

Selecting a Valid Sample Size for Longitudinal and Multilevel Studies in Oral Behavioral Health

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Conflict of Interest

We have no conflicts of interest to declare.

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Learning Objectives

- Learn a conceptual framework for conducting a power analysis.
- Understand how to interact with our free, web-based power and sample size software.
- Write a sample size analysis.

The Sample Size Game

Object of the game: Calculate sample size

- Speakers present information.
- Audience discusses the information in small groups using worksheets.
- Next speaker shows how the information can be used to calculate sample size.

Agenda

How Do we Choose Sample Size and Power for Complex Oral Health Designs?

10:50 - 11:00

Dr. Henrietta Logan

Discussion: Hypothesis, Outcomes, and

11:00 - 11:10

Predictors

Choosing a Hypothesis, Outcomes, and Predictors with Our Free, Web-based

11:10 - 11:20

Software

Dr. Aarti Munjal

Discussion: Mean, Variance, and Correlation 11:20 – 11:30

Agenda

Choosing Means, Variances, and Correlations
with Our Free, Web-based Software
Brandy M. Ringham
Discussion: Sample Size Calculation

11:30 – 11:40

11:40 – 11:50

Wrapping it Up: Writing the Grant 11:50 – 12:00

Deborah H. Glueck

Summary

Discussion: Question and Answer 12:00 – 12:15

How Do we Choose Sample Size and Power for Complex Oral Health Designs?

Dr. Henrietta Logan University of Florida

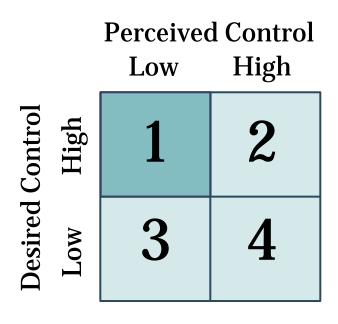


Previous Study on Sensory Focus to Alleviate Pain

- Participants categorized into four coping styles
- Randomized to one of two intervention arms:

sensory focus standard of care

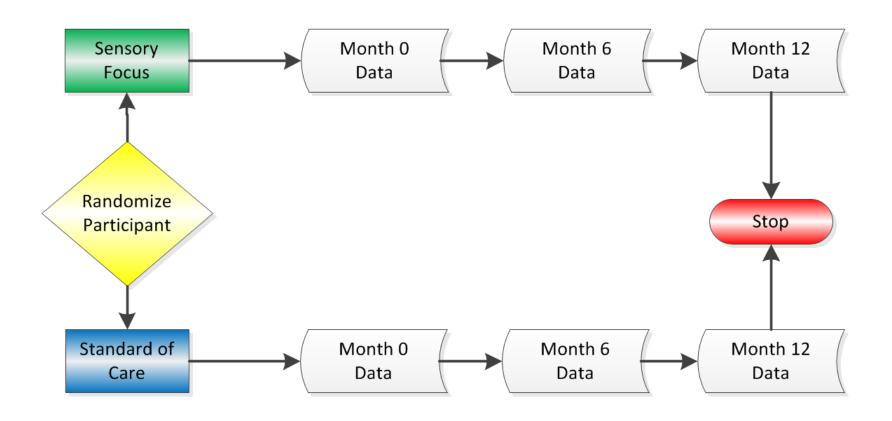
 Measured experienced pain after root canal



(Logan, Baron, Kohout, 1995)

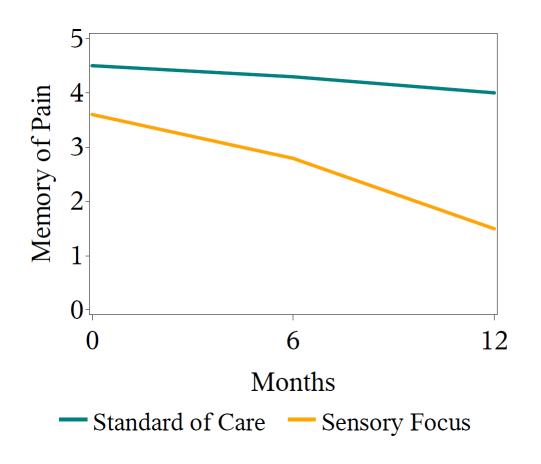


Memory of Pain Trial Study Design





Memory of Pain Trial Research Question





Memory of Pain Trial Study Population

- Recruit participants who have a high desire/low felt coping style
- 30 patients / week
- 40% consent rate for previous studies



Ethics of Sample Size Calculations

- If the sample size is too small, the study may be inconclusive study and waste resources
- If the sample size is too large, then the study may expose too many participants to possible harms due to research



How do we calculate an accurate sample size?



