

MONTHLY CLEANING SCHEDULE FOR GMP PROCESSING AREA																													
GMP AREA	RESPONSIBLE	WEEK 1							WEEK 2							WEEK 3							WEEK 4						
		M	T	W	TH	F	SA	SU	M	T	W	TH	F	SA	SU	M	T	W	TH	F	SA	SU	M	T	W	TH	F	SA	SU
TABLET CORE AREA	PH	x	x	x	x	x			x	x	x	x	x			x	x	x	x	x			x	x	x	x	x		
TABLET FILLING	KB	x		x		x		x	x		x		x		x	x		x		x		x	x		x		x		x
PACKAGING AREA	KH	x			x				x			x				x			x				x			x			
LABELLING AREA	PP	x							x							x							x						
DISPENSING AREA	PH	x	x	x	x	x			x	x	x	x	x			x	x	x	x	x			x	x	x	x	x		
COOL ROOM	SS	x							x							x							x						
STAGING AREA	PP	x			x				x			x				x			x				x			x			
GENERAL WAREHOUSE	KB				x							x							x							x			
CORRIDORS	KH	x		x		x		x	x		x		x		x	x		x		x		x	x		x		x		x
PLANT ROOM	PS				x							x							x							x			
OUTSIDE CLEAN ROOM	PH	x	x	x	x	x			x	x	x	x	x			x	x	x	x	x			x	x	x	x	x		
AIR LOCKS	SS	x		x		x		x	x		x		x		x	x		x		x		x	x		x		x		x
HVAC FILTERS	KB	x														x													
QC LABORATORY	PH	x		x		x		x	x		x		x		x	x		x		x		x	x		x		x		x
MICRO LABORATORY	PP	x		x		x		x	x		x		x		x	x		x		x		x	x		x		x		x
ENG. WORKSHOP	PS															x													
GENERAL OFFICE	KH	x			x				x			x				x			x				x			x			

Gmp Cleaning Schedule

Lei Huang

A red circular graphic with a gradient, appearing as a semi-circle or a partial circle, located to the right of the name bar.

Gmp Cleaning Schedule:

Food and Drink - Good Manufacturing Practice Institute of Food Science and Technology, Louise Manning, 2018-10-22 The latest updated edition of the market leading guide to Good Manufacturing Practice GMP in the food and drink industry This all new 7th edition of Food and Drink Good Manufacturing Practice A Guide to its Responsible Management features a wealth of new information reflecting changes in the industry and advances in science that have occurred since the publication of the last edition back in 2013 They include topics such as Food Safety Culture Food Crime and Food Integrity Management Systems Food Crime Risk Assessment including vulnerability risk assessment and Threat Analysis Critical Control Point TACCP Security and Countermeasures Food Toxins Allergens and Risk Assessment Provenance and authenticity Electronic and digital traceability technologies Worker Welfare Standards Smart Packaging Food Donation Controls and Animal Food Supply Safety Culture Provenance and integrity testing and Sustainability Issues In addition to the new topics mentioned above Food and Drink Good Manufacturing Practice 7th Edition offers comprehensive coverage of information in chapters on Quality Management System Hazard Analysis Critical Control Point HACCP Premises and Equipment Cleaning and Sanitation Product Control Testing and Inspection Heat Preserved Foods Frozen Foods Foods for Catering and Vending Operations and much more Comprises both general guidance and food sector specific requirements for good manufacturing practice Incorporates all the most recent developments and changes in UK and EU law Provides a readable and accessible reference for busy managers in the food industry Food and Drink Good Manufacturing Practice A Guide to its Responsible Management 7th Edition is a valuable reference for anyone in a managerial or technical capacity concerned with the manufacture storage and distribution of food and drink The book is also a must read for the recommended reading lists for food science food technology and food policy undergraduate and postgraduate studies IFST the Institute of Food Science and Technology is the leading qualifying body for food professionals in Europe and the only professional qualifying body in the UK concerned with all aspects of food science and technology **The CDC Handbook - A Guide to Cleaning and Disinfecting Clean**

