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# CENTURY

THE CHALLENGES OF REGULATING  
DRUGS AND NEW TECHNOLOGIES

EDITED BY  
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# FDA in the Twenty-First Century: The Challenges of Regulating Drugs and New Technologies



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In its decades-long effort to assure the safety, efficacy, and protection of medicines and other items, the meals and Drug Administration has struggled with issues of funding, correct associations with industry, and the balance between consumer choice and consumer protection. Dealing honestly and thoroughly with the FDA's successes and failures, these essays rethink the structure, function, and future of the company and the effect policy innovations may possess on regulatory institutions abroad. With essays by leading scholars and authorities and private-industry professionals, FDA in the Twenty-First Hundred years addresses perennial and brand-new problems and the improvements the agency could make to better provide the public good. The book also intervenes in the debate over off-label medication marketing and the correct function of the FDA before and after a medication goes on the market. The collection features essays on effective regulation in an era of globalization, consumer empowerment, and comparative effectiveness, and also questions of data transparency, conflicts of interest, industry responsibility, and innovation policy, all with an focus on pharmaceuticals. Today, these problems are compounded by the pressures of globalization, the intro of novel technology, and fast-evolving threats to community health.



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