#### Chapter 342

## Multi-Arm Equivalence Tests for the Difference Between Treatment and Control Means Allowing Unequal Variance

## Introduction

This module computes power and sample size for multiple equivalence tests of treatment means versus a control mean when no assumption of equal variances is made. Schuirmann's (1987) two one-sided tests (TOST) approach is used to test equivalence. A modified t-test, known as the Aspin-Welch test, Welch's t-test (Welch, 1937), or the Satterthwaite method, is used for the individual tests. The multiplicity is based on the results in Machin, Campbell, Tan, and Tan (2018).

In this design, there are *k* treatment groups and one control group. A mean is measured in each group. A total of *k* hypothesis tests are anticipated, each comparing a treatment group with the common control group using an Aspin-Welch equivalence test.

A Bonferroni adjustment of the type I error rate may be optionally made because several comparisons are being tested using the same data. Making a multiplicity adjustment is usually recommended, but not always. In fact, Saville (1990) advocates not applying it and Machin, Campbell, Tan, and Tan (2018) include omitting it as a possibility.

#### Background

Whether you want to test several doses of a single treatment or several types of treatments, good research practice requires that each treatment be compared with a control. For example, a popular three-arm design consists of three groups: control, treatment A, and treatment B. Two tests are run: treatment A versus control and treatment B versus the same control. This design avoids having to obtain a second control group for treatment B. Besides the obvious efficiency in subjects, it may be easier to recruit subjects if their chances of receiving the new treatment are better than 50-50.

## **Technical Details**

Suppose you want to compare k treatment groups with means  $\mu_i$  and sample sizes  $N_i$  and one control group with mean  $\mu_c$  and sample size  $N_c$ . The total sample size is  $N = N_1 + N_2 + \cdots + N_k + N_c$ .

Let  $\delta = \mu_i - \mu_C$ .

#### **Equivalence Tests**

Measurements are made on individuals that have been randomly assigned to the groups. This *parallel-groups* design may be analyzed by a set of TOST equivalence tests to show that the means of the treatment and control groups do not differ by more than a small amount, either positive or negative. To conduct an equivalence test, you must set upper and lower equivalence limits for the difference between a treatment mean and the control mean. These limits, which will be called EL and EU, establish an interval of equivalence. When the sample mean difference falls between these limits, the null hypothesis of non-equivalence is rejected and the equivalence of the two group means is concluded.

The statistical hypotheses are written as follows.

 $H_{0i}: \mu_i - \mu_C \leq EL$  or  $\mu_i - \mu_C \geq EU$  vs.  $H_{1i}: EL < \mu_i - \mu_C < EU$ 

or, if we define  $\delta_i = \mu_i - \mu_c$ ,

 $H_{0i}: \delta_i \leq EL$  or  $\delta_i \geq EU$  vs.  $H_{1i}: \delta_i < EU$ 

where EL < 0 and EU > 0. Usually, EL = -EU.

#### Two-Sample Unequal-Variance T-Test (Welch's T-Test) Statistic

Welch (1938) proposed the following test statistic when the two variances are not assumed to be equal.

A suitable Type I error probability is chosen for the test (usually 0.05), the data are collected, and a pair of tstatistics are generated using the formulas

$$t_L = \frac{(\bar{x}_i - \bar{x}_c) - EL}{\sqrt{\frac{s_i^2}{N_i} + \frac{s_c^2}{N_c}}}$$
$$t_U = \frac{(\bar{x}_i - \bar{x}_c) - EU}{\sqrt{\frac{s_i^2}{N_i} + \frac{s_c^2}{N_c}}}$$

where

$$\bar{X}_{i} = \frac{\sum_{j=1}^{N_{i}} X_{ij}}{N_{i}}$$
$$s_{i} = \frac{\sum_{j=1}^{N_{i}} (X_{ij} - \bar{X}_{i})^{2}}{N_{i} - 1}$$

These *t*-statistics follow a *t* distribution approximately, with modified degrees of freedom

$$df = \frac{\left(\frac{S_i^2}{N_i} + \frac{S_c^2}{N_c}\right)^2}{\frac{1}{N_i - 1}\left(\frac{S_i^2}{N_i}\right)^2 + \frac{1}{N_c - 1}\left(\frac{S_c^2}{N_c}\right)^2}$$

