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This book constitutes the refereed proceedings of the First International Workshop on Data Integration in the Life Sciences, DILS 2004, held in Leipzig, Germany, in March 2004. The 13 revised full papers and 2 revised short papers presented were carefully reviewed and selected from many submissions. The papers are organized in topical sections on scientific and clinical workflows, ontologies and taxonomies, indexing and clustering, integration tools and systems, and integration techniques.

Never before have two revolutions with so much potential to save and prolong human life occurred simultaneously. The converging, synergistic power of the biochemical and digital revolutions now allows us to read every letter of life's code, create precisely targeted drugs to control it, and tailor their use to individual patients. Cancer, diabetes, Alzheimer's and countless other killers can be vanquished—if we make full use of the tools of modern drug design and allow doctors the use of modern data gathering and analytical tools when prescribing

drugs to their patients. But Washington stands in the way, clinging to outdated drug-approval protocols developed decades ago during medicine's long battle with the infectious epidemics of the past. Peter Huber, an expert in science, technology, and public policy, demonstrates why Washington's one-size-fits-all drug policies can't deal with diseases rooted in the complex molecular diversity of human bodies. Washington is ill-equipped to handle the torrents of data that now propel the advance of molecular medicine and is reluctant to embrace the statistical methods of the digital age that can. Obsolete economic policies, often rationalized as cost-saving measures, stifle innovation and suppress investment in the medicine that can provide the best cures at the lowest cost. In the 1980s, an AIDS diagnosis was a death sentence, until the FDA loosened its throttling grip and began streamlining and accelerating approval of life-saving drugs. *The Cure in the Code* shows patients, doctors, investors, and policy makers what we must now do to capture the full life-saving and cost-saving potential of the revolution in molecular medicine. America has to choose. At stake for America is the power to lead the world in mastering the most free, fecund, competitive, dynamic, and intelligent natural resource on the planet—the molecular code that spawns human life and controls our health.

"I thoroughly enjoyed reading this book as it has taken me on a journey through time, across the globe and through multiple disciplines. Indeed, we need to be thinking about these concepts and applying them every day to do our jobs better." Farah Magrabi, Macquarie University, Australia "The reader will find intriguing not only the title but also the content of the book. I'm also pleased that public health, and even more specifically epidemiology has an important place in this ambitious discussion." Elena Andresen, Oregon Health & Science University, USA "This book is very well written and addresses an important topic. It presents many reasons why basic scientists/researchers should establish collaborations and access information outside traditional means and not limit thinking but rather expand such and perhaps develop more innovative and translational research ventures that will advance science and not move it laterally." Gerald Pepe, Eastern Virginia Medical School, USA "This book gathers logically and presents interestingly (with many examples) the qualities and attitudes a researcher must possess in order to become successful. On the long run, the deep and carefully reexamined research will be the one that lasts." Zoltán Néda, Babeş-Bolyai University, Romania "I really liked the five pillars delineating the components of humanism in research. This book has made a major contribution to the research ethics literature." David Fleming, University of Missouri, USA A comprehensive review of the research phase of life sciences from design to discovery with suggestions to improve innovation This vital resource explores the creative processes leading to biomedical innovation, identifies the obstacles and best practices of innovative laboratories, and supports the production of effective science. *Innovative Research in Life Sciences* draws on lessons from 400 award-winning scientists and research from leading universities. The book explores the innovative process in life sciences and puts the focus on how great ideas are born and become landmark scientific discoveries. The text provides a unique resource for developing professional competencies and applied skills of life sciences researchers. The book examines what happens before the scientific paper is submitted for publication or the innovation becomes legally protected. This phase is the most neglected but most exciting in the process of scientific creativity and innovation. The author identifies twelve competencies of innovative biomedical researchers that described and analyzed. This important resource: Highlights the research phase from design to discovery that precedes innovation disclosure Offers a step by step explanation of how to improve innovation Offers solutions for improving research and innovation productivity in the life sciences Contains a variety of statistical databases and a vast number of stories about individual discoveries Includes a process of published studies and national statistics of biomedical research and reviews the performance of research labs and academic institutions Written for academics and researchers in biomedicine, pharmaceutical science, life sciences, drug discovery, pharmacology, *Innovative Research in Life Sciences* offers a guide to the creative processes leading to biomedical innovation and identifies the best practices of innovative scientists and laboratories.

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