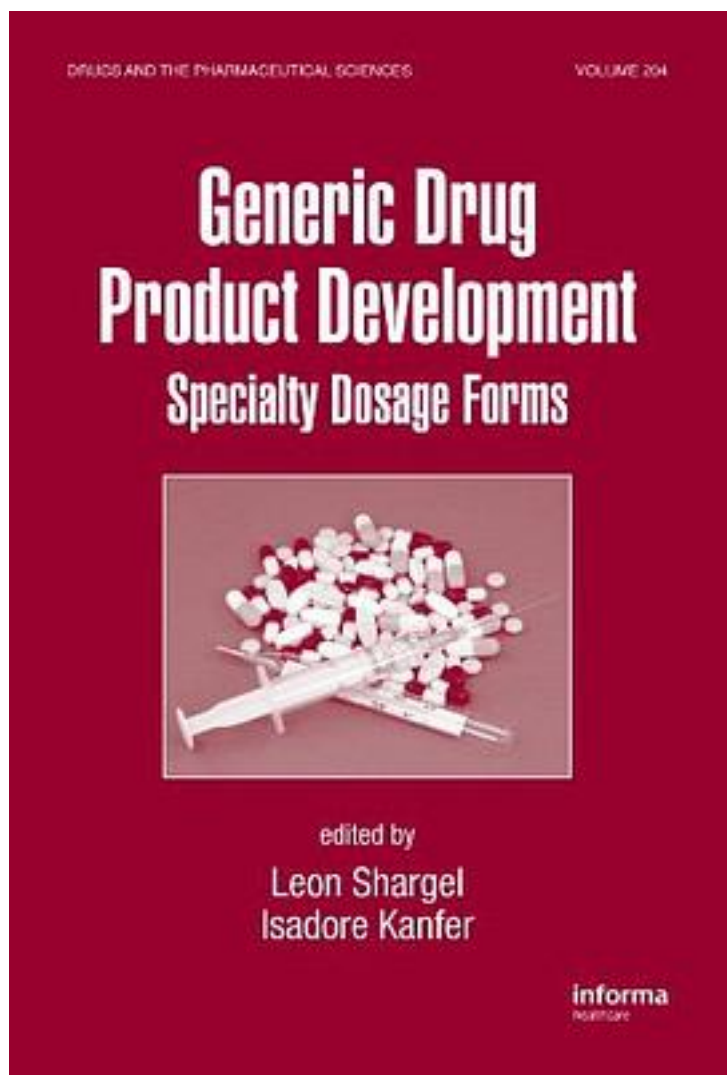


Generic Drug Product Development



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著者:Shargel, Leon/ Kanfer, Isadore

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Generic Drug Product Development: Specialty Dosage Forms explores the issues in providing evidence of pharmaceutical equivalence and bioequivalence for specialty drug products. In addition, the book provides various scientific approaches and regulatory requirements for manufacturers to demonstrate the therapeutic equivalence of generic specialty drug products to brand name alternatives. This book discusses measurement of drug product quality and performance, as well as the regulatory and scientific requirements of topical, nasal and inhalation, generic biologics, and other drug products. The book will be of interest of specialists and researchers in pharmaceutical drug development, regulation, manufacturing, and others in the pharmaceutical sciences.

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