



Good Manufacturing Practices (GMP) Policy Manual

In order to ensure that products are handled safely and effectively at all times whilst within the company premises a Good Manufacturing Practices (GMP) Policy Manual has been developed and implemented.

All employees are responsible to ensure that Company Procedures are followed correctly and that Good Manufacturing Practices (GMP) Policy Manuals being complied with.

Compliance is verified and recorded in accordance with Proc No 115.

Allergen Management

Cross Contamination

Management of Cross Contamination is carried out in accordance with Proc No 140 Allergen Management.

Amenities

Hand Washing

Detergent, sanitiser and hand drying facilities are available in employee amenities room and must be used in accordance with Procedure No 105.

Toilets

Toilets are to be well maintained and cleaned adequately, positioned within walking distance of all food related activities and not open directly to production facilities.

Equipment

Clothing Policy

Rules for wearing, cleaning and changing clothing and PPE are contained in Procedure No's 105 and XXX.

Food Contact Equipment

Equipment in contact with food is to be stored, cleaned and maintained in such a manner as to prevent food contamination.

Food Contact Surfaces

Food contact surfaces are to be in good condition, well maintained and cleaned in such a manner as to prevent food contamination.

Pallets

Pallets are to be kept in good condition so as to prevent either damage to product or foreign object contamination.

Preventive Maintenance

Equipment is maintained in accordance with Proc No 135 Maintenance - Premises and Equipment

Steel Wool

Steel wool is expressly forbidden for use in cleaning of utensils

Gmp Policy Manual

Michael Hoelscher



Gmp Policy Manual:

The Certified Pharmaceutical GMP Professional Handbook Mark Allen Durivage,2016-05-23 The purpose of this handbook is to assist individuals for the Certified Pharmaceutical Good Manufacturing Practices Professional CPGP examination and provide a reference for the practitioner The second edition reflects the Body of Knowledge which was updated in 2015 This edition has also incorporated additional information including updated references The updates reflect the current trends and expectations of the evolving pharmaceutical industry driven by consumer expectations and regulatory oversight This handbook covers compliance with good manufacturing practices GMPs as regulated and guided by national and international agencies for the pharmaceutical industry It covers finished human and veterinary drugs and biologics and combination devices as well as their component raw materials including active pharmaceutical ingredients APIs and excipients and packaging and labeling operations [GMP/ISO Quality Audit Manual for Healthcare Manufacturers and Their Suppliers, \(Volume 1 - With Checklists and Software Package\)](#) Leonard Steinborn,2019-04-23 Volume 1 of this two part package provides a complete set of checklists for internal and contract device and drug manufacturers and developers contract software developers and suppliers of chemical printed material electronic component and general supplies It also includes a simulated QSIT audit and a new product market launch All of these **The ASQ Certified Pharmaceutical GMP Professional Handbook** Mark Allen Durivage,2024-09-30 The ASQ Certified Pharmaceutical GMP Professional Handbook assists candidates preparing for the Certified Pharmaceutical Good Manufacturing Practices Professional CPGP examination and serves as a handy reference guide for practitioners in the field This handbook covers compliance with good manufacturing practices GMPs as regulated and guided by national and international agencies for the pharmaceutical industry *FDA Compliance Policy Guides Manual ,1977* *Sequoia and Kings Canyon National Parks (N.P.), Middle and South Forks of the Kings River and North Fork of the Kern River, General Management Plan ,2006* *Personnel Policy Handbook* William S. Hubbartt,1993 Here s a guide for all managers charged with creating and updating their company s human resource policy manuals This definitive handbook not only covers all areas of employee relations it also tackles the full range of critical contemporary HR issues such as AIDS substance abuse and chemical safety Managers can take advantage of how to instructions to organize and write a manual timesaving checklists and worksheets and invaluable tips on how to write personnel policies that lead to clear understanding and interpretation Alerting the reader to legal pitfalls the handbook covers employment policies leaves of absence pay discipline and discharge benefits union relations and more Plus its 100 helpful illustrations include sample forms flow charts and a complete sample policy manual [Handbook of Pharmaceutical Manufacturing Formulations, Third Edition](#) Sarfaraz K. Niazi,2019-12-06 The Handbook of Pharmaceutical Manufacturing Formulations Third Edition Volume Four Semisolid Products is an authoritative and practical guide to the art and science of formulating drugs for commercial manufacturing With thoroughly revised and expanded content this fourth volume of a six

volume set compiles data from FDA and EMA new drug applications patents and patent applications and other sources of generic and proprietary formulations including author's own experience to cover the broad spectrum of cGMP formulations and issues in using these formulations in a commercial setting. A must have collection for pharmaceutical manufacturers educational institutions and regulatory authorities this is an excellent platform for drug companies to benchmark their products and for generic companies to formulate drugs coming off patent. Features Largest source of authoritative and practical formulations cGMP compliance guidance and self audit suggestions Differs from other publications on formulation science in that it focuses on readily scalable commercial formulations that can be adopted for cGMP manufacturing Tackles common difficulties in formulating drugs and presents details on stability testing bioequivalence testing and full compliance with drug product safety elements Written by a well recognized authority on drug and dosage form development including biological drugs and alternative medicines

Handbook of Pharmaceutical Manufacturing Formulations Sarfaraz K. Niazi, 2016-04-19 While liquid drugs do not share the compression problems of solid dosage forms the filling problems of powder dosage forms or the consistency problems of semisolid dosage forms they do have their own set of considerations in the formulation and manufacturing stages Highlights from Liquid Products Volume Three include practical details in

Food and Drink - Good Manufacturing Practice Institute of Food Science and Technology, Louise Manning, 2012-11-26 Good Manufacturing Practice GMP refers to advice and guidance put in place to outline the aspects of production and testing that can impact the quality and safety of a product In the case of food and drink GMP is aimed at ensuring that products are safe for the consumer and are consistently manufactured to a quality appropriate to their intended use Manufacturers have for several years been driving towards such goals as Total Quality Management TQM lean manufacturing and sustainability GMP is bound up with these issues The ever increasing interest amongst consumers retailers and enforcement authorities in the conditions and practices in food manufacture and distribution increases the need for the food manufacturer to operate within clearly defined policies such as those laid down in GMP The ability to demonstrate that Good Manufacturing Practice has been fully and effectively implemented could in the event of a consumer complaint or a legal action reduce the manufacturer's liability and protect them from prosecution First launched in 1986 IFST's Good Manufacturing Practice Guide has been widely recognized as an indispensable reference work for food scientists and technologists It sets out to ensure that food manufacturing processes deliver products that are uniform in quality free from defects and contamination and as safe as it is humanly possible to make them This 6th edition has been completely revised and updated to include all the latest standards and guidance especially with regard to legislation driven areas such as HACCP The Guide is a must have for anyone in a managerial or technical capacity concerned with the manufacture storage and distribution of food and drink It is also a valuable reference for food education training and for those involved in food safety and enforcement Food scientists in academic and industry environments will value its precision and policy makers and

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