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Chapter 448

Two-Sample T-Tests for Superiority by a Margin Assuming Equal Variance

Introduction

This procedure computes power and sample size for *superiority by a margin* tests in two-sample designs in which the outcome is a continuous normal random variable and the variances of the two groups (populations) are assumed to be equal. Measurements are made on individuals that have been randomly assigned to one of two groups. This is sometimes referred to as a *parallel-groups* design. This design is used in situations such as the comparison of the income level of two regions, the nitrogen content of two lakes, or the effectiveness of two drugs.

The details of sample size calculation for the two-sample design are presented in the Two-Sample T-Tests Assuming Equal Variance chapter and they will not be duplicated here. This chapter only discusses those changes necessary for superiority tests. Sample size formulas for superiority tests of two means are presented in Chow et al. (2018) pages 50-51.

The Statistical Hypotheses

Remember that in the usual *t*-test setting, the null (H0) and alternative (H1) hypotheses for one-sided tests are defined as

$$H_0: \mu_1 - \mu_2 \le \delta_0$$
 versus $H_1: \mu_1 - \mu_2 > \delta_0$

or equivalently

$$H_0: \delta \leq \delta_0$$
 versus $H_1: \delta > \delta_0$.

Rejecting this test implies that the mean difference is larger than the value δ_0 . 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 δ_0 .

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

$$H_0: \mu_1 - \mu_2 \ge \delta_0$$
 versus $H_1: \mu_1 - \mu_2 < \delta_0$

or equivalently

$$H_0: \delta \geq \delta_0$$
 versus $H_1: \delta < \delta_0$.

Superiority by a Margin 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.

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Two-Sample T-Tests for Superiority by a Margin Assuming Equal Variance

$\frac{\textbf{Parameter}}{\mu_1}$	PASS Input/Output Not used	Interpretation Mean of population 1. Population 1 is assumed to consist of those who have received the new treatment.
μ_2	Not used	<i>Mean</i> of population 2. Population 2 is assumed to consist of those who have received the reference treatment.
M_S	SM	Margin of superiority. This is a tolerance value that defines the magnitude of difference that is not of practical importance. This may be thought of as the smallest difference from the reference value that is considered to be of practical significance. This value is assumed to be a positive number.
δ	δ	Actual difference. This is the value of $\mu_1 - \mu_2$, the difference between the means. This is the value at which the power is calculated.

Note that the actual values of μ_1 and μ_2 are not needed. Only their difference is needed for power and sample size calculations.

Superiority by a Margin Tests

A *superiority by a margin test* tests that the treatment mean is better than the reference mean by more than the superiority margin. The actual direction of the hypothesis depends on the response variable being studied.

Case 1: High Values Good

In this case, higher values are better. The hypotheses are arranged so that rejecting the null hypothesis implies that the treatment mean is greater than the reference mean by at least the margin of superiority. The value of δ at which power is calculated must be greater than $|M_S|$. The null and alternative hypotheses with $\delta_0 = |M_S|$ are

$H_0: \mu_1 \le \mu_2 + M_S $	versus	$H_1: \mu_1 > \mu_2 + M_S $
$H_0 \colon \mu_1 - \mu_2 \leq M_S $	versus	$H_1: \mu_1 - \mu_2 > M_S $
$H_0: \delta \leq M_S $	versus	$H_1: \delta > M_S $

Case 2: High Values Bad

In this case, higher values are worse. The hypotheses are arranged so that rejecting the null hypothesis implies that the treatment mean is less than the reference mean by at least the margin of superiority. The value of δ at which power is calculated must be less than $-|M_S|$. The null and alternative hypotheses with $\delta_0 = -|M_S|$ are

$$\begin{split} &H_0 \colon \mu_1 \geq \mu_2 - |M_S| & \text{versus} & H_1 \colon \mu_1 < \mu_2 - |M_S| \\ &H_0 \colon \mu_1 - \mu_2 \geq -|M_S| & \text{versus} & H_1 \colon \mu_1 - \mu_2 < -|M_S| \\ &H_0 \colon \delta \geq -|M_S| & \text{versus} & H_1 \colon \delta < -|M_S| \end{split}$$

Example

A superiority test example will set the stage for the discussion of the terminology that follows. Suppose that a test is to be conducted to determine if a new cancer treatment substantially improves mean bone density. The adjusted mean bone density (AMBD) in the population of interest is 0.002300 gm/cm with a standard deviation of 0.000300 gm/cm. Clinicians decide that if the treatment increases AMBD by more than 5% (0.000115 gm/cm), it provides a significant health benefit.

