1. Record Nr. UNINA9910878043903321

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Titolo	China Food and Drug Law : Policy, Practice, and Future Directions / / by John C. Balzano
Pubbl/distr/stampa	Cham : , : Springer International Publishing : , : Imprint : Springer, , 2024
ISBN	9783031619014 9783031619007
Edizione	[1st ed. 2024.]
Descrizione fisica	1 online resource (399 pages)
Collana	Law for Professionals, , 2662-1428
Disciplina	340.9
Soggetti	Private international law Conflict of laws International law Comparative law Administrative law Food - Safety measures Asia - Politics and government Private International Law, International and Foreign Law, Comparative Law Administrative Law Food Safety Asian Politics
Lingua di pubblicazione	Inglese
Formato	Materiale a stampa
Livello bibliografico	Monografia
Nota di contenuto	1. Introduction and Methology -- 2. Agencies involved in China FDD Regulation -- 3. General Administrative Law Concepts and Enforcement Powers -- 4. Common Principles of China FDD Regulation -- 5. Pharmaceutical Regulation -- 6. Medical Device Regulation -- 7. Biosecurity and FDD Research and Development -- 8. Food and Special Food Regulation -- 9. Cosmetics Regulation -- 10. Advertising and Promotion for Regulated Products.
Sommario/riassunto	This book is an analysis of policy and law governing the development, approval, manufacture, distribution, marketing and post-market

surveillance of human drugs, medical devices, foods and cosmetics in Mainland China (“China FDD Regulation”). It analyzes the policy and general principles behind China FDD Regulation, including the history and jurisdiction of the central and local agencies that administer the laws and regulations, the administrative law structure in which these agencies operate, and other aspects of FDD Regulation interpretation and enforcement. Although it describes practice in China, this book is written in the comparative perspective (i.e., sensitive to assumptions made by those who are steeped in FDD Regulation in the U.S., Japan, and the EU). It includes one chapter on each regulatory space (drugs, medical devices, cosmetics, and food) organized by the lifecycle of products. It also covers subcategories of products, such as vaccines, radiopharmaceuticals, and in vitro diagnostic medical devices. The book includes specialty chapters on areas that are common to multiple types of regulated products, such as biosecurity and advertising. It is the first English language book of its kind, and it can serve as a resource for those in the FDD law and regulatory field to understand the mechanics of developing and marketing products, but also with insights for businesspeople and others, who are developing China-strategies.