

DB Software Laboratory

Phil Trinder

Users Manual for Nevada Test Site Database (NTS-DB) Software Howard Grantham White,1997

Data Collection Procedures for the Software Engineering Laboratory (Sel) Database National Aeronautics and Space Administration (NASA),2018-07-17 This document is a guidebook to collecting software engineering data on software development and maintenance efforts, as practiced in the Software Engineering Laboratory (SEL). It supersedes the document entitled Data Collection Procedures for the Rehosted SEL Database, number SEL-87-008 in the SEL series, which was published in October 1987. It presents procedures to be followed on software development and maintenance projects in the Flight Dynamics Division (FDD) of Goddard Space Flight Center (GSFC) for collecting data in support of SEL software engineering research activities. These procedures include detailed instructions for the completion and submission of SEL data collection forms. Heller, Gerard and Valett, Jon and Wild, Mary Goddard Space Flight Center COMPUTER PROGRAMMING; DATA ACQUISITION; DATA BASES; PROGRAMMING ENVIRONMENTS; SOFTWARE ENGINEERING; AERODYNAMICS; HANDBOOKS; PRODUCT DEVELOPMENT...

LIMS A S Nakagawa,2007-10-31 There is currently a high level of interest in Laboratory Information Management Systems (LIMS), which, when successfully implemented, can revitalize the operations of a laboratory and contribute significantly to the effectiveness and efficiency of the overall enterprise. LIMS describes the strategy, planning, resources, and activities needed to integrate LIMS and its supporting technologies into an organization. It covers all aspects of implementation and management and has the benefit of not being product specific. This book will not date as it is not restricted to a particular software product, hardware platform, or technical automation approach. Instead it deals with the issues, expertise, organization, and resources that contribute to the successful implementation of LIMS. The author has wide experience of automated laboratory systems in the chemical, pharmaceutical, environmental, and biotechnology industries, and for the past 15 years has been intimately involved in every aspect of LIMS implementations including justification, system selection, installation, project management, developing, training, validation, performance optimization, and maintenance. LIMS contains numerous illustrations and tables to highlight concisely the major points and concepts discussed in each chapter. The book is essential reading for laboratory, information systems and project managers responsible for the implementation of LIMS and, as it does not require any previous knowledge of computers or laboratory information management systems, is easily accessible to all.

Databases on Modern Hardware Anastasia Ailamaki,Erietta Liarou,Pinar Tözün,Danica Porobic,Iraklis Psaroudakis,2017-08-14 Data management systems enable various influential applications from high-performance online services (e.g., social networks like Twitter and Facebook or financial markets) to big data analytics (e.g., scientific exploration, sensor networks, business intelligence). As a result, data management systems have been one of the main drivers for innovations in the database and computer architecture communities for several decades. Recent hardware trends require software to take advantage of the abundant parallelism existing in modern and future hardware. The traditional design of the data management systems, however, faces inherent scalability problems due to its tightly coupled components. In addition, it cannot exploit the full capability of the aggressive micro-architectural features of modern processors. As a result, today's most commonly used server types remain largely underutilized leading to a huge waste of hardware resources and energy. In this book, we shed light on the challenges present while running DBMS on modern multicore hardware. We divide the material into two dimensions of scalability: implicit/vertical and explicit/horizontal. The first part of the book focuses on the vertical dimension: it describes the instruction- and data-level parallelism opportunities in a core coming from the hardware and software side. In addition, it examines the sources of under-utilization in a modern processor and presents insights and hardware/software techniques to better exploit the microarchitectural resources of a processor by improving cache locality at the right level of the memory hierarchy. The second part focuses on the horizontal dimension, i.e., scalability bottlenecks of database applications at the level of multicore and multsocket multicore architectures. It first presents a systematic way of eliminating such bottlenecks in online transaction processing workloads, which is based on minimizing unbounded communication, and shows several techniques that minimize bottlenecks in major components of database management systems. Then, it demonstrates the data and work sharing opportunities for analytical workloads, and reviews advanced scheduling mechanisms that are aware of nonuniform memory accesses and alleviate bandwidth saturation.

