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Titolo	Pharmacovigilance : Critique and Ways Forward // edited by I. Ralph Edwards, Marie Lindquist
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Edizione	[1st ed. 2017.]
Descrizione fisica	1 online resource (XXXI, 205 p. 16 illus., 3 illus. in color.)
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Soggetti	Pharmacy Drug Safety and Pharmacovigilance
Lingua di pubblicazione	Inglese
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Livello bibliografico	Monografia
Nota di bibliografia	Includes bibliographical references at the end of each chapters.
Nota di contenuto	1. How to learn Pharmacovigilance, now and in the Future -- 2. Lost in Regulation -- 3. What is the Future of Pharmacovigilance and how can we Make it as Good as Possible? -- 4. Some Other Ideas About the Future of Pharmacovigilance -- 5. Best Safety Practices now and in the Future -- 6. Is There a Benefit from the Medical Literature Monitoring Service of the EMA? -- 7. The Unified Drug Safety-Clinical Database -- 8. Behind the Scenes – 'Silent Factors' Influencing Pharmacovigilance Practice and Decisions -- 9. Shaking up Culture and Communication in PV [V6] -- 10. Broadening the Scope of Pharmacovigilance -- 11. Effective Treatment Matters: Revitalizing Pharmacovigilance -- 12. There are More Weber-Effects -- 13. Herbal and Traditional Medicines, Now and Future -- 14. The Concept of 'Health' -- 15. Impact of Referral Procedures on the Interaction of RMP and PSUR -- 16. Other Sources of Information for Monitoring Drug Safety: Now and in the Future -- 17. Ecopharmacovigilance.
Sommario/riassunto	Written by experienced authors, this book offers expert personal views on what the current problems in pharmacovigilance are and how they should be solved. This book stems from thoughts and ideas discussed in a series of meetings of the International Society of Pharmacovigilance (ISoP), where concerns were raised that the current pharmacovigilance system is not delivering optimally to improve therapeutics in clinical practice. Pharmacovigilance of the future must be an active and integral part of health care delivery, and focus more on science and practices

that support health professionals and patients in day-to-day care situations. To achieve this, a dynamic and sustainable development of vigilance must take precedence over the current excessive preoccupations with data processing and regulations; all aspects of medicines use and their effects need to be considered; and all stakeholders must be involved and engaged in an open and constructive debate. The work is essential reading for anyone who has an interest in safer use of medicines. It is intended to be equally challenging and rewarding, and sets out to stimulate a continuous debate on how pharmacovigilance can better meet the needs of health professionals and patients to achieve the aim of wise therapeutic decision making.
