

Amgen Medical Information

Amgen Inc. (Thousand Oaks, Calif.),David Ewing Duncan

Biotechnology Ronald P. Evens,2020-06-04 The over-riding premise for biotechnology in this book is bringing novel products to market to substantially advance patient care and disease mitigation. Biotechnology, over its relatively brief existence of 40 years, has experienced a mercurial growth. The vast educational need for biotechnology information in this rapidly burgeoning field is a basic rationale here. However a more prominent underpinning is that, bringing biotech products to market for patient care involves success in the following four areas of engagement simultaneously - scientific advances for healthcare technologies, novel and varied products for untreated diseases, regulatory authorities, and biotech companies. Features Comprehensive coverage of biotechnology science topics used in development and manufacturing Addresses all the scientific technologies within biotechnology responsible for products on the market and the pipeline Presents business issues such as marketing and sales of the products, as well as companies engaged, and how biotech business has evolved

The Amgen Story Amgen Inc. (Thousand Oaks, Calif.),David Ewing Duncan,2005 As the most successful biotech company in history and the eighth largest drug-producing company in the world, Amgen has improved the lives of millions of patients worldwide. In 2005, the company celebrates its twenty-fifth anniversary with the publication of The Amgen Story. This stunning illustrated book contains hundreds of archival photos and compelling text from noted biotech writer David Ewing Duncan. It is testament and tribute to the staff, leaders, patients, and science that make each discovery possible.

Telemedicine and E-health Law Lynn D. Fleisher,James C. Dechene,2004 Telemedicine and E-Health Law has the answers that health care providers, hospitals, pharmaceutical companies, insurers and their legal counsel need as medicine enters a new era.

Buyer Beware United States. Congress. Senate. Special Committee on Aging,2002

ASHP Injectable Drug Information American Society of Health-System Pharmacists,2021-02-05 ASHP's New and Expanded Guide to IV Compatibility & Stability For more than 40 years, ASHP has published the most trusted resource for injectable drug information. Our new ASHP® Injectable Drug Information™ now delivers the same high-quality content that you can expect from ASHP with even MORE of the information you need to make informed patient care decisions. For the first time ever, this gold standard reference is available as an eBook with new and expanded information. The 2021 edition features 18 new monographs, and nearly 200 new references for a total of over 24,000 total compatibility pairs. Backed by quality, peer-reviewed published literature, and authored under the editorial authority of ASHP, ASHP® Injectable Drug Information™ is a must-have resource for every pharmacy.

Real-World Evidence Generation and Evaluation of Therapeutics National Academies of Sciences, Engineering, and Medicine,Health and Medicine Division,Board on Health Sciences Policy,Forum on Drug Discovery, Development, and Translation,2017-07-05 The volume and complexity of information about individual patients is greatly increasing with use of electronic records and personal devices. Potential effects on medical product development in the context of this wealth of real-world data could be numerous and varied, ranging from the ability to determine both large-scale and patient-specific effects of treatments to the ability to assess how therapeutics affect patients' lives through measurement of lifestyle changes. In October 2016, the National Academies of Sciences, Engineering, and Medicine held a workshop to facilitate dialogue among stakeholders about the opportunities and challenges for incorporating real-world evidence into all stages in the process for the generation and evaluation of therapeutics. Participants explored unmet stakeholder needs and opportunities to generate new kinds of evidence that meet those needs. This publication summarizes the presentations and discussions from the workshop.

Fiscal Year 1991 Reconciliation Issues Relating to Medicare Waste and Abuse United States. Congress. House. Committee on Ways and Means. Subcommittee on Health,1990

Chronic Disease Nursing Susan Oliver,2006-02-03 Individuals with chronic diseases have to cope with various challenges to their physical and psychological well being as part of their daily lives. In the field of chronic disease management nurses have an integral role in supporting the needs of the patient and identifying the optimum in high quality care. Dealing with the increasing needs of the growing chronic disease and elderly populations presents difficult challenges. Resources need to be used effectively. Chronic Disease Nursing offers support to all nurses but particularly primary and secondary care advanced practitioners wishing to develop an effective system of care for those with a long term medical condition. There is guidance on the practical aspects of setting up a telephone helpline service, identifying the ethical and professional aspects of developing a nurse led clinic, selecting appropriate outcome measures for clinic setting, and suitable biologic therapies used in chronic disease areas and working across professional organisations. This book provides a framework for the development of a chronic disease nursing service, using rheumatology as an example.

Orphan Drug Act United States. Congress. House. Committee on Energy and Commerce. Subcommittee on Health and the Environment,1990

New Scientist ,2002

New Scientist and Science Journal ,2000

Skyrocketing Prescription Drug Prices United States. Congress. Senate. Special Committee on Aging,1990

Medical Affairs Kirk V. Shepard,Charlotte Kremer,Garth Sundem,2024-01-30 Medical Affairs is one of the three strategic pillars of the pharmaceutical and MedTech industries, but while clear career paths exist for Commercial and Research and Development, there is no formal training structure for Medical Affairs professionals. Medical and scientific expertise is a prerequisite for entry into the function, and many people transitioning into Medical Affairs have advanced degrees such as PhD, MD, or PharmD. However, these clinical/scientific experts may not be especially well-versed in aspects of industry such as the drug development lifecycle, crossfunctional collaborations within industry, and digital tools that are transforming the ways Medical Affairs generates and disseminates knowledge. This primer for aspiring and early-career Medical Affairs professionals equips readers with the baseline skills and understanding to excel across roles. Features: Defines the purpose and value of Medical Affairs and provides clear career paths for scientific experts seeking their place within the pharmaceutical and MedTech industries. Provides guidance and baseline competencies for roles within Medical Affairs including Medical Communications, Evidence Generation, Field Medical, Compliance, and many others. Specifies the true north of the Medical Affairs profession as ensuring patients receive maximum benefit from industry innovations including drugs, diagnostics and devices. Presents the purpose and specific roles of Medical Affairs roles across organization types including biotechs, small/medium/large pharma and device/diagnostic companies, taking into account adjustments in the practice of Medical Affairs to meet the needs of developing fields such as rare disease and gene therapy. Leverages the expertise of over 60 Medical Affairs leaders across companies, representing the first unified, global understanding of the Medical Affairs profession.