The hypothesis of interest is whether the mean AMBD in the treated group is more than 0.000115 above that of the reference group. The statistical test will be set up so that if the null hypothesis is rejected, the conclusion will be that the new treatment is superior. The value 0.000115 gm/cm is called the *margin of superiority*.

Two-Sample Equal-Variance T-Test Statistic

Under the null hypothesis, this test assumes that the two groups of data are simple random samples from a single population of normally distributed values that all have the same mean and variance. This assumption implies that the data are continuous, and their distribution is symmetric. The calculation of the test statistic for the case when higher response values are better is as follows.

$$t_{df} = \frac{(\bar{X}_1 - \bar{X}_2) - \delta_0}{s_{\bar{X}_1 - \bar{X}_2}}$$

where

$$\bar{X}_k = \frac{\sum_{i=1}^{n_k} X_{ki}}{n_k},$$

$$s_k = \sqrt{\frac{\sum_{i=1}^{n_k} (X_{ki} - \bar{X}_k)^2}{(n_k - 1)}},$$

$$s_{\bar{X}_1 - \bar{X}_2} = \sqrt{\frac{(n_1 - 1)s_1^2 + (n_2 - 1)s_2^2}{n_1 + n_2 - 2} \left(\frac{1}{n_1} + \frac{1}{n_2}\right)},$$

$$df = n_1 + n_2 - 2$$

and δ_0 is the value of the difference hypothesized by the null hypothesis which depends on the magnitude and sign of M_S .

The null hypothesis is rejected if the computed p-value is less than a specified level (usually 0.05). Otherwise, no conclusion can be reached.

Computing the Power

When $\sigma_1 = \sigma_2 = \sigma$, the power of the equal-variance *t*-test is calculated as follows.

- 1. Find t_{α} such that $1 T_{df}(t_{\alpha}) = \alpha$, where $T_{df}(x)$ is the area to the left of x under a central-t distribution with degrees of freedom, $df = n_1 + n_2 2$.
- 2. Calculate: $\sigma_{\bar{X}} = \sigma \sqrt{\frac{1}{n_1} + \frac{1}{n_2}}$.
- 3. Calculate the noncentrality parameter: $\lambda = \frac{\delta \delta_0}{\sigma_{\bar{x}}}$.
- 4. Calculate: $Power = 1 T'_{df,\lambda}(t_{\alpha})$, where $T'_{df,\lambda}(x)$ is the area to the left of x under a noncentral-t distribution with degrees of freedom, $df = n_1 + n_2 2$, and noncentrality parameter, λ .

When solving for something other than power, **PASS** uses this same power calculation formulation, but performs a search to determine that parameter.

Example 1 - Power Analysis

Suppose that a test is to be conducted to determine if a new cancer treatment improves bone density. The adjusted mean bone density (AMBD) in the population of interest is 0.002300 gm/cm with a standard deviation of 0.000300 gm/cm. Clinicians decide that if the treatment increases AMBD by more than 5% (0.000115 gm/cm), it generates a significant health benefit. They also want to consider what would happen if the margin of superiority is set to 2.5% (0.0000575 gm/cm).

The analysis will be a superiority test at the 0.025 significance level. Power to be calculated assuming that the new treatment has 7.5% improvement on AMBD. Several sample sizes between 10 and 800 will be analyzed. The researchers want to achieve a power of at least 90%. All numbers have been multiplied by 10000 to make the reports and plots easier to read.