Type I error rate:

Desired power:

Loss to follow-up:



• Type I error rate: 0.01

Desired power:

Loss to follow-up:



• Type I error rate:

0.01

Desired power:

0.90

Loss to follow-up:



Type I error rate:

Desired power:

Loss to follow-up:

0.01

0.90

25%



- <u>Hypothesis</u>: the question that the research study is designed to answer
- Outcome: a measureable trait used to answer the research question
- Predictors: factors that may affect the outcome of the study

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11:20 - 11:30

Choosing a Hypothesis, Outcomes, and Predictors with Our Free, Web-based Software

Dr. Aarti Munjal University of Colorado Denver

1. Solving for:

2. Desired power:

3. Type I error rate:

1. Solving for: Sample size (B)

2. Desired power:

3. Type I error rate:

1. Solving for: Sample size (B)

2. Desired power: 0.90 (B)

3. Type I error rate:

1. Solving for: Sample size (B)

2. Desired power: 0.90 (B)

3. Type I error rate: 0.01 (D)



4. Outcome:

5. Predictor:

6. Hypothesis:



4. Outcome: memory of pain (C)

5. Predictor:

6. Hypothesis:



4. Outcome: memory of pain (C)

5. Predictor: intervention group (D)

6. Hypothesis:



4. Outcome: memory of pain (C)

5. Predictor: intervention group (D)

6. Hypothesis: time by intervention interaction (A)



GLIMMPSE

GLIMMPSE is a user-friendly online tool for calculating power and sample size for multilevel and longitudinal studies.

http://glimmpse.samplesizeshop.org/



Salient Software Features

- Free
- Requires no programming expertise
- Allows saving study designs for later use
- Also available on smartphones



Create a Study Design

Start Your Study Design

Welcome to GLIMMPSE. The GLIMMPSE software calculates power and sample size for study designs with normally distributed outcomes. Select one of the options below to begin your power or sample size calculation.

Guided Study Design

Build common study designs including ANOVA, ANCOVA, and regression with guidance from the study design wizard. This mode is designed for applied researchers including physicians, nurses, and other investigators.

Select

Matrix Study Design

Directly enter the matrices for the general linear model. This mode is designed for users with advanced statistical training.

Select

Upload a Study Design

If you have previously saved a study design from GLIMMPSE, you may upload it here. Click browse to select your study design file.

Choose File No f...sen



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Select Guided Mode

GLIMMPSE Solving For

Calculate Start Solving For Desired Power Type I Error Sampling Unit Responses Hypothesis Means Variability Options

Would you like to solve for power or sample size?

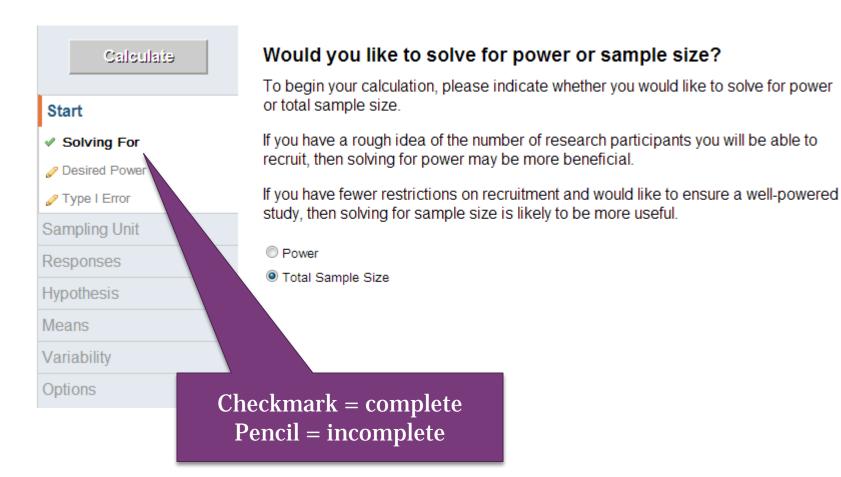
To begin your calculation, please indicate whether you would like to solve for power or total sample size.

