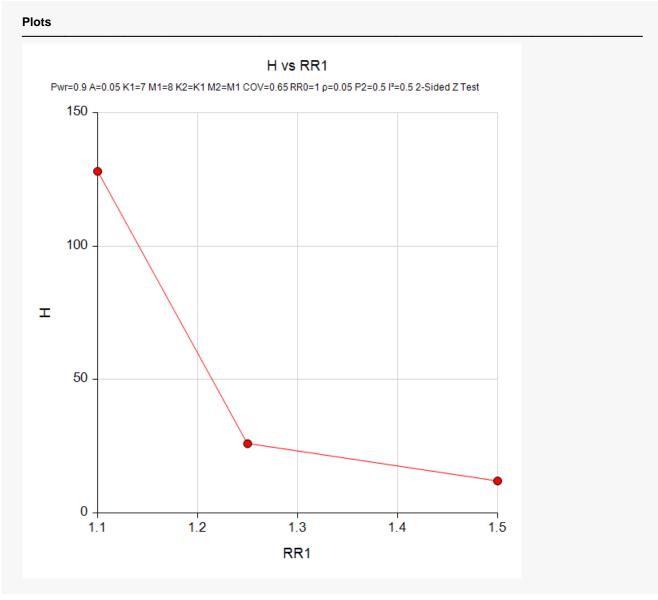
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Valentine, J.C., Pigott, T.D., and Rothstein, H.R. 2010. 'How Many Studies Do You Need? A Primer on Statistical Power for Meta-Analysis', Journal of Education and Behavioral Statistics, Volume 35, No. 2, pages 215-247.

This report shows the sample sizes for the indicated scenarios.

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#### **Plots Section**



This plot represents the required number of studies for various values of RR1.

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

We could not find a validation example for this procedure in the literature so we will validate the procedure using hand calculations.

Suppose a meta-analysis is contemplated with the following characteristics: two-group, equal-allocation trials, two-sided significance level of 0.05 number of studies = 9, average cluster size of all studies is 15, average number of clusters = 10, COV = 0.65,  $\rho = 0.04$ , P2 = 0.5, R = 1,  $RR_1 = 1.2$ , and  $RR_0 = 1.0$ . Calculate the power.

First, we will calculate the power directly using the formulas given above.

Step 1: Find the effective sample sizes.

To do this, first calculate the value of DE. The formula for DE is

$$DE = 1 + \{ [COV^2 + 1]\overline{M} - 1 \} \rho$$

Plugging in COV = 0.65, M = 15, and  $\rho$  = 0.04 gives

$$DE = 1 + \{[0.65^2 + 1]15 - 1\}(0.04) = 1.8135$$

The effective sample sizes are then found as follows.

$$N1 = N2 = \frac{10(15)}{1.8135} = 82.713$$

**Step 2**: Compute the value of *P*1.

P1 = (RR1)P2 = (1.2)0.5 = 0.6

Step 3: Compute the average cell counts.

$$a = (P1)(N1) = 0.6(82.713) = 49.6278$$
  

$$b = (P2)(N2) = 0.5(82.713) = 41.3565$$
  

$$c = (1 - P1)(N1) = 0.4(82.713) = 33.0852$$
  

$$d = (1 - P2)(N2) = 0.5(82.713) = 41.3565$$

**Step 4**: Compute the effect size.

$$\log(RR1) = \log(1.2) = 0.1823216$$

**Step 5**: Compute the within-study variance component.

$$V_W = \frac{1}{a} + \frac{1}{c} - \frac{1}{a+b} - \frac{1}{c+d}$$
$$= \frac{1}{49.6278} + \frac{1}{33.0852} - \frac{1}{49.6278 + 41.3565} - \frac{1}{33.0852 + 41.3565}$$
$$= 0.0259508$$

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**Step 6**: Compute the standard error of the effect size.

$$SE_{\log RR} = \sqrt{\frac{V_W + RV_B}{H}} = \sqrt{\frac{0.0259508 + 0.0259508}{9}} = 0.0759397$$

**Step 7**: Compute *λ*.

$$\lambda = \frac{\log(RR) - \log(RR_0)}{SE_{\log RR}} = \frac{0.1823216 - 0}{0.0759397} = 2.4008714$$

**Step 8**: Compute  $c_{\alpha/2}$ .

$$c_{\alpha/2} = \Phi(1 - 0.025) = 1.95996398$$

Step 9: Compute the power.

$$Power = 1 - \Phi(c_{\alpha/2} - \lambda) + \Phi(-c_{\alpha/2} - \lambda)$$
  
= 1 - \Phi(1.95996398 - 2.4008714) + \Phi(-1.95996398 - 2.4008714)  
= 1 - \Phi(-0.4409074) + \Phi(-4.3608354)  
= 1 - 0.3296400 + 0.0000082  
= 0.6703682

#### 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
Alternative Hypothesis	Two-Sided (H1: RR ≠ RR0)
Alpha	0.05
H (Number of Studies)	9
K1 (Average Number of Clusters)	10
M1 (Average Cluster Size)	15
K2 (Average Number of Clusters)	K1
M2 (Average Cluster Size)	M1
COV of Cluster Sizes	0.65
RR0 (Risk Ratio H0)	1.0
RR1 (Risk Ratio H1)	1.2
P2 (Group 2 Proportion)	0.5
o (Intracluster Correlation, ICC)	0.04
Calculate Between-Study Variance From	mR (Between-Study Var / Within-Study Var

### Output

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

Solve For: Hypotheses: Calculate Between-Study Variance From:			Power H0: RR = RR0 vs. H1: RR ≠ RR0 R (Between-Study Variance / Within-Study Variance)											
	Number of Clusters Number of Studies Average Total H K HK	Clusters		Cluster Size		Sample Size		Risk Ratio		Proportion		Between vs. Within Variance		
Power		Average M	cov	Average N	Total HN	RR H1 RR1	RR H0 RR0	P H1 P.1	P H0 P.0	Ratio Ratio	ICC Ρ	Alpha		
0.67037	9	20	180			300	2700	1.2	1					0.05
[Group 1]		10	90	15	0.65	150	1350			0.6	0.5	1	0.04	
[Group 2]		10	90	15	0.65	150	1350			0.5	0.5	1	0.04	

**PASS** also calculates a power of 0.67037 which validates the procedure.