A Functional Database Phil Trinder,1989

Computer Medical Databases Morris F. Collen,2011-10-20 Chapter 1 offers an overview of the basic computer technology. Each succeeding chapter, describes the problems in medicine, followed by a review in chronological sequence of why and how computers were applied to try to meet these problems. Only the technical aspects of computer hardware, software, and communications are discussed as they are necessary to explain how the technology was applied. This approach generally led to defining the objectives for applications of medical informatics. At the end of each chapter, the author summarizes his personal views and interpretations of the chapter contents. Although the concurrent evolution of medical informatics in Canada, Europe, and Japan certainly influenced workers in the United States, the scope of this historical review is limited to the development of medical informatics within the United States. Furthermore, this review is limited to electronic digital computers; it excludes mechanical, analog, and hybrid computers.

Laboratory Management Denise Harmening,2007 The laboratory environment is ever changing in response to the diverging trends in healthcare. Laboratory managers who can create solutions to today's problems and effectively manage change are in high demand. The second edition of Denise Harmening's Laboratory Management is designed to give a problem-based approach to teaching the principles of laboratory management. the text focuses on presenting underlying managerial concepts and assisting the learner in successfully applying theoretical models to real-life situations.

Query Processing in Database Systems W. Kim,D.S. Reiner,Don Batory,2012-12-06 This book is an anthology of the results of research and development in database query processing during the past decade. The relational model of data provided tremendous impetus for research into query processing. Since a relational query does not specify access paths to the stored data, the database management system (DBMS) must provide an intelligent query-processing subsystem which will evaluate a number of potentially efficient strategies for processing the query and select the one that optimizes a given performance measure. The degree of sophistication of this subsystem, often called the optimizer, critically affects the performance of the DBMS. Research into query processing thus started has taken off in several directions during the past decade. The emergence of research into distributed databases has enormously complicated the tasks of the optimizer. In a distributed environment, the database may be partitioned into horizontal or vertical fragments of relations. Replicas of the fragments may be stored in different sites of a network and even migrate to other sites. The measure of performance of a query in a distributed system must include the communication cost between sites. To minimize communication costs for-queries involving multiple relations across multiple sites, optimizers may also have to consider semi-join techniques.

Laboratory Information Management Systems Christine Paszko,Elizabeth Turner,2018-06-08 Details the most recent advances in Laboratory Information Management Systems. Offers contemporary approaches to system development, design, and installation; system customization; software and hardware compatibility; quality assurance and regulatory requirements; and resource utilization.

FSL In Review, Forecast Systems Laboratory, 1999 - 2000 ,2000

Pathology Informatics, An Issue of the Clinics in Laboratory Medicine Anil V. Parwani,2016-02-10 This issue of the Clinics in Laboratory Medicine, edited by Dr. Anil Parwani, is a special issue is devoted to topics in Pathology Informatics. Topics include but are not limited to: Basics of Information Systems (Hardware, Software); Networks, Interfaces and Communications; Databases; Data Representation, Coding and Communication Standards; Laboratory Information Systems; Enhancing and Customizing Laboratory Information Systems; Laboratory Management and Operations; Specialized Laboratory Information Systems; Middleware and Laboratory Automation; Bar Coding and Tracking; Molecular Pathology Informatics; Pathology Informatics and Project Management; Digital Imaging; Telepathology; Healthcare Information Systems; Data Security and Reliability; Role of Informatics in Patient Safety and Quality Assurance; Role of Pathology Informatics in IT Leadership; Selection and Implementation of New Information Systems; Biomedical Informatics and Research Informatics; Training in Pathology Informatics; and more.

Henry's Clinical Diagnosis and Management by Laboratory Methods E-Book Richard A. McPherson, Matthew R. Pincus, 2021-06-09 For more than 100 years, Henry's Clinical Diagnosis and Management by Laboratory Methods has been recognized as the premier text in clinical laboratory medicine, widely used by both clinical pathologists and laboratory technicians. Leading experts in each testing discipline clearly explain procedures and how they are used both to formulate clinical diagnoses and to plan patient medical care and long-term management. Employing a multidisciplinary approach, it provides cutting-edge coverage of automation, informatics, molecular diagnostics, proteomics, laboratory management, and quality control, emphasizing new testing methodologies throughout. Remains the most comprehensive and authoritative text on every aspect of the clinical laboratory and the scientific foundation and clinical application of today's complete range of laboratory tests. Updates include current hot topics and advances in clinical laboratory practices, including new and extended applications to diagnosis and management. New content covers next generation mass spectroscopy (MS), coagulation testing, next generation sequencing (NGS), transfusion medicine, genetics and cell-free DNA, therapeutic antibodies targeted to tumors, and new regulations such as ICD-10 coding for billing and reimbursement. Emphasizes the clinical interpretation of laboratory data to assist the clinician in patient management. Organizes chapters by organ system for quick access, and highlights information with full-color illustrations, tables, and diagrams. Provides guidance on error detection, correction, and prevention, as well as cost-effective test selection. Includes a chapter on Toxicology and Therapeutic Drug Monitoring that discusses the necessity of testing for therapeutic drugs that are more frequently being abused by users.

