

## Ispe Baseline Pharmaceutical Engineering Guide Volume 5

Pharmaceutical manufacturing can be viewed as a supply chain which spans from the production and purchase of the starting and packaging materials through the manufacture of dosage forms until the safe reception of the finished product by the patient. The entire chain comprises of several processes: auditing, materials purchase (procurement), production, storage, distribution, quality control, and quality assurance. The quality standard for pharmaceutical production is 'current good manufacturing practice (CGMP)', which is applied within the frame of a pharmaceutical quality system (PQS). This implementation, however, requires a scientific approach and has to take into account several elements such as risk assessment, life cycle, patient protection, among other factors. Hence, pharmaceutical manufacturing is a complex subject in terms of regulation, given the technical and managerial requirements. This comprehensive handbook describes CGMP for new professionals who want to understand and apply the elements which build up pharmaceutical quality assurance. The book gives details about basic quality control requirements (such as risk management, quality hazards and management systems, documentation, clean environments, personnel training) and gives guidelines on regulatory aspects. This is an ideal handbook for undergraduates studying pharmaceutical or industrial manufacturing and supply chains as well for entrepreneurs and quality control professionals seeking to learn about CGMP standards and implementing quality assurance systems in the pharmaceutical sector.

A major new work on all aspects of water, the most used raw material ingredient in the pharmaceutical and biotechnology industries-used as an excipient in pharmaceutical formulations, as a cleaning agent, and as a separately packaged product diluent. Drawing on the author's extensive field experience with more than 400 pharmaceutical and related water systems, this book provides a comprehensive overview of the factors, varieties, and applications determining product containment, this concise reference provides authoritative information on containment processes. It reviews the historical context, definition, evolution, and application of containment technology, analyzes a variety of containment techniques in new and renovated facilities, and provides a practical implementation of the lifecycle approach to process validation. This book is an essential reference for pharmaceutical engineers, quality control professionals, and regulatory affairs specialists.

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ISPE Baseline Pharmaceutical Engineering Guide for New and Renovated Facilities  
Biopharmaceutical Manufacturing Facilities  
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Science and Risk-based Approach for the Delivery of Facilities, Systems, and Equipment  
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Water and Steam Systems  
Sterile Product Manufacturing Facilities  
Vol. 3  
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Commissioning and Qualification  
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Pharmaceutical Engineering Guides for New and Renovated Facilities. V.4 : Water and Steam Systems  
Baseline Pharmaceutical Engineering Guide  
Pharmaceutical Engineering Guides for New and Renovated Facilities: V.3: Sterile Manufacturing Facilities  
Risk-based Manufacture of Pharmaceutical Products  
A Guide to Managing Risks Associated with Cross-contamination  
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Technology Transfer  
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Biopharmaceutical Manufacturing Facilities  
Water and Steam Systems  
Good Quality Practice (GQP) in Pharmaceutical Manufacturing: A Handbook  
Bentham Science Publishers

Delivering an encompassing overview of the factors, varieties, and applications determining product containment, this concise reference provides authoritative information on containment processes. It reviews the historical context, definition, evolution, and application of containment technology, analyzes a variety of containment techniques in new

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