

CLINICAL Data

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handbook of Clinical Drug Data

tenth edition

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handbook of Clinical Drug Data

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Assistant Professor, School of Medicine, Section of Clinical Pharmacology, University of Missouri-Kansas City; Adjunct Assistant Professor, School of Pharmacy, University of Missouri-Kansas City, Kansas City, Missouri The Tenth Edition of the *Handbook of Clinical Drug Data* continues a long tradition of providing clinically relevant, well-referenced drug information compiled by expert clinicians and presented in a compact format. The formats of all sections should be familiar to users of the ninth edition. As with recent editions, information in the *Handbook* is divided into three parts.

Drug Monographs in Part I have been updated to include numerous newly marketed and promising investigational drugs. Areas with extensive revisions include the Antivirals reflecting the many new agents for HIV infection, Immunosuppressants, Anticonvulsants, and the Hematologic Drugs. Three new subsections have been added to reflect the growing number of agents for rheumatoid arthritis, glaucoma and osteoporosis: Antiarthritic Drugs in the Analgesic and Anti-inflammatory Drugs section, Ophthalmic Drugs for Glaucoma in the Central Nervous System section, and Bisphosphonates in the Renal and Electrolytes section.

Clinical Drug Information in Part II continues to provide clinically useful information that helps the reader to decide which drug(s) are most likely to have caused adverse reactions or which are the best choices for patients in special populations. All drug-induced diseases sections have been extensively updated, as have the Cytochrome P450 Interactions, Pregnancy, Breastfeeding, Renal Disease, Immunization, and Cardiac Arrest sections. Drug-Laboratory Test Interferences in Part III has also been updated.

In this edition, we welcome several new authors: Dan Baker, Jess Benson, Toy Biederman, Juliana Chan, Paul Cuddy, Rob DiDomenico, Allison Einhorn, Ray Hammond, Patty Marshik, Gary Matzke (a returning author), Renée Mercier, and Anna Taddio (our first "international" author). We would also like to thank the previous authors whose work in most cases served as the basis for revisions of the chapters that appear in this edition by new authors: Andrea Anderson (Drugs and Pregnancy), Lisa Ashton (Respiratory Drugs), Arasb Ateshkadi (Renal and Electrolytes). Rosemary Berardi (Gastrointestinal Drugs), Larry Borgsdorf (anaphylaxis) Larry Davis (NSAIDs and Hematologic Drugs), John Flaherty (Aminoglycosides and β-Lactams), John Gambertoglio (Renal Disease), Millie Gottwald (Antimigraine Drugs and Neurodegenerative Diseases), Amy Guenette (Inotropic Drugs and Nitrates), Brian Kearney (Renal Disease), and Carolyn Zaleon (Gastrointestinal Drugs). We are saddened to report the deaths of Drs. Ateshkadi and Gambertoglio since our last edition. Both will be remembered for their professional dedication and the quality of their work. John Gambertoglio had long-time personal and professional ties to the editors and will be particularly missed.

This edition also marks another major change, being the first edition produced with our new publisher, McGraw-Hill and new editors, Stephen Zollo and

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Nicky Panton. We thank them for their efforts to maintain the high quality of the *Handbook* that we desire and our readers have come to expect.

Philip O. Anderson James E. Knoben William G. Troutman August 2001

How to Use This Book

Part I of this book is organized around 10 major drug categories, which have been subdivided into common therapeutic groups. Within these therapeutic groups, drug information is alphabetically presented in three formats: Monographs, Minimonographs, and Comparison Charts. Monographs and Comparison Charts are grouped together to ensure that related drugs are easy to compare and contrast. Charts are located after the monographs to which they relate. Drug antagonists are grouped together with agonists to simplify organization and accessibility.

Monographs are used for drugs of major importance and prototype agents.

Minimonographs are used for drugs similar to prototype drugs, those of lesser importance within a therapeutic class, and promising investigational agents. Minimonographs contain only selected subheadings of information rather than all subheadings contained in the full monographs.

