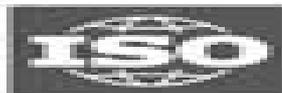

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Environmental Monitoring for Cleanrooms and Controlled Environments Anne Marie Dixon,2016-04-19 A critical technology in the science of contamination control environmental monitoring is a technique that provides important data on the quality of a process processing environment and final product which can aid scientists in identifying and eliminating potential sources of contamination in cleanrooms and controlled environments In response

Good Quality Practice (GQP) in Pharmaceutical Manufacturing: A Handbook Jordi Botet,2015-09-28 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

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Introduction to Contamination Control and Cleanroom Technology Matts Ramstorp,2000-12-27 Contamination control is being used by more and more industries where the highest level of cleanliness and hygiene is of vital importance This book covers the basic principles of contamination control and cleanroom technology from a holistic point of view It deals with cleanliness and hygiene and their effects on the outcome of a process reflecting the latest results from both scientific and practical points of view The following topics are covered contaminants and how they are measured cleanrooms and clean zones cleaning and decontamination cleanroom clothing the impact of people on cleanliness Intended as an introduction to the area of contamination control the text is also an excellent source of knowledge for people with both theoretical and practical experience The Swedish version has been used for a long time within the Nordic countries as a basic training textbook within the pharmaceutical microelectronics food and beverage optics and many other industries

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Semiconductor Manufacturing Handbook Hwaiyu Geng,2005-05-18 This handbook will provide engineers with the principles applications and solutions needed to design and manage semiconductor manufacturing operations Consolidating the many complex fields of semiconductor fundamentals and manufacturing into one volume by deploying a team of world class specialists it allows the quick look up of specific manufacturing reference data across many

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