

1.	Record Nr.	UNIBAS000006026
	Autore	Van Dyke, Henry <1852-1933>
	Titolo	The lost boy / by Henry Van Dike
	Pubbl/distr/stampa	New York ; London : Harper & Brothers, 1914
	Descrizione fisica	68 p., [3] c. di tav. ; 18 cm.
	Disciplina	813.52
	Lingua di pubblicazione	Inglese
	Formato	Materiale a stampa
	Livello bibliografico	Monografia
2.	Record Nr.	UNINA9910830124003321
	Titolo	Stephens' detection of new adverse drug reactions / / edited by John Talbot, Patrick Waller
	Pubbl/distr/stampa	Chichester ; ; Hoboken, NJ, : Wiley, c2004
	ISBN	1-280-26932-4 9786610269327 0-470-09265-3 0-470-01419-9
	Edizione	[5th ed.]
	Descrizione fisica	1 online resource (763 p.)
	Altri autori (Persone)	StephensM. D. B. <1930-> TalbotJ. C. C WallerPatrick
	Disciplina	615.7042 615/.7042
	Soggetti	Drugs - Side effects Drugs - Toxicology
	Lingua di pubblicazione	Inglese
	Formato	Materiale a stampa
	Livello bibliografico	Monografia
	Note generali	Rev. ed. of: Detection of new adverse drug reactions / edited by M.D.B. Stephens, J.C.C. Talbot, and P.A. Routledge. 4th ed. 1998.
	Nota di bibliografia	Includes bibliographical references and index.

Stephens' Detection of New Adverse Drug Reactions Fifth Edition;
Contents; Foreword; Preface; List of Contributors; 1 Introduction;
Mercury; Introduction to pharmacovigilance; The history of
pharmacovigilance; Under-reporting of adverse drug reactions;
Incidence of adverse drug reactions; The financial cost of adverse drug
reactions; Preventability of adverse drug reactions; Risk-benefit ratio;
The changing risks with new drugs; Expression of risk; The risks we are
prepared to take; Definitions; Classification of adverse drug reactions;
Adverse reaction profile
Adverse events in a patient's life Symptoms in healthy persons; Adverse
reactions to placebo; Infectiousness of adverse drug reactions; Herbal
medicines; Some final food for thought; References; Further reading; 2
Adverse Drug Reactions and Interactions: Mechanisms, Risk Factors,
Detection, Management and Prevention; Introduction; Classification of
adverse drug reactions; Risk factors for type A adverse reactions; Risk
factors for type B adverse reactions; Detection of adverse drug
reactions; Management of adverse drug reactions; Prevention of
adverse drug reactions; Drug interactions
Conclusions References; 3 Toxicology and Adverse Drug Reactions;
Introduction; Toxicity testing; Drug development; Data interpretation
and risk assessment; Adverse drug reactions detected after
authorization; Examples of toxicological investigation of adverse drug
reactions; Conclusions; Acknowledgements; References; 4 Clinical
Trials: Collection of Safety Data and Establishing the Adverse Drug
Reaction Profile; Introduction; Adverse events; Final analysis of data;
Inadequate reporting of safety data from clinical trials; Conclusions;
Future aspirations; Acknowledgements; References
5 Clinical Laboratory Safety Data in Drug Studies Introduction; Factors
that influence interpretation of clinical laboratory data; Sample
collection procedure; Analytical variation; Reference ranges; Intra-
individual biological variation; Safety testing in drug development; Test
selection; Exclusion criteria and 'panic levels'; Harmonization of data
from different laboratories; Data analysis and presentation; Conclusion;
References; 6 Statistics: Analysis and Presentation of Safety Data;
Introduction and background; Analysis and presentation of data from
trials
Measures that take time into account Statistical tests utilizing time since
start of treatment; Combining data from several trials: meta-analysis;
Analysis and presentation of data from observational studies; Use of
statistical methods for signal detection with spontaneous reports;
Summary and conclusions; Acknowledgements; References; 7 Causality
and Correlation in Pharmacovigilance; Introduction and historical
background; The notions of necessary and sufficient causes; Factors to
be considered in causality assessment; Methods for causality
assessment; When to assess causality
Assessing causality from multiple information sources: the Bradford-
Hill criteria

Sommario/riassunto

A key text for all those involved in pharmacovigilance. Detection of new
adverse drug reactions is fundamental to the protection of patients
from harm that may occur as a result of medication. This book explores
the methods used to investigate new adverse drug reactions,
discussing all elements from the scientific background and animal
toxicology through to worldwide regulatory and ethical issues.
Stephens' Detection of New Adverse Drug Reactions provides
comprehensive and up-to-date coverage of material fundamentally
important to all those active in the field, whether they w