If you have a rough idea of the number of research participants you will be able to recruit, then solving for power may be more beneficial.

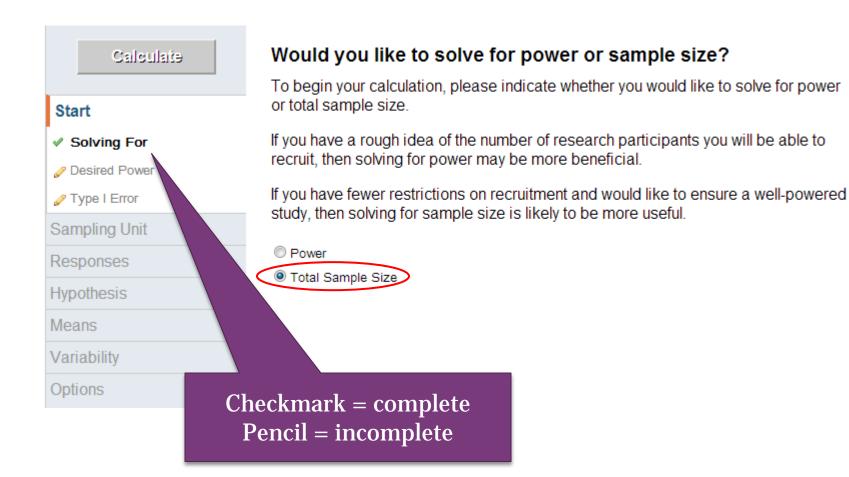
If you have fewer restrictions on recruitment and would like to ensure a well-powered study, then solving for sample size is likely to be more useful.

- Power
- Total Sample Size

GLIMMPSE Solving For



GLIMMPSE Solving For



GLIMMPSE Desired Power

Power Values

Enter the desired power values in the list box below. Power values are numbers between 0 and 1. Higher values correspond to a greater likelihood of rejecting the null hypothesis. Common values are 0.8 or 0.9, although 0.9 or higher is usually preferred.

Type each value into the list box and click "Add". To remove an item, highlight the value and click the "Delete" button.

Power Values:	Add Delete	
0.9		^
		÷

GLIMMPSE Desired Power

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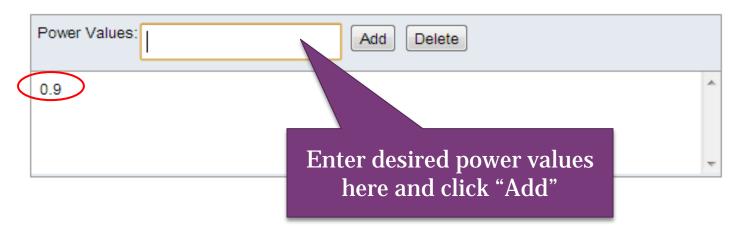


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GLIMMPSE Type I Error Rate

Type I Error

A Type I error occurs when a scientist declares a difference when none is actually present. The Type I error rate is the probability of a Type I error occurring, and is often referred to as α. Type I error rates range from 0 to 1. The most commonly used values are 0.01, 0.05, and 0.1.

Enter each Type I error value into the text box and click "Add". You may enter up to 5 values. To remove a value, select the value in the list box and click the "Delete" button.

Type I Error Values:	Add Delete
0.01	^
	·



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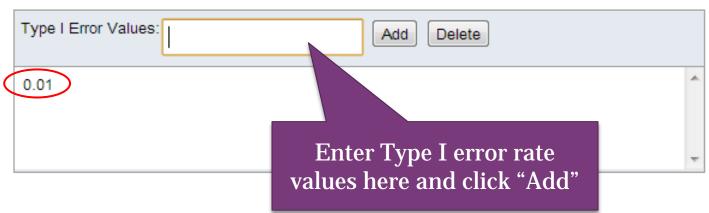


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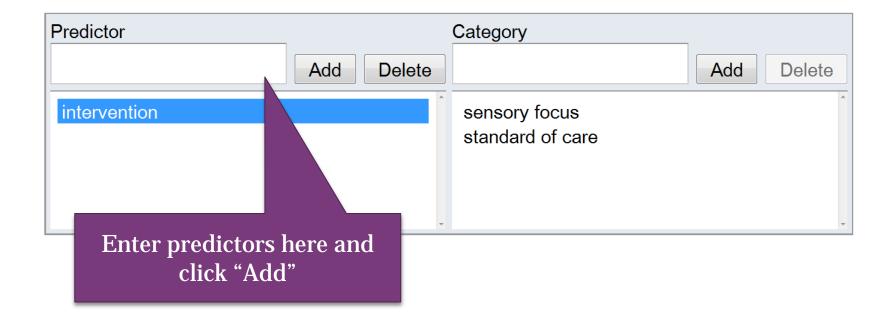


