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justified unforgiveness a moral case against silence neglect and forgetfulness corrects the
 overemphasis on the benefits of forgiveness and shows how unforgiveness is a morally
 appropriate and at times morally necessary response towards unremorseful and unrepentant
 wrongdoers grave wrongs brought about by absolute evil and situations where justice is not
 served in advancing an argument for unforgiveness jennifer m s ang deals with three
 dimensions of forgiveness forgiveness as the letting go of resentment forgiveness as reparation of
 damaged moral relations with the community and forgiveness as letting the past remain in the
 past ang examines these arguments using cases of mass atrocities that are not often discussed in
 the forgiveness literature such as the nanking massacre the japanese military comfort women
 system and the khmer rouge genocide to reflect on the paradigm example of the holocaust as
 absolute evil these discussions also clarify the relationship between time forgiveness and justice
 how time can affect the willingness to forgive how the carrying out of justice over time
 remains vulnerable to politicization and how memory of mass atrocities enables victims to let go
 without forgiving

the idea of christ s substitutionary atonement for sinners is central in both the old and new

testaments from the passover to the prophets to the words of jesus and the apostles in it is well pastors mark dever and michael lawrence demonstrate how the atonement is clearly taught throughout scripture starting with exodus 12 and moving through other key old testament passages into the gospels and the epistles of paul and peter the authors offer careful expositions on fourteen crucial texts as they speak to important issues such as what happens when there is no substitute for sin why god forsook christ jesus perspective on his substitutionary work and the necessity and benefits of the atonement they show how much the doctrine applies to the christian life it is well not only encourages pastors to preach this essential doctrine for the strengthening of the church but it helps individual believers understand and exult in the richness of god s love in christ part of the 9marks series

this book is designed for clinical cardiologists and other physicians working with cardiac patients where specific specialized teams of cardio oncologists are not available and who are called to perform a clinical consultation to evaluate both the cardiac condition and the eligibility for chemotherapy or radiotherapy treatment and to evaluate if a cancer treatment produces toxic effects on a patient treated with chemo or radiotherapy and if appearance of new symptoms is due to this treatment in recent years progress in oncologic therapy has resulted in important developments and the prognostic improvement of patients with malignancy the cornerstone of chemotherapy are the anthracyclines and the analogue mitoxantrone that are direct cellular toxic agents and that are among the most powerful anti neoplastic drugs but their cardiac toxicity is well known significant breakthroughs in cancer therapy have also been achieved with the introduction of signalling inhibitors such as vegf inhibitors herb2 inhibitors or tk inhibitors used alone or in combination with direct cellular toxic drugs however these signalling inhibitors may interact also with cardiovascular signalling and therefore may have functional or structural effects on the myocardium this can be permanent or reversible with impairment of the global conditions and important side effects and increase both morbidity and mortality and worsening of quality of life chemotherapy and radiotherapy can have acute detrimental effects that can be present for years after treatment and that can lead to cardiac consequences also after long periods of no clinic sequelae patient numbers with cancer problems will dramatically increase in the next years and thus every cardiologist will need to have the correct information and the skills to manage these situations in the correct way this book will provide these tools for them

the purpose of this book is to give a concise introduction to development and analysis of pharmaceutical biologics for those in the pharmaceutical industry who are switching focus from small molecules to biologics processing analysis and delivery in order to maintain a limited focus introduction to biologic and biosimilar product development and analysis will deal only with

peptides proteins and monoclonal antibodies

investment in education technology has surged worldwide over the past decade and digital education technologies are now a key resource for oecd education and training systems if used effectively they promise to transform teaching and learning practices to reduce learning inequalities and to create more inclusive and efficient education systems

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translational medicine optimizing preclinical safety evaluation of biopharmaceuticals provides scientists responsible for the translation of novel biopharmaceuticals into clinical trials with a better understanding of how to navigate the obstacles that keep innovative medical research discoveries from becoming new therapies or even making it to clinical trials the book includes sections on protein based therapeutics modified proteins oligonucleotide based therapies monoclonal antibodies antibody drug conjugates gene and cell based therapies gene modified cell based therapies combination products and therapeutic vaccines best practices are defined for efficient discovery research to facilitate a science based efficient and predictive preclinical development program to ensure clinical efficacy and safety key features defines best practices for leveraging of discovery research to facilitate a development program includes general principles animal models biomarkers preclinical toxicology testing paradigms and practical applications discusses rare diseases discusses what why when how highlighting different considerations based upon product attributes includes special considerations for rare diseases about the editors joy a cavagnaro is an internationally recognized expert in preclinical development and regulatory strategy with an emphasis on genetic medicines her 40 year career spans academia government fda and the cro and biotech industries she was awarded the 2019 arnold j lehman award from the society of toxicology for introducing the concept of science based case by case approach to preclinical safety evaluation which became the foundation of ich s6 she currently serves on scientific advisory boards for advocacy groups and companies and consults and lectures in the area of preclinical development of novel therapies mary ellen cosenza is a regulatory toxicology consultant with over 30 years of senior leadership experience in the biopharmaceutical industry in the u s europe and emerging markets she has held leadership position in both the american college of toxicology act and the international union of toxicology iutox and is also an adjunct assistant professor at the university of southern california where she teaches graduate level courses in toxicology and regulation of biologics

every year 10 outstanding research topics are selected as finalists of the frontiers spotlight award

these shortlisted article collections each address a globally important field of research with the potential to drastically impact our future they bring together the latest cutting edge research to advance their fields present new solutions and foster essential large scale collaborations across multiple disciplines and research groups worldwide this international research prize recognizes the most innovative and impactful topics and the winning team of editors receives 100 000 to organize an international scientific conference on the theme of their successful collection

25 additional chapters available online the landmark text on how to achieve optimal patient outcomes through evidence based medication therapy pharmacotherapy a pathophysiologic approach is written to help you advance the quality of patient care through evidence based medication therapy derived from sound pharmacotherapeutic principles the scope of this trusted classic goes beyond drug indications and dosages to include the initial selection proper administration and monitoring of drugs you will find everything you need to provide safe effective drug therapy across the full range of therapeutic categories this edition is enriched by more than 300 expert contributors and every chapter has been revised and updated to reflect the latest evidence based information and recommendations important features such as key concepts at the beginning of each chapter clinical presentation tables that summarize disease signs and symptoms and clinical controversies boxes that examine the complicated issues faced by students and clinicians in providing drug therapy make this text an essential learning tool for students patient focused pharmacists and other health care providers new to this edition a section on personalized pharmacotherapy appears in most sections all diagnostic flow diagrams treatment algorithms dosing guideline recommendations and monitoring approaches have been updated in full color to clearly distinguish treatment pathways new drug monitoring tables have been added most of the disease oriented chapters have incorporated evidence based treatment guidelines that often include ratings of the level of evidence to support the key therapeutic approaches twenty five online only chapters are available at pharmacotherapyonline.com

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