

|                         |  |
|-------------------------|--|
| 1. Record Nr.           | UNINA9910450500603321  |
| Titolo                  | The textbook of pharmaceutical medicine [[electronic resource]]  |
| Pubbl/distr/stampa      | London, : BMJ Books, 2002  |
| ISBN                    | 1-280-19787-0<br>9786610197873<br>1-4051-4606-0<br>1-4237-1777-5   |
| Edizione                | [4th ed. /]  |
| Descrizione fisica      | 1 online resource (896 p.)   |
| Altri autori (Persone)  | GriffinJ. P (John Parry)<br>O'GradyJohn  |
| Disciplina              | 615.1  |
| Soggetti                | Pharmaceutical chemistry<br>Pharmacology<br>Electronic books.  |
| Lingua di pubblicazione | Inglese  |
| Formato                 | Materiale a stampa   |
| Livello bibliografico   | Monografia   |
| Note generali           | Includes index.<br>Previous ed.: / edited by J.P. Griffin, J. O'Grady, P.F. D'Arcy. Belfast : Queen's University of Belfast, c1998.  |
| Nota di contenuto       | Contents; Contributors; Preface; Acknowledgements; The editors; 1: Discovery of new medicines; 2: Pharmaceutical development; 3: Toxicity testing; 4: Exploratory development; 5: Clinical pharmacokinetics; 6: Clinical trials and good clinical practice; 7: Medical statistics; 8: Development of medicines: full development; 9: The medical department; 10: Medical marketing; 11: Information and promotion; 12: The supply of unlicensed medicines for particular patient use; 13: Legal and ethical issues relating to medicinal products; 14: The safety of medicines<br>15: The development of the control of human medicines in Europe from classical times to the year 200016: Technical requirements for registration of pharmaceuticals for human use: the ICH process and the common technical document; 17: The regulation of drug products by the United States Food and Drug Administration; 18: The US FDA in the drug development, evaluation and approval process; 19: Regulatory and clinical trial systems in Japan; 20: Economics of healthcare; 21: |

Controls on NHS medicines prescribing and expenditure in the UK (a historical perspective); Appendix 1; Appendix 2  
PRESCRIPTION MEDICINES CODE OF PRACTICE AUTHORITY Constitution and ProcedureIndex

Sommario/riassunto

This edition has totally revised chapters on drug regulation in the USA, with new contributor, Dr Peter Barton Hutt formerly FDA Special Counsel. New contributions regarding pharmaco-economics are covered in two new chapters.