## Chapter 245

## Non-Inferiority Tests for the Difference Between Two Poisson Rates in a Cluster-Randomized Design

## Introduction

This procedure calculates power and sample size for non-inferiority tests of two rates in a clusterrandomized design in which the outcome variable is a count. It uses the work of Wang, Zhang, and Ahn (2018) which give the power for the case of varying cluster sizes. The analysis uses a simple z-test comparing the two rates.

Cluster-randomized designs are those in which whole clusters of subjects (classes, hospitals, communities, etc.) are put into the treatment group or the control group. Generally speaking, the larger the cluster sizes and the higher the correlation among subjects within the same cluster, the larger will be the overall sample size necessary to detect an effect with the same power.

## **The Statistical Hypotheses**

Non-inferiority tests are examples of directional (one-sided) tests. This program module provides the input and output in formats that are convenient for these types of tests. This section will review the specifics of non-inferiority testing.

Remember that in the usual test setting, the null (H<sub>0</sub>) and alternative (H<sub>1</sub>) hypotheses for one-sided tests are defined as follows, assuming that  $\delta = \mu_1 - \mu_2$  is to be tested.

$$H_0: \delta \leq 0$$
 versus  $H_1: \delta > 0$ 

Rejecting this test implies that the mean difference is larger than the value  $\delta$ . This test is called an *upper-tailed test* because it is rejected in samples in which the difference between the sample means is larger than *D*.

Following is an example of a *lower-tailed test*.

$$H_0: \delta \ge 0$$
 versus  $H_1: \delta < 0$ 

*Non-inferiority* tests are special cases of the above directional tests. It will be convenient to adopt the following specialized notation for the discussion of these tests.

<u>Parameter</u>	PASS Input/Output	<u>Interpretation</u>
$\lambda_1$	$\lambda_1$	<i>Incidence rate</i> of population 1. Population 1 is assumed to consist of those who have received the new treatment.
$\lambda_2$	$\lambda_2$	<i>Incidence rate</i> of population 2. Population 2 is assumed to consist of those who have received the control (standard or reference) treatment.
ε	DO	<i>Margin of non-inferiority.</i> This is a tolerance value that defines the magnitude of the amount that is not of practical importance. This may be thought of as the largest change from the baseline that is considered to be trivial. The absolute value is shown to emphasize that this is a magnitude. The sign of the value will be determined by the specific design that is being used.
δ	D1	<i>True difference</i> . This is the value of $\lambda_1 - \lambda_2$ , the difference between the rates.

## **Non-Inferiority Tests**

A *non-inferiority test* tests that the treatment rate is not worse than the control rate by more than the non-inferiority margin. The actual direction of the hypothesis depends on the nature of the response variable being studied.

### **Case 1: High Values are Better**

In this case, higher values are better. The hypotheses are arranged so that rejecting the null hypothesis implies that the treatment rate is no less than a small amount below the control rate. The value of  $\delta$  is often set to zero. The following are equivalent sets of hypotheses.

$$\begin{split} H_0: \delta &\leq -\varepsilon \quad \text{versus} \quad H_1: \delta > -\varepsilon, \ \varepsilon > 0 \\ H_0: \lambda_1 - \lambda_2 &\leq -\varepsilon \quad \text{versus} \quad H_1: \ \lambda_1 - \lambda_2 > -\varepsilon, \ \varepsilon > 0 \\ H_0: \lambda_1 &\leq \lambda_2 - \varepsilon \quad \text{versus} \quad H_1: \ \lambda_1 > \lambda_2 - \varepsilon, \ \varepsilon > 0 \end{split}$$

#### Case 2: High Values are Worse

In this case, lower values are better. The hypotheses are arranged so that rejecting the null hypothesis implies that the treatment rate is no more than a small amount above the control rate. The value of  $\delta$  is often set to zero. The following are equivalent sets of hypotheses.