Rooms Dr. Tim Sandle, 2012-08-02 The Cleaning and Disinfection handbook is aimed at those working within the pharmaceutical and healthcare sectors around the world as well as providing valuable information for students and for the general reader The book provides comprehensive detail on different types of disinfectants and their modes of action explains the problems of microbial destruction and resistance introduces cleaning techniques and the latest safety regulations expounds upon the application of cleaning within healthcare and pharmaceutical environments noting current national and international standards The book also provides guidance on disinfectant efficacy testing Assembled by expert practitioners the book balances theoretical concepts with sound practical advice and is likely to become the definitive text on keeping contamination in control within clean areas and controlled environments With this second edition the book is fully updated in line with the latest standards and regulations *Handbook of Food Preservation* M. Shafiur Rahman, 2007-07-16 The

processing of food is no longer simple or straightforward but is now a highly interdisciplinary science. A number of new techniques have developed to extend shelf life, minimize risk, protect the environment and improve functional sensory and nutritional properties. The ever-increasing number of food products and preservation techniques create a need for Food Safety Management Programs. Debby Newslow, 2013-12-20. The safety of food products is fundamental. The value of an effective and well-defined, implemented and maintained management system is priceless. When it is integrated into a process, it supplies the necessary foundation and structure to help provide the consumer with a safe product of the highest quality. Food Safety Management Programs Applied. Food Processing Technology. P.J. Fellows, 2016-10-04. Food Processing Technology: Principles and Practice. Fourth Edition has been updated and extended to include the many developments that have taken place since the third edition was published. The new edition includes an overview of the component subjects in food science and technology, processing stages, important aspects of food industry management not otherwise considered, e.g. financial management, marketing, food laws and food industry regulation, value chains, the global food industry and overarching considerations, e.g. environmental issues and sustainability. In addition, there are new chapters on industrial cooking, heat removal, storage and distribution, along with updates on all the remaining chapters. This updated edition consolidates the position of this foundational book as the best single volume introduction to food manufacturing technologies available, remaining as the most adopted standard text for many food science and technology courses. Updated edition completely revised with new developments on all the processing stages and aspects of food industry management not otherwise considered, e.g. financial management, marketing, food laws and food industry regulation and more. Introduces a range of processing techniques that are used in food manufacturing. Explains the key principles of each process, including the equipment used and the effects of processing on microorganisms that contaminate foods. Describes post-processing operations, including packaging and distribution logistics. Includes extra textbook elements such as videos and calculations slides, in addition to summaries of key points in each chapter. *Food Safety Lessons for Cannabis-Infused Edibles*. Kathy Knutson, 2020-04-28. *Food Safety Lessons for Cannabis-Infused Edibles* details the world of cannabis-infused edibles and the way its manufacturing is evolving as the industry moves from isolation to regulatory compliance. The cannabis industry has unique challenges as cannabis-infused edibles are not regulated as food, drugs or dietary supplements at the federal level. Despite these current conditions, the industry is aware of the need to examine the safety of these edibles and prepare for a future of federal compliance. The book looks at the cannabis industry through a scientific lens to increase awareness and expertise in food safety within the field of cannabis-infused edibles. Includes lessons learned by the food industry. Presents unique challenges in the manufacture of cannabis-infused edibles. Provides information of US Federal food safety compliance. Explores the current state of research regarding edibles. **Dictionary of Flavors**. Dolf De Rovira, Sr., 2017-03-14. The third edition of this highly popular scientific reference continues to provide a unique approach to flavors, flavor chemistry and

natural products Dictionary of Flavors features entries on all flavor ingredients granted G R A S status compounds used in the formulation of food flavors and related food science and technology terms Allergies and intolerances are addressed along with strategies to avoid allergenic compounds This latest edition has been fully updated to reflect new ingredients available on the market as well as developments in safety standards and the international regulatory arena Dolf De Rovira applies his extensive experience to make this the most comprehensive guide to flavors available **Process Validation in**