#### **Power Calculation**

The power of the unequal-variance equivalence *t*-test procedure is calculated as

$$\Pr(t_L^* \ge t_{1-\alpha,df} \text{ and } t_U^* \le -t_{1-\alpha,df} | \mu_i, \mu_c, \sigma_i, \sigma_c)$$

where  $t_L^*$  and  $t_U^*$  are distributed as the bivariate, noncentral *t* distribution with noncentrality parameters  $\Delta_L$ and  $\Delta_U$  given by

$$\Delta_L = \frac{\delta_i - E_L}{\sqrt{\frac{\sigma_i^2}{N_i} + \frac{\sigma_c^2}{N_c}}}$$
$$\Delta_U = \frac{\delta_i - E_U}{\sqrt{\frac{\sigma_i^2}{N_i} + \frac{\sigma_c^2}{N_c}}}$$

and

$$df = \frac{\left(\sqrt{\frac{\sigma_{i}^{2}}{N_{i}} + \frac{\sigma_{C}^{2}}{N_{C}}}\right)^{4}}{\frac{\sigma_{i}^{4}}{N_{i}^{2}(N_{i} - 1)} + \frac{\sigma_{C}^{4}}{N_{C}^{2}(N_{C} - 1)}}.$$

When solving for sample size, **PASS** uses this same power calculation formulation, but performs a search to determine the sample size.

#### **Multiplicity Adjustment**

Because *k* t-tests between treatment groups and the control group are run when analyzing the results of this study, many statisticians recommend that the Bonferroni adjustment be applied. This adjustment is easy to apply: the value of alpha that is used in the test is found by dividing the original alpha by the number of tests. For example, if the original alpha is set at 0.05 and the number of treatment (not including the control) groups is five, the individual tests will be conducted using an alpha of 0.01.

The main criticism of this procedure is that if there are many tests, the value of alpha becomes very small. To mitigate against this complaint, some statisticians recommend separating the treatment groups into those that are of primary interest and those that are of secondary interest. The Bonferroni adjustment is made by the using the number of primary treatments rather than the total number of treatments.

There are some who advocate ignoring the adjustment entirely in the case of randomized clinical trials. See for example Saville (1990) and the discussion in chapter 14 of Machin, Campbell, Tan, and Tan (2018).

#### Size of the Control Group

Because the control group is used over and over, some advocate increasing the number of subjects in this group. The standard adjustment is to include  $\sqrt{k}$  subjects in the control group for each subject in one of the treatment groups. See Machin, Campbell, Tan, and Tan (2018, pages 231-232). Note that usually, the treatment groups all have the same size.

## Example 1 – Finding the Sample Size

A parallel-group clinical trial is being designed to determine if any or all of three treatment therapies are equivalent to the standard therapy. Suppose the standard therapy has a mean response of 9.3. They want to consider standard deviations of 2.7 in the control group and 3.5 in the three treatment groups. To investigate the sensitivity of the sample sizes to the values of the standard deviations, additional runs with a set of standard deviations 20% higher than those selected and another set 20% lower will be made. These runs are made using the *standard deviation multiplier* option with K = 0.8, 1.0, and 1.2.

The investigators would like a sample size large enough to find statistical significance at the 0.05 level if the actual mean responses of the three treatments are also 9.3. The power of each test is set to 0.80. The equivalence margin is 20% of 9.3 = 1.86, so the equivalence limits are set to -1.86 and 1.86.

Following standard procedure, the control group multiplier will be set to  $\sqrt{k} = \sqrt{3} = 1.732$  since the control group is used for three comparisons in this design.