GLIMMPSE Predictors

Predictor	7		Category	
	Add	Delete		Add Delete
intervention			sensory focus standard of care	

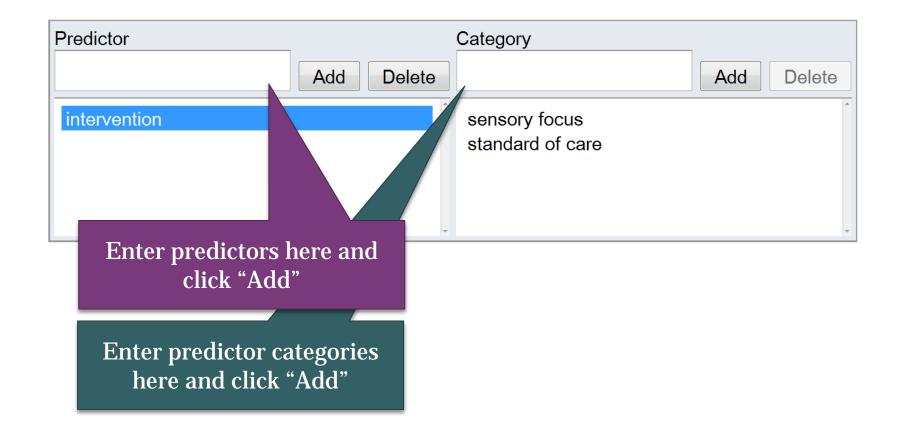


GLIMMPSE Predictors





GLIMMPSE Predictors





GLIMMPSE Outcome

Response Variables

Enter the response variables in the table below. For example, in a study investigating cholesterol-lowering medication, the response variable could be HDL, LDL, and total cholesterol.

Note that repeated measurement information will be addressed on the next screen.





GLIMMPSE Outcome

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GLIMMPSE Outcome

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GLIMMPSE Repeated Measures

Remove Repeated Measures

Units	time
Туре	Numeric
Number of Measurements	3
Spacing	1 2 3
Reset to Equal Spacing	
dd Level Remove Level	



GLIMMPSE Repeated Measures

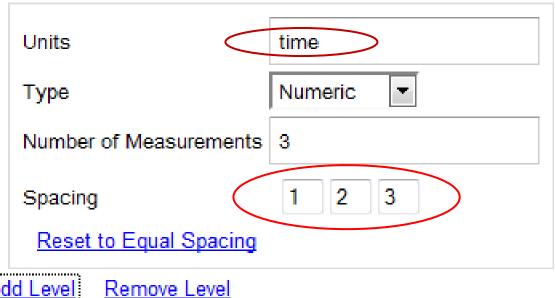
Remove Repeated Measures

Units	time
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GLIMMPSE Repeated Measures

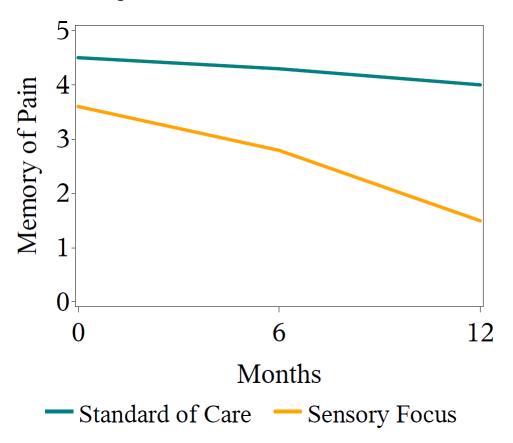
Remove Repeated Measures



Add Level



time by intervention interaction





Grand mean
 Main Effect
 Trend
 Interaction

Select two or more predictors to include in the interaction hypothesis. To test for a trend in a given factor, click the Edit Trend link and select an appropriate trend.

