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baumschlager eberle berlin be berlin was founded in 2010 by dietmar eberle and gerd jäger with a very specific mission in its early years this renowned architecture firm designed global competition bids for baumschlager eberle as a result of its successes especially in residential architecture in and around berlin the berlin office increasingly also took on the work of implementation planning the guiding maxim of be berlin has always been to focus on projects that will stand the test of time this dovetails with dietmar eberle s motto of building for everyday life in addition to about 40 project profiles this book contains an interview with gerd jäger by jürgen tietz an essay on residential construction in germany and especially berlin by gerd jäger and a photo essay by claudia klein monograph on be berlin specialized in the design of subsidized public housing in berlin and internationally supplement to the monograph baumschlager eberle architekten 2010 2020

the 6th edition of a well known and much used standard text in the field this book covers all aspects of the biochemical abnormalities caused by various diseases and how they relate to the biochemical changes in the blood urine cerebrospinal fluid joint fluids other body fluids and in cells the purpose is to provide the fundamental bases for understanding the biochemical changes that occur in disease processes and in turn to provide the rationale for applying this understanding to the diagnosis of the disease process a substantial appendix is provided so that the user can quickly identify the reference ranges for a large number of animal species an appendix is provided in the book so that the user can quickly identify the reference ranges for a large number of animal species explains what biochemical changes occur in disease processes and provides the rationale for applying this understanding to the diagnosis of the disease process

this book provides in depth information about common clinical laboratory assays that are used to evaluate domestic mammals including what assays measure sample or assay conditions that affect results and what results indicate about the physiologic or pathologic state of a patient whenever possible diseases and conditions are grouped by common mechanisms or processes to promote a conceptual understanding of laboratory data that can be generally applied across many species new to the second edition are additional disorders diagnostic tests illustrations images references and pathophysiologic explanations this text has proven valuable to students and veterinarians wanting a fundamental understanding of

veterinary clinical pathology

distilling the available knowledge on ethanol induced liver damage and directly complementing the available bio medical literature ethanol and the liver covers pathogenic and clinical aspects of alcoholic liver disease providing broader coverage of the subject than any available monograph the editors and their panel of experts relate basic scien

this updated and expanded fifth edition of clinical biochemistry of domestic animals brings together in a single comprehensive volume all the pertinent information regarding the biochemistry of disease and non disease states in animals clinical veterinarians and animal scientists now routinely use many of the same diagnostic and therapeutic tools used to identify and treat diseases and metabolic disorders in humans making this book an indispensable teaching learning and application resource for anyone engaged in the care health and welfare of animals this book concentrates on the various rationales and interpretations regarding the biochemistry of disease in animals it includes newly updated chapters with current references and new chapters on clinical toxicology and avian clinical biochemistry key features brings together in a single comprehensive volume all the pertinent information regarding the biochemistry of disease and non disease states in animals includes newly updated chapters with current references contains new chapters on clinical toxicology and avian clinical biochemistry concentrates on the various rationales and interpretations regarding the biochemistry of disease in animals

many aspects of drug safety have become an outstanding and even persistent issue and may occur during the process of both drug discovery and development until 15 years ago drug discovery and evaluation was primarily a sequential process starting with the selection of the most pharmacologically active compound from a series of newly synthesized small molecule chemical series by means of distinctive pharmacological assays safety aspects were addressed by evaluation of the selected compound at high doses in a series of specific studies directed at indications other than the intended indication of the new compound these tests are then followed by pharmacokinetic studies which are primarily conducted to confirm whether the selected compound possesses a suitable half life for sufficient exposure and efficacy and whether it has the desired properties specificity to the intended route of administration safety aspects relied predominantly on the conduct of single and repeat toxicologydose studies which inform changes in organ structure rather than organ function both toxicological and pharmacokinetic studies are

adapted to the progress of studies in clinical pharmacology and clinical trials the new edition of this well and broadly accepted reference work contains several innovative and distinguished chapters this sequential strategy has been abandoned with this new version of the book for several reasons of the possible multitude of negative effects that novel drugs may impart on organ function e g ventricular tachy arrhythmia many are detected too late in non clinical studies to inform clinicians on the other hand negative findings in chronic toxicity studies in animals may turn out to be irrelevant for human beings new scientific approaches e g high throughput screening human pluripotent stem cells transgenic animals knock out animals in silico models pharmaco genomics and pharmaco proteomics as well as artificial intelligence ai methods offered new possibilities there are several examples that show that the druggability of compounds was considerably underestimated when the probability of success of a new project was assessed the success rate in the pharmaceutical industry and the introduction of new chemical entities to the market per year dropped dramatically whereas the development time for a new compound increased sometimes exceeding the patent protection research and development scientists involving the following changes therefore adopted a change of strategy parallel instead of sequential involvement of the various disciplines multidimensional compound optimization the term safety pharmacology was coined the international conference on harmonization ich founded a safety pharmacology working group and the safety pharmacology society sps was launched the discipline provided for evaluation development and validation of a multitude of safety tests outlined in the core battery of studies characterizing the exposure profile of a drug by conducting pharmacokinetic studies that evaluates the absorption distribution metabolism and excretion should to be investigated at an early stage of development as results contribute to the selection of a compound for further development advancements in toxicology were achieved by the introduction of new methods e g in silico methods genetic toxicology computational toxicology and ai the book is a landmark in the continuously changing world of drug research and developments as such it is essential reading for many groups not only for all students of pharmacology and toxicology but also for industry scientists and physicians especially those involved in clinical trials of drugs and for pharmacists who must know the safety requirements of drugs the book is essential for scientists and managers in the pharmaceutical industry who are involved in drug discovery drug development and decision making in the development process in particular the book will be of use to government institutions and committees working on official guidelines for drug evaluation worldwide

the new edition of this successful reference offers both cutting edge and classic pharmacological methods thoroughly revised and expanded to

two volumes it offers an updated selection of the most frequently used assays for reliably detecting the pharmacological effects of potential drugs every chapter has been updated and numerous assays have been added each of the more than 1 000 assays comprises a detailed protocol outlining purpose and rationale and a critical assessment of the results and their pharmacological and clinical relevance

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