#### 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.

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Solve For	Sample Size
Power of Each Test	•
Overall Alpha	
Bonferroni Adjustment	
-	Enter Group Allocation Pattern, solve for group sample sizes
EU (Upper Equivalence Limit)	
EL (Lower Equivalence Limit)	Upper Limit
Control Mean	9.3
Control Standard Deviation	2.7
Control Sample Size Allocation	1.732
Set A Number of Groups	
Set A Mean	9.3
Set A Standard Deviation	3.5
Set A Sample Size Allocation	1
Set B Number of Groups	0
Set C Number of Groups	0
Set D Number of Groups	0
More	Unchecked
Add sets of standard deviations with different magnitudes, but identical ratio patterns	Checked
K (σ Multiplier)	09112

#### Output

κ

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

#### **Numeric Reports**

Numeric Results	
Solve For:	Sample Size
Group Allocation:	Enter Group Allocation Pattern, solve for group sample sizes
Test Type:	Unequal-Variance T-Test
Hypotheses:	H0:δ≤EL orδ≥EU vs. H1:EL<δ <eu< td=""></eu<>
Number of Groups:	4
Bonferroni Adjustment:	Standard Bonferroni (Divisor = 3)

			0.					alence nits	Standa	rd Deviation		Alpha
Comparison	Target	ower Actual	Ni	Allocation	Mean µi	Difference δi	Lower EL	Upper EU	Value σi	Multiplier K	Overall	Bonferroni- Adjusted
Control			64	1.732	9.3				2.16	0.8		
vs A1	0.8	0.80311	37	1.000	9.3	0	-1.86	1.86	2.80	0.8	0.05	0.01667
vs A2	0.8	0.80311	37	1.000	9.3	0	-1.86	1.86	2.80	0.8	0.05	0.01667
vs A3	0.8	0.80311	37	1.000	9.3	0	-1.86	1.86	2.80	0.8	0.05	0.01667
Total			175									
Control			99	1.732	9.3				2.70	1.0		
vs A1	0.8	0.80240	57	1.000	9.3	0	-1.86	1.86	3.50	1.0	0.05	0.01667
vs A2	0.8	0.80240	57	1.000	9.3	0	-1.86	1.86	3.50	1.0	0.05	0.01667
vs A3	0.8	0.80240	57	1.000	9.3	0	-1.86	1.86	3.50	1.0	0.05	0.01667
Total			270									
Control			140	1.732	9.3				3.24	1.2		
vs A1	0.8	0.80064	81	1.000	9.3	0	-1.86	1.86	4.20	1.2	0.05	0.01667
vs A2	0.8	0.80064	81	1.000	9.3	0	-1.86	1.86	4.20	1.2	0.05	0.01667
vs A3	0.8	0.80064	81	1.000	9.3	0	-1.86	1.86	4.20	1.2	0.05	0.01667
Total			383									
Comparison		The group	that	is involved i	n the co	mparison be	etween t	he treatr	ment and	l control disp	played on	this report
T						ng the differ			he we at the s	- '- ( ( - '		<b>.</b>
Target Powe	er						cting a fa	alse null	hypothe	sis for this c	ompariso	n. This power
				parison shov		is line only.						
Actual Powe				ally achieve								
Ni				subjects in t ividual group			tal samp	ole size s	shown be	elow the grou	ups is equ	ual to the
Allocation							aroup	The valu	10 00 03	ch row repre	conte the	rolativo
Allocation				pie size allo			i gioup.	The value		chilow lepie		relative
μi							s compu	ited. The	e first row	contains µc	c, the con	trol group
δί		The difference compute		between the	ith trea	tment mean	and the	control	mean (µ	i - µc) at whi	ch the po	wer is
EL		treatme	nt gro		control g					ive mean dif n that the tre		etween each Iroup is
EU		The upper betweer	r equi n each	valence limi	t for the group a	nd the contro				e difference n the conclu		erence the treatment
σί						onses within	this aro	up.				

The multiplier that was applied to form the group standard deviations shown on this line. The probability of rejecting at least one of the comparisons in this experiment when each null hypothesis Overall Alpha is true.

Bonferroni Alpha The adjusted significance level at which each individual comparison is made.

#### **Summary Statements**

A parallel, 4-group design (with one control group and 3 treatment groups) will be used to test whether the mean for each treatment group is equivalent to the control group mean, with equivalence difference bounds of -1.86 and 1.86 (H0:  $\delta \le -1.86$  or  $\delta \ge 1.86$  versus H1: -1.86 <  $\delta < 1.86$ ,  $\delta = \mu i - \mu c$ ). Each of the 3 equivalence comparisons will be made using two one-sided, two-sample, Bonferroni-adjusted, unequal-variance (Welch's) t-tests. The overall (experiment-wise) Type I error rate ( $\alpha$ ) is 0.05. The group standard deviations (beginning with the control group) are assumed to be 2.16, 2.8, 2.8, and 2.8. The control group mean is assumed to be 9.3. To detect the treatment means 9.3, 9.3, and 9.3 with at least 80% power for each test, the control group sample size needed will be 64 and the number of needed subjects for the treatment groups will be 37, 37, and 37 (totaling 175 subjects overall).

