

SAS® Press Series

Validating Clinical Trial Data Reporting with SAS®

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Validating Clinical Trial Data Reporting with SAS Carol I. Matthews, Brian C. Shilling, 2008 This indispensable guide focuses on validating programs written to support the clinical trial process from after the data collection stage to generating reports and submitting data and output to the Food and Drug Administration *Validating Clinical Trial Data Reporting with SAS (Hardcover Edition)* Carol I. Matthews, Brian C. Shilling, 2008-03-17 Validation is a critical component to programming clinical trial analysis Essential to effective validation is the programmer's understanding of the data with which they'll be working If you don't understand how the data is arranged the values that are reasonable for each variable and the way the data should behave you cannot ensure that the final result of your programming effort is complete or even appropriate Therefore to be a successful programmer in the pharmaceutical industry you need to understand validation requirements and to learn how to make the code do the bulk of the work so that your programs are efficient as well as accurate This indispensable guide focuses on validating programs written to support the clinical trial process from after the data collection stage to generating reports and submitting data and output to the Food and Drug Administration FDA Authors Carol Matthews and Brian Shilling provide practical examples explanations for why different techniques are helpful and tips for avoiding errors in your output Topics addressed include Validation and pharmaceutical industry overviews Documentation and maintenance requirements discussions General techniques to facilitate validation Data importing and exporting Common data types Reporting and statistics *Validating Clinical Trial Data Reporting with SAS* is designed for SAS programmers who are new to the pharmaceutical industry as well as for those seeking a good foundation for validation in the SAS programming arena Readers should have a working knowledge of Base SAS and a basic understanding of programming tasks in the pharmaceutical industry **SAS Programming in the Pharmaceutical Industry, Second Edition** Jack Shostak, 2014-03-01 This comprehensive resource provides on the job training for statistical programmers who use SAS in the pharmaceutical industry This one stop resource offers a complete review of what entry to intermediate level statistical programmers need to know in order to help with the analysis and reporting of clinical trial data in the pharmaceutical industry *SAS Programming in the Pharmaceutical Industry Second Edition* begins with an introduction to the pharmaceutical industry and the work environment of a statistical programmer Then it gives a chronological explanation of what you need to know to do the job It includes information on importing and massaging data into analysis data sets producing clinical trial output and exporting data This edition has been updated for SAS 9.4 and it features new graphics as well as all new examples using CDISC SDTM or ADaM model data structures Whether you're a novice seeking an introduction to SAS programming in the pharmaceutical industry or a junior level programmer exploring new approaches to problem solving this real world reference guide offers a wealth of practical suggestions to help you sharpen your skills This book is part of the SAS Press program **Data Preparation for Analytics Using SAS** Gerhard Svolba, 2006-11-01 Text addresses such tasks as viewing

analytic data preparation in the context of its business environment identifying the specifics of predictive modeling for data mart creation understanding the concepts and considerations of data preparation for time series analysis and using SAS procedures for scoring

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Clinical Data Quality 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 Dictionary Compliance *Drug Discovery and Clinical Research* SK Gupta, 2011-06 The Drug Discovery and Clinical 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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and validating SDTM and ADaM data determining solutions for legacy data conversions and preparing data for regulatory submission The book covers products such as Base SAS SAS Clinical Data Integration and the SAS Clinical Standards Toolkit as well as JMP Clinical Topics included in this new edition include an implementation of the Define XML 2.0 standard new SDTM domains validation with Pinnacle 2.1 software event narratives in JMP Clinical and of course new versions of SAS and JMP software Any manager or user of clinical trial data in this day and age is likely to benefit from knowing how to either put data into a CDISC standard or analyzing and finding data once it is in a CDISC format If you are one such person a data manager clinical and or statistical programmer biostatistician or even a clinician then this book is for you [InfoWorld](#)

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Cancer Treatment Dirk Van Gestel,Jan Baptist Vermorken,2021-03-30

Risk-Based Monitoring and Fraud Detection

in Clinical Trials Using JMP and SAS Richard C. Zink,2014-07-01 Improve efficiency while reducing costs in clinical trials with centralized monitoring techniques using JMP and SAS International guidelines recommend that clinical trial data should be actively reviewed or monitored the well being of trial participants and the validity and integrity of the final analysis results are at stake Traditional interpretation of this guidance for pharmaceutical trials has led to extensive on site monitoring including 100% source data verification On site review is time consuming expensive estimated at up to a third of the cost of a clinical trial prone to error and limited in its ability to provide insight for data trends across time patients and clinical sites In contrast risk based monitoring RBM makes use of central computerized review of clinical trial data and site metrics to determine if and when clinical sites should receive more extensive quality review or intervention Risk Based Monitoring and Fraud Detection in Clinical Trials Using JMP and SAS presents a practical implementation of methodologies within JMP Clinical for the centralized monitoring of clinical trials Focused on intermediate users this book describes analyses for RBM that incorporate and extend the recommendations of TransCelerate Biopharm Inc methods to detect potential patient or investigator misconduct snapshot comparisons to more easily identify new or modified data and other novel visual and analytical techniques to enhance safety and quality reviews Further discussion highlights recent regulatory guidance documents on risk based approaches addresses the requirements for CDISC data and describes methods to supplement analyses with data captured external to the study database Given the interactive dynamic and graphical nature of JMP Clinical any individual from the clinical trial team including clinicians statisticians data managers programmers regulatory associates and monitors can make use of this book and the numerous examples contained within to streamline accelerate and enrich their reviews of clinical trial data The analytical methods described in Risk Based Monitoring and Fraud Detection in Clinical Trials Using JMP and SAS enable the clinical trial team to take a proactive approach to data quality and safety to streamline clinical development activities and address shortcomings while the study is ongoing This book is part of the SAS

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