Between Participant Factors

Within Participant Factors

■ time Edit trend : None



Grand mean Main Effect Trend Interaction

Select two or more predictors to include in the interaction hypothesis. To test for a trend in a given factor, click the Edit Trend link and select an appropriate trend.

Between Participant Factors

■ intervention Edit trend : None

Within Participant Factors

■ time Edit trend : None



Select two or more predictors to include in the interaction hypothesis. To test for a trend in a given factor, click the Edit Trend link and select an appropriate trend.

Between Participant Factors

intervention Edit trend: None

Within Participant Factors

time Edit trend: None



Select two or more predictors to include in the interaction hypothesis. To test for a trend in a given factor, click the Edit Trend link and select an appropriate trend.

Between Participant Factors

intervention Edit trend: None

Within Participant Factors

time Edit trend: None



Where Can I Find Means, Variances, and Correlations?

- Pilot study
- Similar published research
- Unpublished internal studies
- Clinical experience



- <u>Mean</u>: a measure of the size of the intervention effect
- <u>Variance</u>: a measure of the variability of the outcome
- <u>Correlation</u>: a measure of the association between the repeated measures

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Discussion: Mean, Variance, and Correlation

11:20 - 11:30

Agenda

Choosing Means, Variances, and Correlations with Our Free, Web-based Software Brandy M. Ringham	11:30 – 11:40
Discussion: Sample Size Calculation Summary	11:40 – 11:50
Wrapping it Up: Writing the Grant Deborah H. Glueck	11:50 – 12:00
Discussion: Question and Answer	12:00 – 12:15

Choosing Means, Variances, and Correlations with Our Free, Web-based Software

Brandy Ringham University of Colorado Denver

Correlation Between Outcomes Over Time

Correlation Between Outcomes Over Time

Correlation Between Outcomes Over Time

Correlation at 6 months apart

(A)

Correlation at 12 months apart

(B)



Correlation at 6 months apart



Correlation at 12 months apart

(B) **0.4**



<u>Correlation Between Outcomes Over Time</u>



<u>Correlation Between Outcomes Over Time</u>

Correlation at 6 months apart



Correlation at 12 months apart

(B) **0.4**

Correlation at 6 months apart

(A) **0.5**

Correlation at 12 months apart

(B) **0.4**

Standard Deviation of the Outcome

Logan, Baron, and Kohout (1995) examined whether sensory focus therapy during a root canal procedure could reduce a patient's experienced pain. The investigators assessed experienced pain on a 5 point scale both immediately and at one week following the procedure. The standard deviation of the measurements was 0.98.

Standard Deviation of the Outcome

Logan, Baron, and Kohout (1995) examined whether sensory focus therapy during a root canal procedure could reduce a patient's experienced pain. The investigators assessed experienced pain on a 5 point scale both immediately and at one week following the procedure. The standard deviation of the measurements was 0.98.

Standard deviation of memory of pain



Standard deviation of memory of pain

(C) 0.98

Intervention	Baseline	6 Months	12 Months
Sensory Focus (SF)	3.6	2.8	0.9
Standard of Care (SOC)	4.5	4.3	3.0

Intervention Difference (SF - SOC)

(D)

(E)

(F)

Net Difference Over Time (12 Months - Baseline)

Intervention	Baseline	6 Months	12 Months
Sensory Focus (SF)	3.6	2.8	0.9
Standard of Care (SOC)	4.5	4.3	3.0

Intervention Difference (SF - SOC)

(D) -0.9

(E)

(F)

Net Difference Over Time (12 Months - Baseline)

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(D) -0.9

(E) -1.5

(F)

Net Difference Over Time (12 Months - Baseline)

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Intervention Difference (SF - SOC)

(D) -0.9

(E) -1.5

(F) -2.1

Net Difference Over Time (12 Months - Baseline)

Intervention	Baseline	6 Months	12 Months
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Standard of Care (SOC)	4.5	4.3	3.0

Intervention Difference (SF - SOC)

(D) -0.9

(E) -1.5

(F) -2.1

Net Difference Over Time (12 Months - Baseline)

(G) -1.2

GLIMMPSE Means Specifying a Mean Difference

treatment	memory of pain
sensory focus	-1.2
standard of care	0

Select the time (location, etc.) from the list(s) below. This will allow you to edit the means at the selected time (location, etc.). time 3

GLIMMPSE Means Specifying a Mean Difference

intervention	memory of pain
sensory focus	-1.2
standard of care	0

Select the time (location, etc.) from the list(s) below. This will allow you to edit the means at the selected time (location, etc.).