Manufacturing of Biopharmaceuticals Anurag Singh Rathore, Hal Baseman, Scott Rudge, 2023-12-18 The fourth edition of Process Validation in Manufacturing of Biopharmaceuticals is a practical and comprehensive resource illustrating the different approaches for successful validation of biopharmaceutical processes A pivotal text in its field this new edition provides guidelines and current practices contains industrial case studies and is expanded to include in depth analysis of the new Process Validation PV guidance from the US FDA Key Features Offers readers a thorough understanding of the key concepts that form the basis of a good process validation program for biopharmaceuticals Includes case studies from the various industry leaders that demonstrate application of these concepts Discusses the use of modern tools such as multivariate analysis for facilitating a process validation exercise Covers process characterization techniques for scaling down unit operations in biopharmaceutical manufacturing including chromatography chemical modification reactions ultrafiltration and microfiltration and practical methods to test raw materials and in process samples Providing a thorough understanding of the key concepts that form the basis of a good process validation program this book will help readers ensure that PV is carried out and exceeds expectations Fully illustrated this is a much needed practical guide for biopharmaceutical manufacturers *Handbook of Validation in Pharmaceutical Processes, Fourth Edition* James

Agalloco, Phil DeSantis, Anthony Grilli, Anthony Pavell, 2021-10-28 Revised to reflect significant advances in pharmaceutical production and regulatory expectations Handbook of Validation in Pharmaceutical Processes Fourth Edition examines and blueprints every step of the validation process needed to remain compliant and competitive This book blends the use of theoretical knowledge with recent technological advancements to achieve applied practical solutions As the industry's leading source for validation of sterile pharmaceutical processes for more than 10 years this greatly expanded work is a comprehensive analysis of all the fundamental elements of pharmaceutical and bio pharmaceutical production processes Handbook of Validation in Pharmaceutical Processes Fourth Edition is essential for all global health care manufacturers and pharmaceutical industry professionals Key Features Provides an in depth discussion of recent advances in sterilization Identifies obstacles that may be encountered at any stage of the validation program and suggests the newest and most advanced solutions Explores distinctive and specific process steps and identifies critical process control points to reach acceptable results New chapters include disposable systems combination products nano technology rapid microbial methods contamination control in non sterile products liquid chemical sterilization and medical device manufacture *Cell Therapy*

Adrian P. Gee,2021-11-10 This new edition presents a fully updated and expanded look at current Good Manufacturing Practice cGMP for cell therapy products It provides a complete discussion of facility design and operation including details specific to cord blood banking cell processing vector production and qualification of a new facility Several chapters cover facility infrastructure including cleaning and maintenance vendor qualification writing a Standard Operating Procedure staff training and process validation The detailed and invaluable product information covers topics like labelling release and administration transportation and shipment et al Further chapters cover relevant topics like writing and maintaining investigational new drug applications support opportunities in North America and the European Union commercial cell processing and quality testing services and financial considerations for academic GMP facilities A chapter on future directions rounds out Cell Therapy cGMP Facilities and Manufacturing making it essential reading for any cell therapy professional involved in the development use or management of this type of facility *Quality Assurance G*

Welty,2013-06-30 Quality assurance is necessary to maintain quality and services in the pharmaceutical and life science industries Quality assurance demonstrates that the logic and practice of problem solving can integrate both program efficacy and regulatory compliance This title is divided into three parts the first part discusses the process by which a problem in regulated industry is identified for example a manufacturing deviation that leads to an adulterated drug product and reviews the decision making steps involved in remedying the problem The second part delves into the staff training requirements of procedures that are thereby revised The third part expands on this discussion by considering piloting the proposed training module preparing assessments of trainee proficiency evaluating the training module including integrating rigorous evaluative designs with formative program improvement and documenting the entire effort Presents a comprehensive view of the field of quality assurance An approach grounded in direct experience Uses diagrams and figures to clarify analytical points