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

Group	Dropout Rate	Sample Size Ni	Dropout- Inflated Enrollment Sample Size Ni'	Expected Number of Dropouts Di
1	20%	64	80	16
2	20%	37	47	10
3	20%	37	47	10
4	20%	37	47	10
Total		175	221	46
1	20%	99	124	25
2	20%	57	72	15
3	20%	57	72	15
4	20%	57	72	15
Total		270	340	70
1	20%	140	175	35
2	20%	81	102	21
3	20%	81	102	21
4	20%	81	102	21
Total		383	481	98

Group Lists the group numbers.

 Dropout Rate
 The percentage of subjects (or items) that are expected to be lost at random during the course of the study and for whom no response data will be collected (i.e., will be treated as "missing"). Abbreviated as DR.

 Ni
 The evaluable sample size for each group at which power is computed (as entered by the user). If Ni subjects are evaluated out of the Ni' subjects that are enrolled in the study, the design will achieve the stated power.

 Ni
 The number of subjects that should be enrolled in each group in order to obtain Ni evaluable subjects, based on the assumed dropout rate. Ni' is calculated by inflating Ni using the formula Ni' = Ni / (1 - DR), with Ni' always rounded up. (See Julious, S.A. (2010) pages 52-53, or Chow, S.C., Shao, J., Wang, H., and Lokhnygina, Y. (2018) pages 32-33.)

 Di
 The expected number of dropouts in each group. Di = Ni' - Ni.

#### Dropout Summary Statements

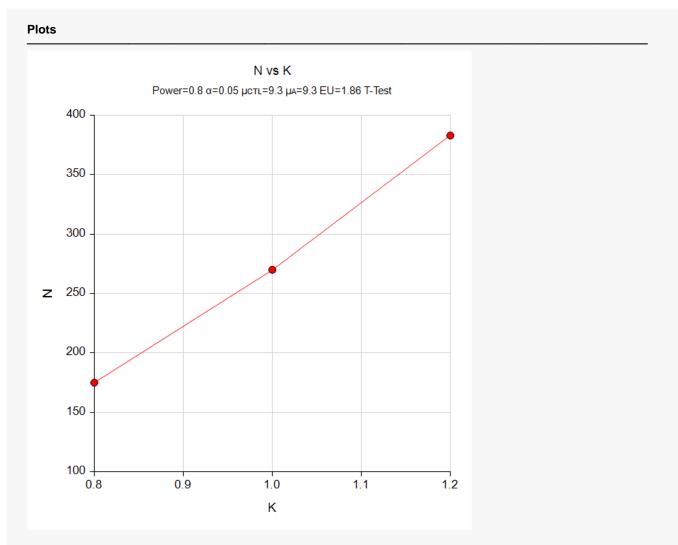
Anticipating a 20% dropout rate, group sizes of 80, 47, 47, and 47 subjects should be enrolled to obtain final group sample sizes of 64, 37, 37, and 37 subjects.

#### References

Blackwelder, W.C. 1998. 'Equivalence Trials.' In Encyclopedia of Biostatistics, John Wiley and Sons. New York. Volume 2, 1367-1372.
Chow, S.C., Shao, J., Wang, H., and Lokhnygina, Y. 2018. Sample Size Calculations in Clinical Research, 3rd
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Machin, D., Campbell, M.J., Tan, S.B, and Tan, S.H. 2018. Sample Sizes for Clinical, Laboratory, and
Epidemiology Studies, 4th Edition. Wiley Blackwell.
Welch, B.L. 1938. 'The significance of the difference between two means when the population variances are
unequal.' Biometrika, 29, 350-362.
Zar, Jerrold H. 1984. Biostatistical Analysis (Second Edition). Prentice-Hall. Englewood Cliffs, New Jersey.