Establishing A CGMP Laboratory Audit System David M. Bliesner, 2006-04-28 The first systematic, hands-on auditing guide for today's pharmaceutical laboratories In today's litigious environment, pharmaceutical laboratories are subject to ever stricter operational guidelines as mandated by the FDA, and must be able to establish and demonstrate sustainable operational practices that ensure compliance with the current good manufacturing practice (CGMP) regulations. David Bliesner's *Establishing a CGMP Laboratory Audit System: A Practical Guide* is designed to provide laboratory supervisors and personnel with a step-by-step, hands-on audit system that they can rely on to ensure their facility remains compliant with all current and future requirements. Focusing on a team approach, the author uses detailed flowcharts, checklists, and descriptions of the auditing process to help readers develop a new audit system or upgrade their current system in order to: * Improve current compliance * Demonstrate sustainable compliance * Produce data for federal inspections * Avoid regulatory action Enhanced with detailed checklists and a wealth of practical and flexible auditing tools on CD-ROM, this book provides an ideal resource for new and future laboratory personnel, and an excellent means for keeping existing industry practitioners up to date on the nuances of operating a consistently compliant pharmaceutical laboratory.

Database and Expert Systems Applications Dimitris Karagiannis, 2013-11-11 The Database and Expert Systems Applications - DEXA - conferences are dedicated to providing an international forum for the presentation of applications in the database and expert systems field, for the exchange of ideas and experiences, and for defining requirements for the future systems in these fields. After the very promising DEXA 90 in Vienna, Austria, we hope to have successfully established with this year's DEXA 91 a stage where scientists from diverse fields interested in application-oriented research can present and discuss their work. This year there was a total of more than 250 submitted papers from 28 different countries, in all continents. Only 98 of the papers could be accepted. The collection of papers in these proceedings offers a cross-section of the issues facing the area of databases and expert systems, i.e., topics of basic research interest on one hand and questions occurring when developing applications on the other. Major credit for the success of the conference goes to all of our colleagues who submitted papers for consideration and to those who have organized and chaired the panel sessions. Many persons contributed numerous hours to organize this conference. The names of most of them will appear on the following pages. In particular we wish to thank the Organization Committee Chairmen Johann Gordes, A Min Tjoa, and Roland Wagner, who also helped establishing the program. Special thanks also go to Gabriella Wagner and Anke Ruckert. Dimitris Karagiannis General Conference Chairman Contents Conference Committee.

Clinical Pathology, An Issue of the Clinics in Laboratory Medicine E-Book Geza S Bodor, 2018-08-19 This issue of Clinics in Laboratory Medicine will focus on Clinical Pathology and is edited by Geza S. Bodor. Topics include, but are not limited to, Steroid measurement / Salivary cortisol measurement, Protein testing by LCMSMS, LCMSMS in the Clinical Laboratory, Laboratory Standards for Clinical LCMSMS, The need to teach LCMSMS to clinical laboratory scientists, MALDI-TOF in the clinical laboratory, MALDI TOF MS in the clinical microbiology laboratory, LCMSMS method development consideration in clinical laboratory practice, Cancer diagnosis using mass spectrometry, Adulteration and LCMSMS drug testing, Diagnosis of inherited metabolic disorders using LCMSMS, Harmonization of LCMSMS protein assays, Vitamin D testing by LCMSMS versus by immunoassay, Pain management testing by LCMSMS, and Development of FDA approved clinical mass spectrometer.

RDF Database Systems Olivier Curé, Guillaume Blin, 2014-11-24 RDF Database Systems is a cutting-edge guide that distills everything you need to know to effectively use or design an RDF database. This book starts with the basics of linked open data and covers the most recent research, practice, and technologies to help you leverage semantic technology. With an approach that combines technical detail with theoretical background, this book shows how to design and develop semantic web applications, data models, indexing and query processing solutions. Understand the Semantic Web, RDF, RDFS, SPARQL, and OWL within the context of relational database management and NoSQL systems Learn about the prevailing RDF triples solutions for both relational and non-relational databases, including column family, document, graph, and NoSQL Implement systems using RDF data with helpful guidelines and various storage solutions for RDF Process SPARQL queries with detailed explanations of query optimization, query plans, caching, and more Evaluate which approaches and systems to use when developing Semantic Web applications with a helpful description of commercial and open-source systems