time 3

Choose a timepoint

GLIMMPSE Means Specifying a Mean Difference

intervention	memory of pain		
sensory focus	-1.2		
standard of care	0		

Select the time (location, etc.) from the list(s) bere. This will allow you to edit the means at the selected time (location, etc.). time 3

Choose a timepoint

Enter the expected net mean difference

GLIMMPSE Variability Entering Standard Deviation of the Outcome

time

Responses

Enter the standard deviation you expect to observe for each response. Note that GLIMMPSE currently assumes that the standard deviation is constant across repeated measurements.

memory of pain 0.98

GLIMMPSE Variability **Entering Standard Deviation of the Outcome**

time

Responses

Enter the standard deviation you expect to observe for each response. Note that GLIMMPSE currently assumes that the standard deviation is constant across repeated measurements.

memory of pain 0.98

Enter the standard deviation of the outcome variable



GLIMMPSE Variability **Specifying Correlations**

Responses time

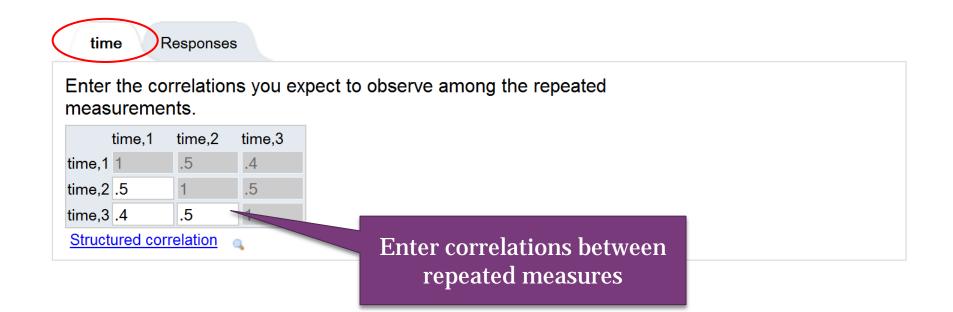
Enter the standard deviation you expect to observe for each response. Note that GLIMMPSE currently assumes that the standard deviation is constant across repeated measurements.

memory of pain 0.98

Enter the standard deviation of the outcome variable



GLIMMPSE Variability Specifying Correlations





GLIMMPSE Hypothesis Test

Statistical Tests

Select the statistical tests to include in your calculations. For study designs with a single outcome, power is the same regardless of the test selected.

Note that only the Hotelling-Lawley Trace and the Univariate Approach to Repeated Measures are supported for designs which include a baseline covariate.

Click here to learn more about selecting an appropriate test.

- Hotelling-Lawley Trace
- Pillai-Bartlett Trace
- Wilks Likelihood Ratio
- Univariate Approach to Repeated Measures with Box Correction
- Univariate Approach to Repeated Measures with Geisser-Greenhouse Correction
- Univariate Approach to Repeated Measures with Huynh-Feldt Correction
- Univariate Approach to Repeated Measures, uncorrected



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- Univariate Approach to Repeated Measures, uncorrected



GLIMMPSE Calculate Button





Power Results

Power	Total Sample Size	Target Power	Test	Type I Error Rate	Means Scale Factor	Variability Scale Fac
0.901	44	0.900	HLT	0.01	1	1
0.925	26	0.900	HLT	0.01	1	0.5
0.905	84	0.900	HLT	0.01	1	2
4				111		(b)

Save to CSV View Matrices



Power Results

Power	Total Sample Size	Target Power	Test	Type I Error Rate	Means Scale Factor	Variability Scale Fac
0.901	44	0.900	HLT	0.01	1	1
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0.905	84	0.900	HLT	0.01	1	2
4				111		>

Save to CSV View Manises

Total sample size to achieve at least 90% power



Power Results

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0.901	44	0.900	HLT	0.01	1	1
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0.905	84	0.900	HLT	0.01	1	2
4				111		

Save to CSV View Manises

Scale the standard deviation to ½ and 2 times to see how it affects sample size

Total sample size to achieve at least 90% power



Power Results

Power	Total Sample Size	Target Power	Test	Type I Error Rate	Means Scale Factor	Variability Scale Fac
0.901	44	0.900	HLT	0.01	1	1
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4				111		D