Cleaning Validation Manual Syed Imtiaz Haider,2010-05-24 During the past decades enormous progress and enhancement of pharmaceutical manufacturing equipment and its use have been made And while there are support documents books articles and online resources available on the principles of cleaning and associated processing techniques none of them provides a single database with convenient ready to use training tools Until now Cleaning Validation Manual A Comprehensive Guide for the Pharmaceutical and Biotechnology Industries elucidates how to train the man power involved in development manufacturing auditing and validation of bio pharmaceuticals on a pilot scale leading to scale up production With over 20 easy to use template protocols for cleaning validation of extensively used equipments this book provides technical solutions to assist in fulfilling the training needs of finished pharmaceutical manufacturers Drawing on the authors more than two decades of experience in the pharmaceutical and biotech industries the text offers hands on training based on current approaches and techniques The book does not merely provide guidelines or thought processes rather it gives ready to use formulas to develop Master Plan SOPs and validation protocols It includes cleaning procedures for the most commonly

used equipment in various manufacturing areas and their sampling points using a pharmaceutical manufacturing site with both sterile and non sterile operations as the case facility It also provides the training guidelines on downloadable resources to enable users to amend or adopt them as necessary Grounded in practicality the book s applicability and accessibility set it apart It can be used as a guide for implementing a cleaning validation program on site without the help of external consultants making it a resource that will not be found collecting dust on a shelf but rather referred to again and again

Guide to Quality Management Systems for the Food Industry Ralph Early,2012-12-06 Whenever I step into an aeroplane I cannot avoid considering the risks associated with flying Thoughts of mechanical failure pilot error and terrorist action fill my mind I try to reassure myself with statistics which tell me there is greater chance of injury crossing the road The moment the plane takes off I am resigned to my fate placing faith in pilots who are highly qualified and superbly trained for the task of delivering me safely to my destination To be a passenger in an aeroplane is to express faith in the systems used by the airline It is to express a faith in the quality of the airline s organisation and the people who work within it The same is true of surgery Thoughts of mortality are difficult to avoid when facing the surgeon s knife However faith in the surgeon s training and skill faith in the anaesthetist and theatre technicians faith in the efficient resources and quality of the hospital all help to convince that there is little need to worry Apart from flying and surgery there are many facets of life which entail risk but knowing the risks we willingly place our confidence in others to deliver us safely In the consumption of food however few of us consider the risks Everyday if we are fortunate we eat food Food sustains and gives us pleasure Food supports our social interactions

Good Manufacturing Practices for Pharmaceuticals, Seventh Edition Graham P. Bunn,2019-02-04 This book provides insight into the world of pharmaceutical quality systems and the key elements that must be in place to change the business and organizational dynamics from task oriented procedure based cultures to truly integrated quality business systems that are self detecting and correcting Chapter flow has been changed to adopt a quality systems organization approach and supporting chapters have been updated based on current hot topics including the impact of the worldwide supply chain complexity and current regulatory trends Key Features Presents insight into the world of pharmaceutical quality systems Analyzes regulatory trends and expectations Includes approaches and practices used in the industry to comply with regulatory requirements Discusses recent worldwide supply chain issues Delivers valuable information to a worldwide audience regarding the current GMP practices in the industry

