

N.F. Muller • R.P. Dessing (Eds.)

# European Drug Index

Fourth  
Edition



European Society of Clinical Pharmacy



CRC Press  
Taylor & Francis Group

# European Drug Index

**National Library of Medicine (U.S.)**



## **European Drug Index:**

*European Drug Index* Niels F. Muller, Rudolf P. Dessing, 2022-03-07 Many health care providers are frequently dealing with problems related to the identification and interpretation of medicines and prescriptions of foreign origin Health authorities customs and travel agencies also encounter such problems which are related to the increasing mobility of the European population Thus the need for a European Drug Index is obvious The EDI provides extended information for practitioners confronted with the enormous number of drug names available on the European pharmaceutical market This market is increasing due to the rapidly changing palette of countries and economic restrictions in Europe The listings have been derived from drug data sources from the increased number of participating countries in this second edition Each item starts with a trade name in alphabetical order followed by depending on the original source dosage forms strength volume if applicable and generic name s of the active principle s in a random sequence The item is concluded by the Anatomical Therapeutic Chemical ATC classification when made available by the original source and a code for the country of origin

**European Drug Index** Niels F. Muller, Rudolf P. Dessing, 1997      **Drug Information** Bonnie Snow, 1999-11-17  
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Contemporary Drug Information Claude J. Gaebelein, Brenda L. Gleason, 2008 This innovative textbook teaches the basics of drug information literature evaluation and biostatistics and relates these topics to evidence based pharmaceutical care Readers will learn what to look for in studies how to critique them and how to apply them in clinical pharmacy practice A major focus is critical appraisal of evidence derived from different types of studies cases cohorts surveys randomized controlled clinical trials pharmacoeconomic studies and systematic reviews Concluding chapters discuss clinical decision making using evidence from studies      *European Drug Index* ,1992      **European drug index** ,1992      **WHO Drug**

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also highlighting the social impact which drugs have in modern societies Taking an evidence based approach to the problem the practice of clinical pharmacology and pharmacotherapy in the developing as well as the developed world is examined For this purpose the book covers general clinical pharmacology pharmacology of various drug groups and the treatments specific to various diseases the book gives guidance on how doctors should act so that drugs can be used effectively and safely and it encourages the rational use of drugs in society This publication brings together a large amount of excellent content that will be invaluable for anyone working within or associated with the field of clinical pharmacology and pharmacotherapy undergraduates postgraduates regulatory authorities and the pharmaceutical industry

**WHO Drug Information**, 2021-04-08 *Index of NLM Serial Titles* National Library of Medicine (U.S.), 1984 A keyword listing of serial titles currently received by the National Library of Medicine Information Resources in Toxicology P.J. Bert Hakkinen, Asish Mohapatra, Steven G. G. Gilbert, 2009-08-19 This latest version of Information Resources in Toxicology IRT continues a tradition established in 1982 with the publication of the first edition in presenting an extensive itemization review and commentary on the information infrastructure of the field This book is a unique wide ranging international annotated bibliography and compendium of major resources in toxicology and allied fields such as environmental and occupational health chemical safety and risk assessment Thoroughly updated the current edition analyzes technological changes and is rife with online tools and links to Web sites IRT IV is highly structured providing easy access to its information Among the hot topics covered are Disaster Preparedness and Management Nanotechnology Omics the Precautionary Principle Risk Assessment and Biological Chemical and Radioactive Terrorism and Warfare are among the designated International in scope with contributions from over 30 countries Numerous key references and relevant Web links Concise narratives about toxicologic sub disciplines Valuable appendices such as the IUPAC Glossary of Terms in Toxicology Authored by experts in their respective sub disciplines within toxicology Impact of written drug information in patient package inserts. Acceptance and benefit/risk perception Robert H. Vander Stichele, 2004 Drug Information Patrick M. Malone, Karen L. Kier, John Stanovich, 2010-05-12 Extensive coverage of the Internet as a source of and distribution means for drug information and detailed sections on evaluating medical literature from clinical trials Audience includes Pharmacists Pharmacy students and Pharmacy schools Updated to include using PDAs for medication information Covers the ethical and legal aspects of drug information management Nothing else like it on the market **Current Catalog** National Library of Medicine (U.S.), 1993 First multi year cumulation covers six years 1965 70 *National Library of Medicine Current Catalog* National Library of Medicine (U.S.), 1988 **Index Medicus**, 2001-06 Vols for 1963 include as pt 2 of the Jan issue Medical subject headings **The Challenges of the New EU Pharmaceutical Legislation** José Luis Valverde, Paul Weissenberg, 2005 In the domain of public policy on pharmaceuticals protecting public health requires a dual strategy robust regulation on the one hand and stimulation of competitiveness and innovation on the other Regulation must be robust to

ensure that only medicines meeting exact standards of safety quality and efficacy are authorised for human and animal use At the same time competitiveness and innovation must be stimulated Without innovation in pharmaceuticals the incurable diseases of today will remain incurable Competitiveness drives innovation and innovation saves lives Increased competitiveness of the pharmaceutical sector will not only better protect public health but will also create high quality jobs and create growth In this context the implementation of the G10 recommendations particularly regarding the pricing and reimbursement of medicines by Member States remains a considerable challenge

**Dictionary of Pharmaceutical Medicine** Gerhard Nahler, 2017-03-17 This dictionary defines various terms typically used in pharmaceutical medicine A new 4th edition includes adaptations of the text to the steadily increasing regulatory requirements particularly in the area of genetics gene therapy product quality e g protection against falsified medicines and of product safety pharmacovigilance Further evolving areas that are covered by the 4th edition are typical grey zones health effects often borderline to medicinal products such as cosmetics and dietary supplements where misleading information is prohibited on one hand but where any health claims need formal authorisation on the other These but also other areas are reviewed and presented in an updated and if justified in an enlarged form

**Detection and Prevention of Adverse Drug Events** Régis Beuscart, Werner Hackl, Christian Nøhr, 2009 Presents papers from the workshop Patient Safety through Intelligent Procedures in medication focusing on identification of ADE and medication errors in hospital settings the role of human and organizational factors on ADE and medication errors and information and communication technologies to prevent or correct ADEs and medication errors

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