$$\begin{split} H_0: \delta &\geq \varepsilon \quad \text{versus} \quad H_1: \delta < \varepsilon, \ \varepsilon > 0 \\ H_0: \lambda_1 - \lambda_2 &\geq \varepsilon \quad \text{versus} \quad H_1: \ \lambda_1 - \lambda_2 < \varepsilon, \ \varepsilon > 0 \\ H_0: \lambda_1 &\geq \lambda_2 + \varepsilon \quad \text{versus} \quad H_1: \ \lambda_1 < \lambda_2 + \varepsilon, \ \varepsilon > 0 \end{split}$$

## **Technical Details**

The following discussion summarizes the results in Wang, Zhang, and Ahn (2018).

Suppose you are interested in comparing the incidence rates of two groups (treatment and control) with a non-inferiority test. Further suppose that the response is known to be related to other covariates (such as age, race, or gender) and so their impact needs to be adjusted for. This may be accomplished by stratifying on the covariates and forming hypotheses about a common mean difference across all clusters and strata. Often, the stratification is based on cluster size, but this is not required.

Let  $Y_{jki}$  be the count outcome of the *i*th  $(i = 1, ..., M_{jk})$  subject in the *k*th  $(k = 1, ..., K_j)$  cluster of the *j*th (j = 1, 2) group. Assuming that  $Y_{iki}$  follows a Poisson model with

$$\lambda_j = \mathrm{E}(Y_{jki}) = \mathrm{Var}(Y_{jki})$$

and a common intracluster correlation coefficient (ICC)  $\rho = \operatorname{corr}(Y_{iki}, Y_{iki'})$  for  $i \neq i'$ .

## **Test Statistic**

An unbiased estimator of  $\lambda_i$  is

$$\hat{\lambda}_{j} = \frac{\sum_{k=1}^{K_{j}} \sum_{i=1}^{M_{jk}} Y_{jki}}{\sum_{k=1}^{K_{j}} M_{jk}}$$

with estimated variance

$$s_j^2 = \frac{\hat{\lambda}_j \sum_{k=1}^{K_j} M_{jk} [1 + (M_{jk} - 1)\hat{\rho}]}{\left(\sum_{k=1}^{K_j} M_{jk}\right)^2}$$

Using a one-sided hypothesis  $H0: \lambda_1 = \lambda_2$  versus  $H1: \lambda_1 < \lambda_2$ , H0 is rejected if

$$\frac{\hat{\lambda}_1 - \hat{\lambda}_2}{\sqrt{s_1^2 + s_2^2}} > z_{1-\alpha}$$

## Sample Size and Power

Wang, Zhang, and Ahn (2018) provide the following formula for estimating  $K_2$  (the number of clusters in the control group) when there are an unequal number of clusters in each group.

$$K_{2} = \frac{\left(z_{1-\frac{\alpha}{2}} + z_{1-\beta}\right)^{2} \left(\frac{\lambda_{1}}{R} + \lambda_{2}\right)}{(\lambda_{1} - \lambda_{2})^{2}} \left[(1-\rho)\frac{1}{\theta} + \rho + \rho\gamma^{2}\right]$$

where  $R = K_1/K_2$ ,  $M = E(M_{jk})$ ,  $\tau^2 = Var(M_{jk})$ , and  $\gamma = \tau/M$ . Here M is the average cluster size of all clusters in the study and  $\gamma$  is the coefficient of variation of the cluster sizes. Note that  $z_{\chi} = \Phi(\chi)$  is the standard normal distribution function.

This equation can easily be rearranged to provide a formula for power  $(1 - \beta)$ .

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

A non-inferiority study is being planned to investigate whether a new intervention will decrease the incidence rate of a certain disease over the rate achieved by the current intervention. This response is a count. For a number of reasons, the researchers decide to administer the intervention to whole clusters (clinics) rather than randomize the treatment to individuals within a cluster. The number of clinics receiving each treatment will be balanced.

The average number of subjects per clinic is 21. The coefficient of variation of the cluster sizes is 0.42.