This report shows the numeric results of this power study. Notice that the results are shown in blocks of three rows at a time. Each block represents a single design.

#### **Plots Section**



This plot gives a visual presentation of the results in the Numeric Report. We can quickly see the impact on the sample size of changing the standard deviation magnitude.

# Example 2 – Validation using a Previously Validated Procedure

We could not find a validation result in the statistical literature, so we will use a previously validated **PASS** procedure (**Two-Sample T-Tests for Equivalence Allowing Unequal Variance**) to produce the results for the following example.

A parallel-group clinical trial is being designed to determine if any or all of three treatment therapies are equivalent to the standard therapy. Suppose the standard therapy has a mean response of 9.3. They want to consider standard deviations of 2.7 in the control group and 3.5 in the three treatment groups. The investigators would like a sample size large enough to find statistical significance at the 0.05 level if the actual mean responses of the three treatments are also 9.3. The power of each test is set to 0.80. The equivalence margin is 20% of 9.3 = 1.86, so the equivalence limits are set to -1.86 and 1.86.

The sample sizes of all groups will be equal.

The Two-Sample T-Tests for Equivalence Allowing Unequal Variance procedure is set up as follows.

Design Tab	
Solve For	Sample Size
Power	0.8
Alpha	<b>0.01667</b> (which is Alpha / k)
Group Allocation	Equal (N1 = N2)
EU (Upper Equivalence Limit)	1.86
EL (Lower Equivalence Limit)	Upper Limit
δ (Actual Difference)	0
σ1 (Standard Deviation of Group 1)	3.5
σ2 (Standard Deviation of Group 2)	2.7

This set of options generates the following report.

#### **Numeric Results**

Solve For:	Sample Size
Difference:	δ = μ1 - μ2 = μΤ - μR
Hypotheses:	H0: $\delta \leq EL \text{ or } \delta \geq EU \text{ vs. } H1: EL < \delta < EU$
Test Type:	Two One-Sided Unequal-Variance T-Tests

Pov	ver	Sa	ample S	ize	Equivaler	nce Limits	Actual		ndard ation	
Target	Actual	 N1	N2	N	Lower EL	Upper EU	Difference δ	<u>σ1</u>	σ2	Alpha
0.8	0.81252	68	68	136	-1.86	1.86	0	3.5	2.7	0.01667

In order to maintain a power of 80% for all three groups, it is apparent that the groups will all need to have a sample size of 68 per group. This table contains the validation values. We will now run these values through the current procedure and compare the results with these values.

#### 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
Power of Each Test	0.80
Overall Alpha	0.05
Bonferroni Adjustment	Standard Bonferroni
Group Allocation	Equal (Nc = N1 = N2 =)
EU (Upper Equivalence Limit)	1.86
EL (Lower Equivalence Limit)	Upper Limit
Control Mean	9.3
Control Standard Deviation	2.7
Set A Number of Groups	3
Set A Mean	9.3
Set A Standard Deviation	3.5
Set B Number of Groups	0
Set C Number of Groups	0
Set D Number of Groups	0
More	Unchecked
Add sets of standard deviations with different magnitudes, but identical ratio patterns	Unchecked

#### Output

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

Solve For: Group Allocatio Test Type: Hypotheses: Number of Gro Bonferroni Adju	on: E L H ups: 4	Sample Size Equal (Nc = N Jnequal-Varia H0: $\delta \le EL$ or H Standard Bon	ance T-Test δ≥EU vs.		δ <eu< th=""><th></th><th></th><th></th><th></th><th></th></eu<>						
			0 - maile				alence nits	Oton doub		Alpha	
Comparison	P	ower	Sample Size Ni	Size					Standard		
Comparison	Target	Actual		Mean µi	Difference δi	Lower EL	Upper EU	Deviation σi	Overall	Bonferroni- Adjusted	
Comparison Control	Target	Actual							Overall		
· · · · · · · · · · · · · · · · · · ·	Target	<b>Actual</b> 0.81249	Ni	μi				σi	Overall		
Control			<b>Ni</b> 68	<b>µі</b> 9.3	δί	EL	EU	<b>σi</b> 2.7		Adjusted	

As you can see, the sample sizes are all 68. This matches the sample size found in the validation run above. The procedure is validated.