HANDBOOK OF

Aquequs Solubility Data

SECOND EDITION

Samuel H. Yalkowsky Yan He Parijat Jain

(GRC Press

Handbook Of Aqueous Solubility Data Second Edition

Atun Roy Choudhury, Sankar Ganesh Palani

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guidance for human health consideration Discusses the latest advances and trends in solvent technology including modern methods of cleaning contaminated soils selection of gloves suits and respirators Handbook of Bioequivalence Testing, Second Edition Sarfaraz K. Niazi, 2014-10-29 As the generic pharmaceutical industry continues to grow and thrive so does the need to conduct adequate efficient bioequivalence studies In recent years there have been significant changes to the statistical models for evaluating bioequivalence In addition advances in the analytical technology used to detect drug and metabolite levels have made bioequivalence testing more complex The second edition of Handbook of Bioequivalence Testing has been completely updated to include the most current information available including new findings in drug delivery and dosage form design and revised worldwide regulatory requirements New topics include A historical perspective on generic pharmaceuticals New guidelines governing submissions related to bioequivalency studies along with therapeutic code classifications Models of noninferiority Biosimilarity of large molecule drugs Bioequivalence of complementary and alternate medicines Bioequivalence of biosimilar therapeutic proteins and monoclonal antibodies New FDA guidelines for bioanalytical method validation Outsourcing and monitoring of bioequivalence studies The cost of generic drugs is rising much faster than in the past partly because of the increased costs required for approval including those for bioequivalence testing There is a dire need to re examine the science behind this type of testing to reduce the burden of development costs allowing companies to develop generic drugs faster and at a lower expense The final chapter explores the future of bioequivalence testing and proposes radical changes in the process of biowaivers It suggests how the cost of demonstrating bioequivalence can be reduced through intensive analytical investigation and proposes that regulatory agencies reduce the need for bioequivalence studies in humans Backed by science and updated with the latest research this book is destined to spark continued debate on the efficacy of the current bioequivalence testing paradigm Handbook of Physical-Chemical Properties and Environmental Fate for Organic Chemicals, Second Edition Donald Mackay, Wan-Ying Shiu, Kuo-Ching Ma, Sum Chi Lee, 2006-03-14 Transport and transformation processes are key for determining how humans and other organisms are exposed to chemicals These processes are largely controlled by the chemicals physical chemical properties This new edition of the Handbook of Physical Chemical Properties and Environmental Fate for Organic Chemicals is a comprehensive series in four volumes that serves as a reference source for environmentally relevant physical chemical property data of numerous groups of chemical substances The handbook contains physical chemical property data from peer reviewed journals and other valuable sources on over 1200 chemicals of environmental concern The handbook contains new data on the temperature dependence of selected physical chemical properties which allows scientists and engineers to perform better chemical assessments for climatic conditions outside the 20 25 degree range for which property values are generally reported This second edition of the Handbook of Physical Chemical Properties and Environmental Fate for Organic Chemicals is an essential reference for university libraries regulatory agencies consultants and industry professionals particularly those

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the index and glossary have been greatly improved

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