Prior studies have obtained an incidence rate of 0.35 for the current treatment and an ICC of 0.07. The noninferiority margin is set to 0.05. The researchers want to compare the necessary sample size when the actual incidence rate of the new treatment is 0.3, 0.25, 0.2, and 0.15.

The one-sided significance level is set to 0.025 and the power is set to 0.8.

## 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.

K1 (Number of Clusters in Group 1)
Worse (H1: λ1 - λ2 < D0, where D0 > 0)
0.80
0.025
K1
21
0.42
Differences (D0 and D1)
0.05
0.2 -0.15 -0.1 -0.05
0.35
0.07

## Output

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

## **Numeric Reports**

Numeric Results			
Solve For: Groups:	K1 (Number of Clusters in Group 1) 1 = Treatment, 2 = Control		
Higher Rates Are:			
Hypotheses:	H0: $\lambda 1 - \lambda 2 \ge D0$ vs. H1: $\lambda 1 - \lambda 2 < D0$		
		Incidence Rate	

	Numbo	r of Cluste	ore	Cluster	Sizo	Total	Treatn	nent		Differe	ence		
Power	Treatment K1	Control K2	Total K	Average M	CV	Sample Size N	Non- Inferiority λ1.0	Actual λ1.1	Control λ2	Non- Inferiority D0	Actual D1	Intracluster Correlation ρ	Alpha
0.80060	219	219	438	21	0.42	9198	0.4	0.15	0.35	0.05	-0.20	0.07	0.025
0.80075	239	239	478	21	0.42	10038	0.4	0.20	0.35	0.05	-0.15	0.07	0.025
0.80087	259	259	518	21	0.42	10878	0.4	0.25	0.35	0.05	-0.10	0.07	0.025
0.80098	279	279	558	21	0.42	11718	0.4	0.30	0.35	0.05	-0.05	0.07	0.025

Power	The probability of rejecting a false null hypothesis when the alternative hypothesis is true.
K1, K2, and K	The number of clusters in groups 1, 2, and both, respectively.
M	The average cluster size. It is the anticipated average number of subjects per cluster.
CV	The coefficient of variation of the cluster sizes. It is the standard deviation of cluster sizes divided by the average cluster size.
Ν	The total sample size, i.e., the total number of subjects from all clusters.
λ1.0	The non-inferiority bound on incidence rate in the treatment group.
λ1.1	The actual incidence rate of the treatment group at which the power is computed.
λ2	The incidence rate in the control group.
D0	The difference in incidence rates assumed by the null hypothesis. This is often called the non-inferiority margin. D0 = $\lambda 1.0 - \lambda 2$ .
D1	The difference in incidence rates assumed by the alternative hypothesis. This is the difference at which the power is computed. D1 = $\lambda$ 1.1 - $\lambda$ 2.
ρ	The intracluster correlation coefficient. This is the correlation between any two subjects within a particular cluster.
Alpha	The probability of rejecting a true null hypothesis.