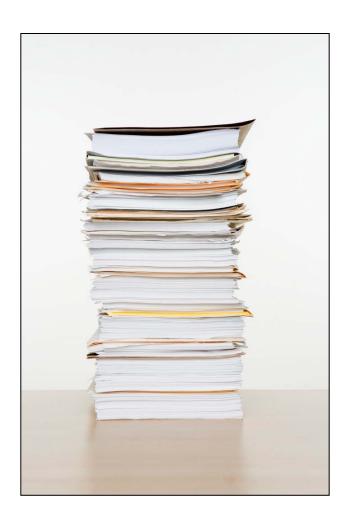
Save to CSV View Manises

Scale the standard deviation to ½ and 2 times to see how it affects sample size

Total sample size to achieve at least 90% power



Funding the Planned Study





Worksheet 3

Sample Size Calculation Summary

- Summarize the sample size calculation
- Include the following information:
 - Type I error rate
 - Desired power
 - Hypothesis
 - Hypothesis test used
 - Analysis method
 - Means, variances, correlation with justification
 - Calculated sample size

Agenda

Choosing Means, Variances, and Correlations with Our Free, Web-based Software

11:30 - 11:40

Brandy M. Ringham

Discussion: Sample Size Calculation

11:40 - 11:50

Summary

Wrapping it Up: Writing the Grant

11:50 - 12:00

Deborah H. Glueck

Discussion: Question and Answer

12:00 - 12:15

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Choosing Means, Variances, and Correlations with Our Free, Web-based Software

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11:40 - 11:50

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11:50 - 12:00

Deborah H. Glueck

Discussion: Question and Answer

12:00 - 12:15

Wrapping it Up: Writing the Grant

Dr. Deborah Glueck University of Colorado Denver

Outline Writing the Grant

- Aligning power analysis with data analysis
- Justifying the power analysis
- Accounting for uncertainty
- Handling missing data
- Demonstrating enrollment feasibility
- Planning for multiple aims

Worksheet 3Sample Size Calculation Summary

We plan a repeated measures ANOVA using the Hotelling-Lawley Trace to test for a time by intervention interaction.

Worksheet 3Sample Size Calculation Summary

We plan a <u>repeated measures ANOVA</u> using the <u>Hotelling-Lawley Trace</u> to <u>test for a time</u> by intervention interaction.



Aligning Power Analysis with Data Analysis

- Type I error rate
 - $\alpha = 0.01$
- Hypothesis test
 - Wrong: power = intervention data analysis = time x intervention
 - Right: power = time x intervention
 data analysis = time x intervention

Worksheet 3Sample Size Calculation Summary

Based on previous studies, we predict memory of pain measures will have a standard deviation of 0.98 and the correlation between baseline and 6 months will be 0.5. Based on clinical experience, we believe the correlation will decrease slowly over time, for a correlation of 0.4 between pain recall measures at baseline and 12 months.

Worksheet 3Sample Size Calculation Summary

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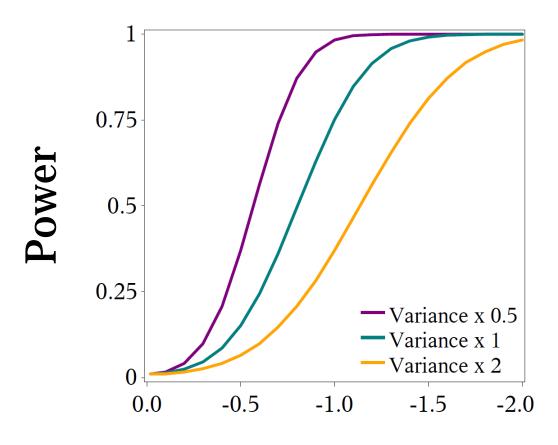
Justifying the Power Analysis

- Give all the values needed to recreate the power analysis
- Provide appropriate citation

For a desired power of 0.90 and a Type I error rate of 0.01, we estimated that we would need 44 participants to detect a clinically meaningful mean difference of 1.2.