The Laboratory Rabbit, Guinea Pig, Hamster, and Other Rodents Mark A. Suckow, Karla A. Stevens, Ronald P. Wilson, 2012-01-09 This is a single volume, comprehensive book sanctioned by the American College of Laboratory Animal Medicine (ACLAM), covering the rabbit, guinea pig, hamster, gerbil and other rodents often used in research. This well illustrated reference includes basic biology, anatomy, physiology, behavior, infectious and noninfectious diseases, husbandry and breeding, common experimental methods, and use of the species as a research model. It is a resource for advancements in the humane and responsible care of: rabbit, guinea pig, hamster, gerbil, chinchilla, deer mouse, kangaroo rat, cotton rat, sand rat, and degu Includes up-to-date, common experimental methods. Organized by species for easy access during bench research.

Metadata-driven Software Systems in Biomedicine Prakash M. Nadkarni, 2011-05-27 While the use of database technology is ubiquitous throughout IT (and health IT in particular), it is not generally appreciated that, as a database increases in scope, certain designs are far superior to others. In biomedical domains, new knowledge is being generated continually, and the databases that must support areas such as clinical care and research must also be able to evolve while requiring minimal or no logical / physical redesign. Appropriately designed metadata, and software designed to utilize it effectively, can provide significant insulation against change. Many of the larger EMR or clinical research database vendors have realized this, but their designs are proprietary and not described in the literature. Consequently, numerous misconceptions abound among individuals who have not had to work with large-scale biomedical systems, and graduates of a health or bioinformatics program may find that they need to unlearn what they were taught in database and software design classes in order to work productively with such systems. A working knowledge of such systems is also important for individuals who are not primarily software developers, such as health informaticians, medical information officers and data analysts. This book is, in a sense, intended to prepare all of the above individuals for the real world.

Prudent Practices in the Laboratory National Research Council, Division on Earth and Life Studies, Board on Chemical Sciences and Technology, Committee on Prudent Practices in the Laboratory: An Update, 2011-03-25 Prudent Practices in the Laboratory--the book that has served for decades as the standard for chemical laboratory safety practice--now features updates and new topics. This revised edition has an expanded chapter on chemical management and delves into new areas, such as nanotechnology, laboratory security, and emergency planning. Developed by experts from academia and industry, with specialties in such areas as chemical sciences, pollution prevention, and laboratory safety, Prudent Practices in the Laboratory provides guidance on planning procedures for the handling, storage, and disposal of chemicals. The book offers prudent practices designed to promote safety and includes practical information on assessing hazards, managing chemicals, disposing of wastes, and more.

Prudent Practices in the Laboratory will continue to serve as the leading source of chemical safety guidelines for people working with laboratory chemicals: research chemists, technicians, safety officers, educators, and students.

Laboratory Control System Operations in a GMP Environment David M. Bliesner,2020-04-21 Develop an understanding of FDA and global regulatory agency requirements for Laboratory Control System (LCS) operations In *Laboratory Control System Operations in a GMP Environment*, readers are given the guidance they need to implement a CGMP compliant Laboratory Control System (LCS) that fits within Global Regulatory guidelines. Using the Quality Systems Approach, regulatory agencies like the FDA and the European Medicine Agency have developed a scheme of systems for auditing pharmaceutical manufacturing facilities which includes evaluating the LCS. In this guide, readers learn the fundamental rules for operating a CGMP compliant Laboratory Control System. Designed to help leaders meet regulatory standards and operate more efficiently, the text includes chapters that cover Laboratory Equipment Qualification and Calibration, Laboratory Facilities, Method Validation and Method Transfer, Laboratory Computer Systems, Laboratory Investigations as well as Data Governance and Data Integrity. The text also includes chapters related to Laboratory Managerial and Administrative Systems, Laboratory Documentation Practices and Standard Operating Procedures and General Laboratory Compliance Practices. Additionally, a chapter outlining Stability Program operations is included in the text. In addition to these topics, it includes LCS information and tools such as: ● End of chapter templates, checklists, and LCS guidance to help you follow the required standards ● Electronic versions of each tool so users can use them outside of the text ● An In-depth understanding of what is required by the FDA and other globally significant regulatory authorities for GMP compliant systems For quality assurance professionals working within the pharmaceutical or biopharma industries, this text provides the insight and tools necessary to implement government-defined regulations.

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