Auditing in the Food Industry M Dillon,C Griffith,2001-07-12 The food industry faces an unprecedented level of scrutiny Consumers are not only concerned with the safety and quality of food products but also the way in which they are produced At the same time the food industry has developed new ways of assuring appropriate standards for its products and their methods of production developing systems such as TQM and HACCP to identify and manage key steps in production These new methods require new skills in auditing Auditing in the food industry provides an authoritative guide to the range of standards and the auditing

skills they demand Part one sets the scene with an introductory chapter reviewing developments in standards affecting the food industry There then follows chapters on how retailers audit their suppliers and how governments have moved from a traditional inspection role to one of regulatory verification with its emphasis on auditing the robustness of a business s own systems for managing safety and quality Part two examines the key aspects of safety and quality A first chapter reviews the ways retailers assess supplier HACCP systems There is then a chapter reviewing TQM systems that provides a context for a discussion of auditing techniques for HACCP based quality systems A final chapter looks at standards governing the analytical methods used in safety and quality control Part three considers newer standards that are becoming increasingly important in the food industry There are chapters on benchmarking an organisation against others as a way of improving performance auditing the impact of food processing operations on the environment and auditing organic food processing Auditing in the food industry is a valuable guide to the range of standards facing the food industry and the ways it can audit and thus improve the quality of its performance

Pharmaceutical Manufacturing Formulations Dr. Priyanka Gupta Manglik,2024-08-15 This book provides detailed insight into the various aspects of pharmaceutical manufacturing covering formulations process design technology and regulatory requirements essential for professionals in the pharma industry

Handbook of Pharmaceutical Manufacturing Formulations Sarfaraz K. Niazi,2016-04-19 Providing methodologies that can serve as a reference point for new formulations the second volume covers uncompressed solids which include formulations of powders capsules powders ready for reconstitution and other similar products Highlights from

Uncompressed Solid Products Volume Two include the fundamental issues of good manufacturin *Process Chemistry in the Pharmaceutical Industry* Kumar Gadamasetti,1999-05-06 Providing guidance for chemists and other scientists entering pharmaceutical discovery and development this up to the minute reference presents contributions from an international group of nearly 50 renowned researchers offering a solid grounding in synthetic and physical organic chemistry and clarifying the roles of various special

FSMA and Food Safety Systems Jeffrey T. Barach,2017-01-24 The FDA s Food and Drug Administration FSMA Food Safety Modernization Act is the most sweeping reform of United States food safety laws in more than 70 years The key to successful implementation of FSMA rules depends on building a comprehensive Food Safety System with effective prerequisite programs in place and a well designed Food Safety Plan that incorporates risk based preventive controls to mitigate hazards This book provides essential guidance for small to mid sized businesses on how to design implement and maintain a world class Food Safety Plan that conforms to FSMA regulations With practical and up to date advice the author offers a straight forward approach for readers to successfully migrate into FSMA The inclusion of fully developed Food Safety Plans as well as examples of hazards and preventative controls make this a must read not only for those that are new to the regulations but also those with a plan already in place **FSMA and Food Safety Systems A Guide to Understanding and Implementing the Rules** is an indispensable resource for all those managing the manufacture of FDA

regulated products food safety regulators and educators as well as scientists and students of food science and technology

HACCP in Meat, Poultry, and Fish Processing A. M. Pearson, T. R. Dutson, 2012-12-06 The RACCP hazard analysis critical control point concept for food products was an outgrowth of the US space program with the demand for a safe food supply for manned space flights by the National Aeronautics and Space Administration NASA The original work was carried out by the Pillsbury Company under the direction of Roward E Bauman who as the author of chapter 1 describes the evolution of the RACCP system and its adaptation to foods The second chapter discusses the adoption of RACCP principles and explains how they fit into the USDA and FDA meat poultry and seafood inspection systems The next chapter discusses how RACCP principles can be extended to production of meat poultry and seafoods a most important area involved in producing a safe food supply Chapter 4 deals with the use of RACCP in controlling hazards encountered in slaughtering and distribution of fresh meat and poultry while chapter 5 discusses the problem both spoilage and hazards involved in processing and distribution of meat poultry and seafood products Chapter 6 covers the entire area of fish and seafoods including both fresh and processed products from the standpoints of spoilage and hazards

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