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Preventing and treating missing data in longitudinal clinical trials: A practical guide / Craig Mallinckrodt. pages cm â€œ (Practical guides to biostatistics and epidemiology)

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the potential impact and treatment of missing data. 2 Investigators, sponsors, and regulators should design clinical trials consistent with the goal of maximizing the number of participants who are maintained on the protocol-specified intervention until the outcome data are collected.

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Quick summary. Material presented by Craig Mallinckrodt and Russ Wolfinger in 2013 at the FDA workshop on missing data. Download. The pdf file is available in zip format at [training_2013FDA_Workshop](#)

A Practical Guide to Preventing and Treating Missing Data

This book focuses on the prevention and treatment of missing data in longitudinal clinical trials. Based on his extensive experience with missing data, the author offers advice on choosing analysis methods and on ways to prevent missing data through appropriate trial design and conduct.

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"Dr. Mallinckrodt is, as usual, a paragon of clear writing and even clearer thinking. His commonsense, practical approach further elucidates what could otherwise be intractably co

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Recent research has fostered new guidance on preventing and treating missing data, most notably the landmark expert panel report from the National Research Council (NRC) that was commissioned by FDA. One of the findings from that panel was the need for better software tools to conduct missing data sensitivity analyses and frameworks for drawing inference from them.

Recent Developments in the Prevention and Treatment of

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