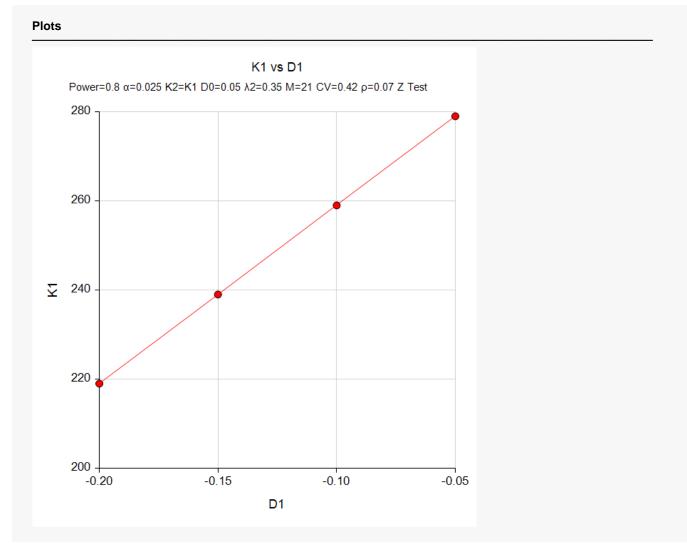
#### Summary Statements

A parallel two-group cluster-randomized design will be used to test whether the Group 1 (treatment) incidence rate ( $\lambda$ 1) is non-inferior to the Group 2 (control) incidence rate ( $\lambda$ 2), with an incidence rate difference non-inferiority margin of 0.05 (H0:  $\delta \ge 0.05$  versus H1:  $\delta < 0.05$ ,  $\delta = \lambda 1 - \lambda 2$ ). The comparison will be made using a one-sided incidence rate difference Z-test with a Type I error rate ( $\alpha$ ) of 0.025. The intracluster correlation coefficient is assumed to be 0.07. The control group incidence rate is assumed to be 0.35. The individual cluster sizes (the number of subjects per cluster) are assumed to vary according to a discrete distribution with mean 21 and coefficient of variation 0.42. To detect an incidence rate difference ( $\lambda$ 1 -  $\lambda$ 2) of -0.2 (or treatment group incidence rate of 0.15) with 80% power, with an average of 21 subjects per cluster, the number of clusters needed will be 219 in Group 1 (treatment) and 219 in Group 2 (control) (for an overall total of 9198 subjects).

#### References

Wang, J., Zhang, S., and Ahn, C. 2018. 'Sample size calculation for count outcomes in cluster randomization trials with varying cluster sizes.' Communications in Statistics--Theory and Methods, DOI: 10.1080/03610926.2018.1532004.

## **Plots Section**



The values from the Numeric Results report are displayed in this plot.

# Example 2 – Validation using a Previously Validated Procedure

We could not find a validation example in the literature so we will use a previously validated procedure in **PASS** to validate this procedure.

Suppose in a non-inferiority design, higher rates are worse, power = 0.9, alpha = 0.025, K2 = K1, M = 50, CV = 0.2,  $\lambda$ 1.0 = 0.6,  $\lambda$ 1.1 = 0.5,  $\lambda$ 2 = 0.5, and  $\rho$  = 0.002. Solve for K1.

This scenario can be solved using the procedure "Tests for the Difference Between Two Poisson Rates in a Cluster-Randomized Design" with the following settings.

Set the alternative hypothesis to "One-Sided (H1:  $\delta > 0$ )", power = 0.9, alpha = 0.025, K2 = K1, M = 50, CV = 0.2,  $\lambda 1 = 0.6$ ,  $\lambda 2 = 0.5$ , and  $\rho = 0.002$ . The solution is K1 = K2 = 26.

## 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.

Design Tab	
Solve For	K1 (Number of Clusters in Group 1)
Higher Rates Are	Worse (H1: λ1 - λ2 < D0, where D0 > 0)
Power	0.9
Alpha	0.025
K2 (Clusters in Group 2)	K1
M (Average of Cluster Sizes)	50
CV (Coef of Variation of Cluster Sizes)	0.2
Incidence Rate Input Type	Incidence Rates ( $\lambda$ 1.0 and $\lambda$ 1.1)
λ1.0 (Non-Inferiority Incidence Rate)	0.6
λ1.1 (Actual Incidence Rate)	0.5
λ2 (Incidence Rate of Control Group)	0.5
ρ (Intracluster Correlation, ICC)	0.002

## Output

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

				Incidence Rate									
		umber of Clusters			Clusters Cluster Size Total -					D'//			
	Num	per of Cluste	ers	Cluster	Size	Total	Treatm	nent		Differe	nce		
					Size	Total Sample	Non-			Non-		Intracluster	
Power	Numl Treatmen K	t Control		Cluster Average M	Size CV			Actual λ1.1	Control λ2	Non-	Actual D1	Correlation	Alpha

**PASS** has also obtained K1 = K2 = 26. Thus, the procedure is validated.