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	Power
Higher Means Are	Better (H1: δ > SM)
Alpha	0.025
Group Allocation	Equal (N1 = N2)
Sample Size Per Group	10 50 100 200 300 500 600 800
SM (Superiority Margin)	0.575 1.15
δ (Actual Difference to Detect)	1.725
σ (Standard Deviation)	3

Output

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

Numeric Results

Solve For: Power

Test Type: Two-Sample Equal-Variance T-Test

Difference: $\delta = \mu 1 - \mu 2 = \mu T - \mu R$

Higher Means Are: Better

Hypotheses: $H0: \delta \leq SM \text{ vs. } H1: \delta > SM$

	Sample Size		Superiority Mean Margin Difference		Standard Deviation			
Power	N1	N2	N	SM			Alpha	
0.12553	10	10	20	0.575	1.725	3	0.025	
0.47524	50	50	100	0.575	1.725	3	0.025	
0.76957	100	100	200	0.575	1.725	3	0.025	
0.96885	200	200	400	0.575	1.725	3	0.025	
0.99681	300	300	600	0.575	1.725	3	0.025	
0.99998	500	500	1000	0.575	1.725	3	0.025	
1.00000	600	600	1200	0.575	1.725	3	0.025	
1.00000	800	800	1600	0.575	1.725	3	0.025	
0.06013	10	10	20	1.150	1.725	3	0.025	
0.15601	50	50	100	1.150	1.725	3	0.025	
0.27052	100	100	200	1.150	1.725	3	0.025	
0.48089	200	200	400	1.150	1.725	3	0.025	
0.64940	300	300	600	1.150	1.725	3	0.025	
0.85769	500	500	1000	1.150	1.725	3	0.025	
0.91295	600	600	1200	1.150	1.725	3	0.025	
0.96943	800	800	1600	1.150	1.725	3	0.025	

Power The probability of rejecting a false null hypothesis when the alternative hypothesis is true.

N1 The sample size from group 1.

N2 The sample size from group 2.

N The total sample size from both groups. N = N1 + N2.

SM The magnitude and direction of the margin of superiority. Since higher means are better, this value is positive and is

the minimum distance above µ2 that µ1 must be to conclude that group 1 is superior to group 2.

δ The difference between the group means at which power and sample size calculations are made. $\delta = \mu 1 - \mu 2$.

σ The assumed standard deviation for each of the two groups.

Alpha The probability of rejecting a true null hypothesis.

Summary Statements

A parallel two-group design will be used to test whether the Group 1 (treatment) mean (μ 1) is superior to the Group 2 (reference) mean (μ 2) by a margin, with a superiority margin of 0.575 (H0: $\delta \leq$ 0.575 versus H1: $\delta >$ 0.575, $\delta = \mu$ 1 - μ 2). The comparison will be made using a one-sided, two-sample equal-variance t-test, with a Type I error rate (α) of 0.025. The common standard deviation for both groups is assumed to be 3. To detect a difference in means of 1.725, with sample sizes of 10 in Group 1 and 10 in Group 2, the power is 0.12553.

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Dropout-Inflated Sample Size

	Sample Size			I	Dropout-Inflated Enrollment Sample Size			Expected Number of Dropouts		
Dropout Rate	N1	N2	N	N1'	N2'	N'	D1	D2	D	
20%	10	10	20	13	13	26	3	3	6	
20%	50	50	100	63	63	126	13	13	26	
20%	100	100	200	125	125	250	25	25	50	
20%	200	200	400	250	250	500	50	50	100	
20%	300	300	600	375	375	750	75	75	150	
20%	500	500	1000	625	625	1250	125	125	250	
20%	600	600	1200	750	750	1500	150	150	300	
20%	800	800	1600	1000	1000	2000	200	200	400	
Dropout Rate N1, N2, and N	The evaluable are evaluate	n no respo sample si d out of th	onse data will zes at which p	be collected (i	i.e., will be to outed (as en	treated as "mi tered by the u	ssing"). Abbı ıser). If N1 a	reviated as nd N2 sub	DR. jects	
N1', N2', and N'	formulas N1	f subjects sed on the ' = N1 / (1	assumed dro - DR) and N2	e enrolled in the opout rate. N1' ' = N2 / (1 - DF S.C., Shao, J.,	and N2' ard R), with N1'	e calculated by and N2' alway	y inflating N1 ys rounded u	I and N2 υ ιρ. (See Ju	ising the ulious,	
D1, D2, and D	The expected	number of	duamanta D4	NAL NA D	O NIOL NI	0 ID D4	. DO			

