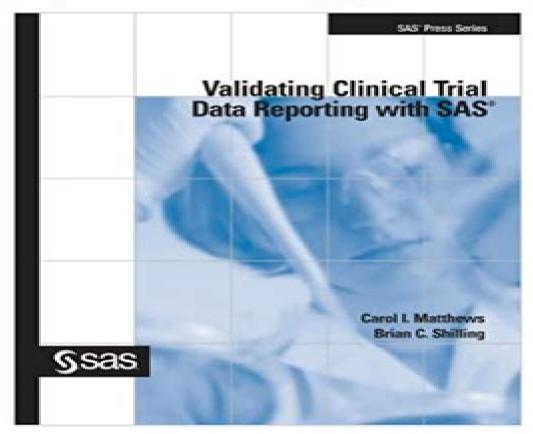
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Validating Clinical Trial Data Reporting With Sas

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Validating Clinical Trial Data Reporting With Sas:

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Checks for CDISC Compliance Using SAS Sunil Gupta, 2019-09-23 Clinical Data Quality Checks for CDISC Compliance using SAS is the first book focused on identifying and correcting data quality and CDISC compliance issues with real world innovative SAS programming techniques such as Proc SQL metadata and macro programming Learn to master Proc SQL s subqueries and summary functions for multi tasking process Drawing on his more than 25 years experience in the pharmaceutical industry the author provides a unique approach that empowers SAS programmers to take control of data quality and CDISC compliance This book helps you create a system of SDTM and ADaM checks that can be tracked for continuous improvement How often have you encountered issues such as missing required variables duplicate records invalid derived variables and invalid sequence of two dates With the SAS programming techniques introduced in this book you can start to monitor these and more complex data and CDISC compliance issues With increased standardization in SDTM and ADaM specifications and data values codelist dictionaries can be created for better organization planning and maintenance This book includes a SAS program to create excel files containing unique values from all SDTM and ADaM variables as columns In addition another SAS program compares SDTM and ADaM codelist dictionaries with codelists from define xml specifications Having tools to automate this process greatly saves time from doing it manually Features SDTMs and ADaMs Vitals SDTMs and ADaMs Data CDISC Specifications Compliance CDISC Data Compliance Protocol Compliance Codelist **Drug Discovery and Clinical Research** SK Gupta, 2011-06 The Drug Discovery and Clinical **Dictionary Compliance** Research bandwagon has been joined by scientists and researchers from all fields including basic sciences medical sciences biophysicists biotechnologists statisticians regulatory officials and many more The joint effort and contribution from all is translating into the fast development of this multi faceted field At the same time it has become challenging for all stakeholders to keep abreast with the explosion in information The race for the finish line leaves very little time for the researchers to update themselves and keep tabs on the latest developments in the industry To meet these challenges this book entitled Drug Discovery and Clinical Research has been compiled All chapters have been written by stalwarts of the field who have their finger on the pulse of the industry The aim of the book is to provide succinctly within one cover an update on all aspects of this wide area Although each of the chapter dealt here starting from drug discovery and development clinical development bioethics medical devices pharmacovigilance data management safety monitoring patient recruitment etc are topics for full fledged book in themselves an effort has been made via this book to provide a bird s eye view to readers and help them to keep abreast with the latest development despite constraints of time It is hoped that the book will contribute to the growth of readers which should translate into drug discovery and clinical research industry s growth

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