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- Event Location: _____

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Richard Bailey



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Letavec, 2006-09-15 The effective management of project portfolios has become a growing challenge for industry government and other institutions This text provides expert advice information concepts and working models for all organisations wishing to improve standardise and optimise their project management Clinical Trials Tom Brody, 2011-11-11 Clinical Trials Study Design Endpoints and Biomarkers Drug Safety and FDA and ICH Guidelines is a practical guidebook for those engaged in clinical trial design This book details the organizations and content of clinical trials including trial design safety endpoints subgroups HRQoL consent forms and package inserts It provides extensive information on both US and international regulatory guidelines and features concrete examples of study design from the medical literature This book is intended to orient those new to clinical trial design and provide them with a better understanding of how to conduct clinical trials It will also act as a guide for the more experienced by detailing endpoint selection and illustrating how to avoid unnecessary pitfalls This book is a straightforward and valuable reference for all those involved in clinical trial design Provides extensive coverage of the study schema and related features of study design Offers a hands on reference that contains an overview of the process but more importantly details a step by step account of clinical trial design Features examples from the medical literature to highlight how investigators choose the most suitable endpoint s for clinical trial and includes graphs from real clinical trials to help explain each concept in study design Integrates clinical trial design pharmacology biochemistry cell

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