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Developing and Validating Test Items Thomas M. Haladyna,Michael C. Rodriguez,2013-07-18 Since test items are the building blocks of any test learning how to develop and validate test items has always been critical to the teaching learning process As they grow in importance and use testing programs increasingly supplement the use of selected response multiple choice items with constructed response formats This trend is expected to continue As a result a new item writing book is needed one that provides comprehensive coverage of both types of items and of the validity theory underlying them This book is an outgrowth of the author s previous

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Innovative Strategies, Statistical Solutions and Simulations for Modern Clinical Trials Mark Chang, John Balser, Jim Roach, Robin Bliss, 2019-03-20 This is truly an outstanding book It brings together all of the latest research in clinical trials methodology and how it can be applied to drug development Chang et al provide applications to industry supported trials This will allow statisticians in the industry community to take these methods seriously Jay Herson Johns Hopkins University The pharmaceutical industry s approach to drug discovery and development has rapidly transformed in the last decade from the more traditional Research and Development R D approach to a more innovative approach in which strategies are employed to compress and optimize the clinical development plan and associated timelines However these strategies are generally being considered on an individual trial basis and not as part of a fully integrated overall development program Such optimization at the trial level is somewhat near sighted and does not ensure cost time or development efficiency of the overall program This book seeks to address this imbalance by establishing a statistical framework for overall global clinical development optimization and providing tactics and techniques to support such optimization including clinical trial simulations Provides a statistical framework for achieve global optimization in each phase of the drug development process Describes specific techniques to support optimization including adaptive designs precision medicine survival endpoints dose finding and multiple testing Gives practical approaches to handling missing data in clinical trials using SAS Looks at key controversial issues from both a clinical and statistical perspective Presents a generous number of case studies from multiple therapeutic areas that help motivate and illustrate the statistical methods introduced in the book Puts great emphasis on software implementation of the statistical methods with multiple examples of software code both SAS and R It is important for statisticians to possess a deep knowledge of the drug development process beyond statistical considerations For these reasons this book incorporates both statistical and clinical medical perspectives

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