Dropout Summary Statements

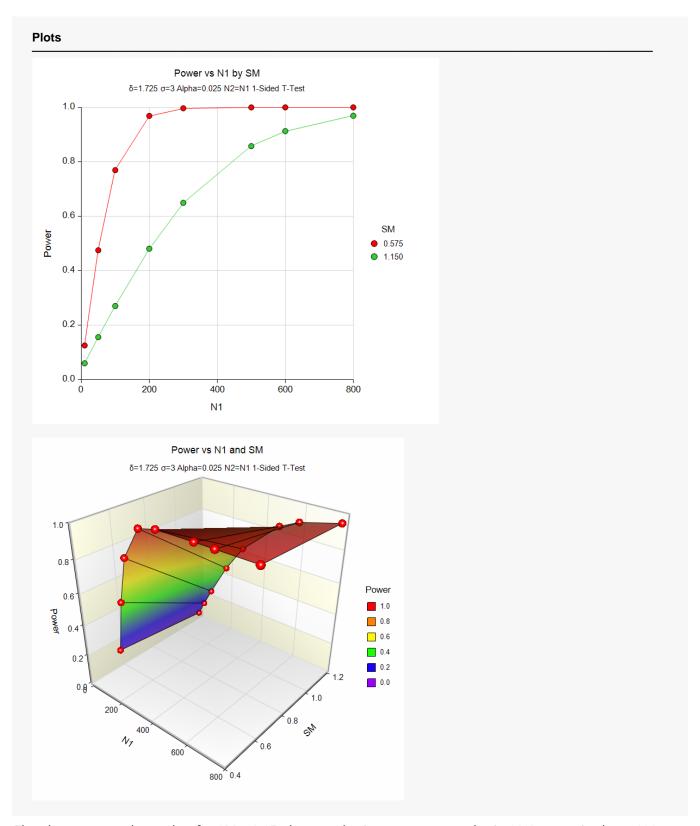
Anticipating a 20% dropout rate, 13 subjects should be enrolled in Group 1, and 13 in Group 2, to obtain final group sample sizes of 10 and 10, respectively.

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.

Julious, Steven A. 2004. 'Tutorial in Biostatistics. Sample sizes for clinical trials with Normal data.' Statistics in Medicine, 23:1921-1986.

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The above report shows that for SM = 1.15, the sample size necessary to obtain 90% power is about 600 per group. However, if SM = 0.575, the required sample size is only about 150 per group.

Example 2 - Finding the Sample Size

Continuing with Example 1, the researchers want to know the exact sample size for each value of SM to achieve 90% power.

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	Sample Size
Higher Means Are	Better (H1: δ > SM)
Power	0.90
Alpha	0.025
Group Allocation	Equal (N1 = N2)
SM (Superiority Margin)	0.575 1.15
δ (Actual Difference to Detect)	1.725
σ (Standard Deviation)	3

Output

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

		_	
Num	eric	RASI	ılte

Solve For: Sample Size

Test Type: Two-Sample Equal-Variance T-Test

Difference: $\delta = \mu 1 - \mu 2 = \mu T - \mu R$

Higher Means Are: Better

Hypotheses: H0: δ ≤ SM vs. H1: δ > SM

Power		Sample Size		Superiority Margin	Mean Difference	Standard Deviation		
Target	Actual	N1	N2	N	SM	δ	σ	Alpha
0.9	0.90004	144	144	288	0.575	1.725	3	0.025
0.9	0.90036	573	573	1146	1.150	1.725	3	0.025

This report shows the exact sample size requirement for each value of SM.

Example 3 – Validation

This procedure uses the same mechanics as the *Two-Sample T-Tests for Non-Inferiority Assuming Equal Variance* procedure. We refer the user to Examples 3 and 4 of Chapter 450 for the validation.