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Astra Hatchback, Saloon & Estate, and Zafira MPV, inc. special/limited editions. Does NOT cover automatic transmission variants, or new range introduced May 2004 Diesel: 1.7 litre (1686 & 1700cc) & 2.0 litre (1995cc) turbo diesel. Does NOT cover 1.7 litre CDTi common rail engines.

An Ultra-Convenient, Safety-Focused Drug Guide Written for RNs by RNs Nursing Spectrum Drug Handbook delivers instant access to concise, need-to-know information on more than 3,000 brand-name and 1,000 generic drugs. Carefully updated and based on feedback from nurses worldwide, the Handbook is an essential tool in ensuring the safest, most effective drug administration possible. Features Organized alphabetically by generic name Covers each drug's mechanism of action, indication, dosage, contraindications, and more Up-to-date overview of ophthalmic drugs, drug classes, vitamins, minerals, herbs and supplements Color-coded " clinical alert " icons throughout Full-color patient safety and drug identification inserts Useful photo gallery of common tablets and capsules NEW treatment algorithms Vital coverage of life-threatening adverse reactions Valuable appendices Ribbon bookmark to keep your place Off-label uses for many drugs PDA download of the ENTIRE text Online companion site (www.nursedrughandbook.com) with drug updates, downloadable patient education information, and more

For decades researchers and programmers have used SAS to analyze, summarize, and report clinical trial data. Now Chris Holland and Jack Shostak have updated their popular Implementing CDISC Using SAS, the first comprehensive book on applying clinical research data and metadata to the Clinical Data Interchange Standards Consortium (CDISC) standards. Implementing CDISC Using SAS: An End-to-End Guide, Revised Second Edition, is an all-inclusive guide on how to implement and analyze the Study Data Tabulation Model (SDTM) and the Analysis Data Model (ADaM) data and prepare clinical trial data for regulatory submission. Updated to reflect the 2017 FDA mandate for adherence to CDISC standards, this new edition covers creating and using metadata, developing conversion specifications, implementing and validating SDTM and ADaM data, determining solutions for legacy data conversions, and preparing data for regulatory submission. The book covers products such as Base SAS, SAS Clinical Data Integration, and the SAS Clinical Standards Toolkit, as well as JMP Clinical. Topics included in this edition include an implementation of the Define-XML 2.0 standard, new SDTM domains, validation with Pinnacle 21 software, event narratives in JMP Clinical, SDTM and ADAM metadata spreadsheets, and of course new versions of SAS and JMP software. The second edition was revised to add the latest C-Codes from the most recent release as well as update the make_define macro that accompanies this book in order to add the capability to handle C-Codes. The metadata spreadsheets were updated accordingly. Any manager or user of clinical trial data in this day and age is likely to benefit from knowing how to either put data into a CDISC standard or analyzing and finding data once it is in a CDISC format. If you are one such person—a data manager, clinical and/or statistical programmer, biostatistician, or even a clinician—then this book is for you.

Present day insights into the relationship between gender and health emphasise the necessity of taking sex and gender differences into consideration. Sex refers to biological characteristics such as chromosomes, physiology and anatomy that distinguish females and males. Gender refers to the array of socially constructed roles and relationships, personality traits, attitudes, behaviours and values that society ascribes to the two sexes on a differential basis. In health-related research both sex differences and differences resulting from gender are implied. Sex can determine differential propen.

A monumental work. This surely has the potential of becoming the sound designer's bible!"---Kees van den Doel, Scientific Computing Laboratory, University of British Columbia--

In this book, world-renowned experts in the field express well-reasoned opinions on a range of issues and controversies relating to haploidentical transplantation with the aim of providing practicing hematologists with clinically relevant and readily applicable information. Among the areas covered are graft manipulation and methods to control T-cell alloreactivity, the nature of the ideal graft and donor, haploidentical transplantation in pediatric and adult patients with malignant and nonmalignant diseases, immunologic reconstitution following transplantation, complications, and the prevention and treatment of relapse post transplantation. Attention is drawn to the implications of high-impact clinical trials whenever such trials are available. The readily intelligible text is complemented by numerous helpful tables, algorithms, and figures. The book will provide practical support for hematologists and transplant physicians as they attempt to provide optimal care in this exciting but increasingly complex medical specialty.

Antibody–drug conjugates (ADCs) represent one of the most promising and exciting areas of anticancer drug discovery. Five ADCs are now approved in the US and EU [i.e., ado-trastuzumab emtansine (Kadcyla™), brentuximab vedotin (Adcetris™), inotuzumab ozogamicin (Besponsa™), gemtuzumab ozogamicin (Mylotarg™) and moxetumomab pasudotox-tdfk (Lumoxiti®)] and over 70 others are in various stages of clinical development, with impressive interim results being reported for many. The technology is based on the concept of delivering a cytotoxic payload selectively to cancer cells by attaching it to an antibody targeted to antigens on the cell surfaces. This approach has several advantages including the ability to select patients as likely responders based on the presence of antigen on the surface of their cancer cells and a wider therapeutic index, given that ADC targeting enables a more efficient delivery of cytotoxic agents to cancer cells than can be achieved by conventional chemotherapy, thus minimising systemic toxicity. Although there are many examples of antibodies that have been developed for this purpose, along with numerous linker technologies used to attach the cytotoxic agent to the antibody, there is presently a relatively small number of payload molecules in clinical use. The purpose of this book is to describe the variety of payloads used to date, along with a discussion of their advantages and disadvantages and to provide information on novel payloads at the research stage that may be used clinically in the future.

Melanoma is one of the most types of cancer. When melanoma is detected at an early stage, treatment is highly successful, but outcomes can be poor when the disease is advanced. There has been significant progress in our understanding of the molecular biology, genetics, and immunology of melanoma over the past decade. This has been accompanied by rapid advances in therapeutic strategies for patients with melanoma. This book provides the clinician and the researcher with a broad understanding of the molecular and cellular pathogenesis of melanoma, explores the clinical characteristics and criteria for clinical and pathological staging of the disease, and provides an overview of current and evolving treatment strategies in the adjuvant, metastatic, and preventive settings. The treatment of special populations and rare variants of melanoma that often present particular clinical challenges is also covered. Authored by international experts in melanoma biology and clinical management, this volume concisely explains how to diagnose, treat, and prevent melanoma while reviewing advances in basic science and providing an overview of innovative approaches still under development.

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