J. T. Carstensen, Ph.D.

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# **Pharmaceutical Principles Of Solid Dosage Forms**

Jens T. Carstensen

#### **Pharmaceutical Principles Of Solid Dosage Forms:**

**Pharmaceutical Principles of Solid Dosage Forms** Jens T. Carstensen, 1993-08-16 This comprehensive guide presents the principles underlying the formation properties and manufacturing of solid dosage forms Developed by a leading pharmaceutical expert with 25 years of experience this book is a precise handbook to powder dosage forms hard shell capsules milling tablet granulation and drying stability crystalline solids DSC and amorphates hygroscopicity packaging micromeritics sustained release and more Designed for use by development pharmacists analytical chemists production pharmacists and pharmacy professors and students this book emphasizes U S validation principles the interrelationship between formulation manufacturing and existing USP and FDA guidelines and aspects of pharmaceutics important to IND and NDA submission and subsequent supplements **Developing Solid Oral Dosage Forms** Yihong Qiu, Yisheng Chen, Geoff G.Z. Zhang, Lirong Liu, William Porter, 2009-03-10 Developing Solid Oral Dosage Forms is intended for pharmaceutical professionals engaged in research and development of oral dosage forms It covers essential principles of physical pharmacy biopharmaceutics and industrial pharmacy as well as various aspects of state of the art techniques and approaches in pharmaceutical sciences and technologies along with examples and or case studies in product development The objective of this book is to offer updated or current knowledge and skills required for rational oral product design and development The specific goals are to provide readers with Basics of modern theories of physical pharmacy biopharmaceutics and industrial pharmacy and their applications throughout the entire process of research and development of oral dosage forms Tools and approaches of preformulation investigation formulation process design characterization and scale up in pharmaceutical sciences and technologies New developments challenges trends opportunities intellectual property issues and regulations in solid product development The first book ever that provides comprehensive and in depth coverage of what s required for developing high quality pharmaceutical products to meet international standards It covers a broad scope of topics that encompass the entire spectrum of solid dosage form development for the global market including the most updated science and technologies practice applications regulation intellectual property protection and new development trends with case studies in every chapter A strong team of more than 50 well established authors co authors of diverse background knowledge skills and experience from industry academia and regulatory agencies Instrumentation and Data Analysis for Solid Dosage R&D (Seminar Notes - Apr. 1993) C R C Press LLC, Carstensen, 1992-01-01 Handbook of **Stability Testing in Pharmaceutical Development** Kim Huynh-Ba,2008-11-16 This handbook is the first to cover all aspects of stability testing in pharmaceutical development Written by a group of international experts the book presents a scientific understanding of regulations and balances methodologies and best practices Handbook of Pharmaceutical Granulation Technology Dilip M. Parikh, 2016-04-19 The Third Edition presents all pharmaceutical industry personnel and those in academia with critical updates on the recent advances in granulation technology and changes in FDA regulatory

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