Comparison Charts are used to present clinically useful information on members of the same pharmacologic class and different drugs with a similar therapeutic use, as well as to present clinically relevant information on certain other topics.

The preferred method to gain access to complete information on a *particular brand* or *generic drug* is to use the index at the end of the book. The index may also direct the user to *other pertinent information* on the drug.

MONOGRAPH FORMAT

CLASS INSTRUCTIONS

This is an optional heading at the beginning of each drug class. It consists of patient instructions that apply to more than one of the drug monographs in this subcategory. If all drugs are not identical in their instructions, only the common information is found here. The Patient Instructions section of each monograph that is affected states, "See Class Instructions" as the opening phrase.

GENERIC DRUG NAME

Brand Name(s)

The *nonproprietary (generic)* name is listed on the left, followed by common brand names listed on the right. Brand-name products listed are not necessarily superior or preferable to other brand-name or generic products; "Various" indicates the availability of additional brand and/or generic products.

Pharmacology. A description of the chemistry, major mechanisms of action, and human pharmacology of the drug in clinical application.

Administration and Adult Dosage. Route of administration, indications, and usual adult dosage range are given for the most common labeled uses. Dosages correspond

to those in the product labeling or in standard reference sources. "Dose" refers to a single administration and "dosage" to a cumulative amount (eg, daily dosage).

Special Populations. Dosages in patient populations other than the typical adult are listed:

Pediatric Dosage (given by age or weight range)

Geriatric Dosage (given by age range)

Other Conditions (renal failure, hepatic disease, obesity, etc.)

Dosage Forms. The most commonly used dosage forms and available strengths are listed, as well as popular combination product dosage forms. Prediluted IV piggyback or large-volume parenteral containers are not listed unless this is the only commercially available product.

Patient Instructions. Key information that should be provided to the patient when prescribing or dispensing medication is presented. When introductions apply to an entire drug category, see "Class Instructions" at the beginning of that subcategory.

Missed Doses. What the patient should do if one or more doses are missed.

Pharmacokinetics. Data are presented as the mean \pm the standard deviation. Occasionally the standard error of the mean (SE) is the only information available on variability, and it is identified as such.

Onset and Duration (time course of the pharmacologic or therapeutic effect)

Serum Levels (therapeutic and toxic plasma concentrations are given)

Fate (The course of the drug in the body is traced. Pharmacokinetic parameters are generally provided as total body weight normalized values. The volume of distribution is either a V_d in a one-compartment system or V_c and $V_{d\beta}$ or V_{dss} in a two-compartment system.)

t½ (terminal half-life is presented)

Adverse Reactions. Reactions known to be dose related are usually given first, then other reactions in decreasing order of frequency. Reaction frequency is classified into three ranges. However, percentages of reactions may be provided for reactions that occur more frequently than 1%.

frequent (>1/100 patients) occasional (1/100 to 1/10,000 patients)

rare (<1/10,000 patients)

Contraindications. Those listed in product labeling are given. "Hypersensitivity" is not listed as a contraindication because it is understood that patients should usually not be given a drug to which they are allergic or hypersensitive—exceptions are noted.

Precautions. Warnings for use of the drug in certain disease states and/or patient populations, together with any cross-sensitivity with other drugs. Part II, Chapter 3, "Drug Use in Special Populations," should be consulted for more information, particularly regarding pregnancy and breastfeeding.

Drug Interactions. The most important drug interactions are listed.

Parameters to Monitor. Important clinical signs and/or laboratory tests to monitor to ensure safe and effective use are presented. The frequency of monitoring may also be given; however, for many drugs the optimal frequency has not been determined.

Notes. Distinguishing characteristics, therapeutic usefulness, or relative efficacy of the drug are presented, as well as unique or noteworthy physicochemical properties, handling, storage, or relative cost.

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