Proceedings of the ... Public Health Conference on Records and Statistics ,

Transforming Clinical Research in the United States Institute of Medicine,Board on Health Sciences Policy,Forum on Drug Discovery, Development, and Translation,2010-10-22 An ideal

health care system relies on efficiently generating timely, accurate evidence to deliver on its promise of diminishing the divide between clinical practice and research. There are growing indications, however, that the current health care system and the clinical research that guides medical decisions in the United States falls far short of this vision. The process of generating medical evidence through clinical trials in the United States is expensive and lengthy, includes a number of regulatory hurdles, and is based on a limited infrastructure. The link between clinical research and medical progress is also frequently misunderstood or unsupported by both patients and providers. The focus of clinical research changes as diseases emerge and new treatments create cures for old conditions. As diseases evolve, the ultimate goal remains to speed new and improved medical treatments to patients throughout the world. To keep pace with rapidly changing health care demands, clinical research resources need to be organized and on hand to address the numerous health care questions that continually emerge. Improving the overall capacity of the clinical research enterprise will depend on ensuring that there is an adequate infrastructure in place to support the investigators who conduct research, the patients with real diseases who volunteer to participate in experimental research, and the institutions that organize and carry out the trials. To address these issues and better understand the current state of clinical research in the United States, the Institute of Medicine's (IOM) Forum on Drug Discovery, Development, and Translation held a 2-day workshop entitled Transforming Clinical Research in the United States. The workshop, summarized in this volume, laid the foundation for a broader initiative of the Forum addressing different aspects of clinical research. Future Forum plans include further examining regulatory, administrative, and structural barriers to the effective conduct of clinical research; developing a vision for a stable, continuously funded clinical research infrastructure in the United States; and considering strategies and collaborative activities to facilitate more robust public engagement in the clinical research enterprise.

Science Lessons Gordon M. Binder,Philip Bashe,2008 Under Gordon Binder's leadership, Amgen became the world's largest and most successful biotech company in the world. This text describes what it really takes to manage risk, financing, creative employees, and intellectual property on the international stage.

Fulfilling the Potential of Cancer Prevention and Early Detection National Research Council,Institute of Medicine,National Cancer Policy Board,2003-05-07 Cancer ranks second only to heart disease as a leading cause of death in the United States, making it a tremendous burden in years of life lost, patient suffering, and economic costs. Fulfilling the Potential for Cancer Prevention and Early Detection reviews the proof that we can dramatically reduce cancer rates. The National Cancer Policy Board, part of the Institute of Medicine, outlines a national strategy to realize the promise of cancer prevention and early detection, including specific and wide-ranging recommendations. Offering a wealth of information and directly addressing major controversies, the book includes: A detailed look at how significantly cancer could be reduced through lifestyle changes, evaluating approaches used to alter eating, smoking, and exercise habits. An analysis of the intuitive notion that screening for cancer leads to improved health outcomes, including a discussion of screening methods, potential risks, and current recommendations. An examination of cancer prevention and control opportunities in primary health care delivery settings, including a review of interventions aimed at improving provider performance. Reviews of professional education and training programs, research trends and opportunities, and federal programs that support cancer prevention and early detection. This in-depth volume will be of interest to policy analysts, cancer and public health specialists, health care administrators and providers, researchers, insurers, medical journalists, and patient advocates.

Ethics and Information Technology James G. Anderson,Kenneth Goodman,2007-05-28 This series is directed to health care professionals who are leading the tra- formation of health care by using information and knowledge. Launched in 1988 as Computers in Health Care, the series offers a broad range of titles: some addressed to specific professions such as nursing, medicine, and health administration; others to special areas of practice such as trauma and radi- ogy. Still other books in the series focus on interdisciplinary issues, such as the computer-based patient record, electronic health records, and networked health care systems. Renamed Health Informatics in 1998 to reflect the rapid evolution in the discipline now known as health informatics, the series will continue to add titles that contribute to the evolution of the field. In the series, eminent - perts, serving as editors or authors, offer their accounts of innovations in health informatics. Increasingly, these accounts go beyond hardware and so- ware to address the role of information in influencing the transformation of healthcare delivery systems around the world. The series also increasingly focuses on “peopleware” and the organizational, behavioral, and societal changes that accompany the diffusion of information technology in health services environments.

Physicians' Desk Reference ,2009

Biotechnology Ronald P. Evens,2020-06-04 The over-riding premise for biotechnology in this book is bringing novel products to market to substantially advance patient care and disease mitigation. Biotechnology, over its relatively brief existence of 40 years, has experienced a mercurial growth. The vast educational need for biotechnology information in this rapidly burgeoning field is a basic rationale here. However a more prominent underpinning is that, bringing biotech products to market for patient care involves success in the following four areas of engagement simultaneously - scientific advances for healthcare technologies, novel and varied products for untreated diseases, regulatory authorities, and biotech companies. Features Comprehensive coverage of biotechnology science topics used in development and manufacturing Addresses all the scientific technologies within biotechnology responsible for products on the market and the pipeline Presents business issues such as marketing and sales of the products, as well as companies engaged, and how biotech business has evolved

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