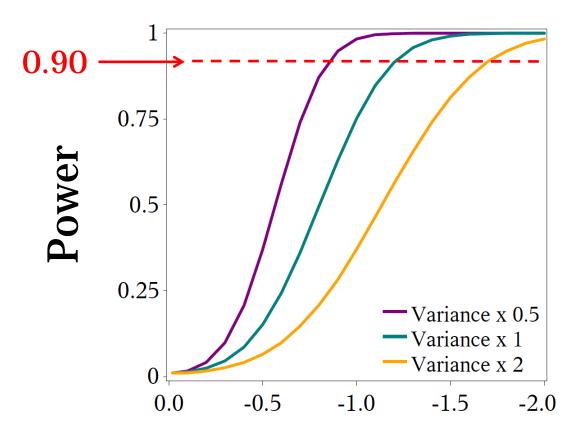
For a <u>desired power of 0.90</u> and a <u>Type I</u> <u>error rate of 0.01</u>, we estimated that we would need <u>44 participants</u> to detect a clinically meaningful <u>mean difference of 1.2</u>.

Accounting for Uncertainty



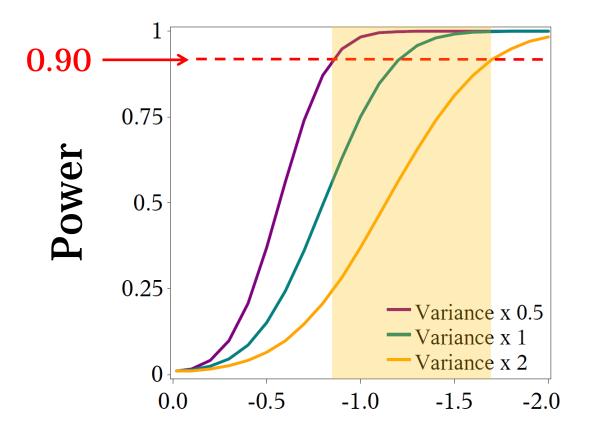
Mean Difference

Accounting for Uncertainty



Mean Difference

Accounting for Uncertainty



Mean Difference



We plan a <u>repeated measures ANOVA</u> using the <u>Hotelling-</u> <u>Lawley Trace</u> to test for a <u>time by intervention interaction</u>. Based on previous studies, we predict measures of pain recall will have a standard deviation of 0.98. correlation in pain recall between baseline and 6 months will be 0.5. Based on clinical experience, we predict that the correlation will decrease slowly over time. Thus, we anticipate a correlation of 0.4 between pain recall measures at baseline and 12 months. For a desired power of 0.90 and a Type I error rate of 0.01, we need to enroll 44 participants to detect a clinically meaningful mean difference of 1.2.



Handling Missing Data

- 25% loss to follow-up
- Account for missing data by increasing the sample size

$$44 / 0.75 = 59$$



Handling Missing Data

- 25% loss to follow-up
- Account for missing data by increasing the sample size

 $44 / 0.75 \approx 60$

Over 12 months, we expect 25% loss to follow up. To account for attrition, we will increase the sample size to 60 participants, or 30 participants per intervention arm.

Over 12 months, we expect <u>25% loss to follow up</u>. To account for attrition, we will increase the sample size to <u>60 participants</u>, or 30 participants per intervention arm.



Demonstrating Enrollment Feasibility

- Is the target population sufficiently large?
- Can recruitment be completed in the proposed time period?



Planned Sample Size vs. Available Sample Size

- 30 patients per week with a high desire / low felt coping style
- 40% consent rate

Sample size needed 60

Sample size available



Planned Sample Size vs. Available Sample Size

- 30 patients per week with a high desire / low felt coping style
- 40% consent rate

3 week enrollment period

Sample size needed 60

Sample size available 36



Planned Sample Size vs. Available Sample Size

- 30 patients per week with a high desire / low felt coping style
- 40% consent rate

5 week enrollment period

Sample size needed 60

Sample size available 60

The clinic treats 30 patients per week with the high desire/low felt coping style. Based on recruitment experience for previous studies, we expect a 40% consent rate. At an effective enrollment of 12 participants per week, we will reach the enrollment goal of 60 participants in 5 weeks time.

The clinic treats <u>30 patients per week</u> with the high desire/low felt coping style. Based on recruitment experience for previous studies, we expect a <u>40% consent rate</u>. At an effective enrollment of 12 participants per week, we will reach the enrollment goal of <u>60 participants in 5 weeks time</u>.

Planning for Multiple Aims

- Aims typically represent different hypotheses
- Maximum of the sample sizes calculated for each aim

Questions?



Question & Answer

- How do I find GLIMMPSE?
- How can I put it on my smartphone?
- Can you review a point from the example power analysis?

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