

## SAMPLE SIZE DETERMINATION IN CLINICAL TRIALS WITH MULTIPLE ENDPOINTS%0A

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Bushi Wang and Naitee Ting, Sample size determination with familywise control of both type I and type II errors in clinical trials, *Journal of Biopharmaceutical Statistics*, 10.1080/10543406.2016.1148706, 26, 5, (951-965), (2016).

POWER AND SAMPLE SIZE DETERMINATION IN CLINICAL TRIALS ...

POWER AND SAMPLE SIZE DETERMINATION IN CLINICAL TRIALS WITH MULTIPLE PRIMARY CONTINUOUS CORRELATED ENDPOINTS Pierre Lafaye de Micheaux<sup>1</sup>, Benoit Liqueur<sup>2,3,4</sup>, S bastien Marque<sup>5</sup>, and J r m ie Riou<sup>2,3,4</sup> <sup>1</sup>Department of Mathematics and Statistics, Universit de Montr al, Quebec, Canada <sup>2</sup>University of Bordeaux, ISPED, INSERM, Bordeaux, France <sup>3</sup>INSERM, ISPED, Bordeaux, France <sup>4</sup>The University of

Sample Size Estimation in Clinical Trial

Every clinical trial should be planned. This plan should include the objective of trial, primary and secondary endpoint, method of collecting data, sample to be included, sample size with scientific justification, method of handling data, statistical methods and assumptions.

SAMPLE SIZE DETERMINATION IN SUPERIORITY CLINICAL TRIALS ...

to multiple endpoints should be handled in the design and analysis of clinical trials. Specially, at the design stage, in addition to incorporating the prede ned strategy for statistical analysis into a trial protocol, statisticians should

consider providing

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Sample Size Determination for Clinical Trials Paivand

Jalalian Advisor: Professor Kelly McConville May 17,

2014 Abstract An important component of clinical trials is determining the smallest sample size