

Guideline for Equipment and System Qualification

[pharmabeginners](#) / October 7, 2020

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Standard Operating Procedure (SOP) and Guideline for preparation of Equipment / System Qualification (URS, IQ, OQ, PQ, FAT, SAT, etc.) documents, execution of Qualification activities, Review and Compilation of data, Assessment and Interpretation of Qualification & validation activity results.

Equipment and System Qualification

1.0 Purpose :

To lay down the procedure for preparation of Qualification documents, execution of Qualification activities, Review and Compilation of Data, Assessment, and Interpretation of Qualification & validation activity results.

2.0 Scope :

The standard operating procedure applicable for the preparation of qualification and re-qualification documents for all equipment & system and execution of qualification activities performed.

3.0 Responsibility :

Officer and above of User, Engineering and QA Department shall be responsible for the preparation and execution of the equipment and system Qualification and Re-qualification activities.

4.0 Accountability

Head- User, Engineering, and Quality Assurance shall be responsible for compliance.

5.0 Procedure – Equipment and System Qualification :

- **Qualification Approach – Equipment and System :**
- The universal “V model” approach shall be followed for all the Qualification and Validation activities.
- The model is as follows:

Equipment System Verification Qualification

Mark Allen Durivage



Equipment System Verification Qualification:

Sterile Manufacturing Sam A. Hout, 2021-07-04 This book highlights key ideas and factors to coach and guide professionals involved in learning about Sterile Manufacturing and operational requirements It covers regulations and guidelines instituted by the FDA ISPE EMA MHRA and ICH emphasizing good manufacturing practice and inspection requirements in the manufacturing of medicinal products Additionally this book provides the fundamentals of aseptic techniques quality by design risk assessment and management in support of sterile operations applications It creates a link to the implementation of business practices in drug manufacturing and healthcare and forms a correlation between design strategies including a step by step process to ensure reliability safety and efficacy of healthcare products for human and animal use The book also provides a connection between drug production and regulated applications by offering a review of the basic elements of sterile processing and how to remain viable with solid strategic planning The book is a concise reference for professionals and learners in the field of sterile operations that governs primarily pharmaceutical and medical device space but can also extend to food and cosmetics that require clean aseptic manufacturing applications It also helps compounding pharmacists and GMP inspectors and auditors

Pharmaceutical Analysis for Small Molecules Behnam Davani, 2017-08-01 A comprehensive introduction for scientists engaged in new drug development analysis and approvals Each year the pharmaceutical industry worldwide recruits thousands of recent science graduates especially chemistry analytical chemistry pharmacy and pharmaceutical majors into its ranks However because of their limited background in pharmaceutical analysis most of those new recruits find making the transition from academia to industry very difficult Designed to assist both recent graduates as well as experienced chemists or scientists with limited regulatory compendial or pharmaceutical analysis background make that transition Pharmaceutical Analysis for Small Molecules is a concise yet comprehensive introduction to the drug development process and analysis of chemically synthesized small molecule drugs It features contributions by distinguished experts in the field including editor and author Dr Behnam Davani an analytical chemist with decades of technical management and teaching experience in compendial regulatory and industry This book provides an introduction to pharmaceutical analysis for small molecules non biologics using commonly used techniques for drug characterization and performance tests The driving force for industry to perform pharmaceutical analyses is submission of such data and supporting documents to regulatory bodies for drug approval in order to market their products In addition related required supporting studies including good laboratory documentation practices including analytical instrument qualification are highlighted in this book Topics covered include Drug Approval Process and Regulatory Requirements private standards Pharmacopeias and Compendial Approval Process public standards Common methods in pharmaceutical analysis typically compendial Common Calculations for assays and impurities and other specific tests Analytical Method Validation Verification Transfer Specifications including how to handle out of specification OOS and out of trend OOT Impurities

including organic inorganic residual solvents and elemental impurities Good Documentation Practices for regulatory environment Management of Analytical Laboratories Analytical Instrument Qualifications including IQ OQ PQ and VQ Due to global nature of pharmaceutical industry other topics on both regulatory ICH and Compendial harmonization are also highlighted Pharmaceutical Analysis for Small Molecules is a valuable working resource for scientists directly or indirectly involved with the drug development process including analytical chemists pharmaceutical scientists pharmacists and quality control quality assurance professionals It also is an excellent text reference for graduate students in analytical chemistry pharmacy pharmaceutical and regulatory sciences

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

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

Handbook of Aseptic Processing and Packaging Jairus R. D. David, Pablo M. Coronel, Josip Simunovic, 2022-09-09 Nine years have passed since the second edition of the Handbook of Aseptic Processing and Packaging was published Significant changes have taken place in several aseptic processing and packaging areas These include aseptic filling of plant based beverages for non refrigerated shelf stable formats for longer shelf life and sustainable packaging along with cost of environmental benefits to leverage savings on energy and carbon footprint In addition insight into safe processing of particulates using two and three dimensional thermal processing followed by prompt cooling is provided In the third edition the editors have compiled contemporary topics with information synthesized from internationally recognized authorities in their fields In addition to updated information 12 new chapters have been added in this latest release with content on Design of the aseptic processing system and thermal processing Thermal process equipment and technology for heating and cooling Flow and residence time distribution RTD for homogeneous and heterogeneous fluids Thermal process and optimization of aseptic processing containing solid particulates

Aseptic filling and packaging equipment for retail products and food service Design of facility infrastructure and utilities
 Cleaning and sanitization for aseptic processing and packaging operations Microbiology of aseptically processed and
 packaged products Risk based analyses and methodologies Establishment of validated state for aseptic processing and
 packaging systems Quality and food safety management systems for aseptic and extended shelf life ESL manufacturing
 Computational and numerical models and simulations for aseptic processing Also there are seven new appendices on original
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 of combined experience encompassing manufacturing innovation in processing and packaging R D quality assurance and
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 processing industry The future of aseptic processing and packaging of foods and beverages will be driven by customer facing
 convenience and taste use of current and new premium clean label natural ingredients use of multifactorial preservation or
 hurdle technology for maximizing product quality and sustainable packaging with claims and messaging

Validation of Pharmaceutical Processes James P. Agalloco, Frederick J. Carleton, 2007-09-25 Completely revised and updated to reflect the
 significant advances in pharmaceutical production and regulatory expectations this third edition of *Validation of
 Pharmaceutical Processes* examines and blueprints every step of the validation process needed to remain compliant and
 competitive The many chapters added to the prior compilation examine va

The Standard Handbook for Aeronautical and Astronautical Engineers Mark Davies, 2003 Designed as a one stop reference for engineers of all disciplines in
 aeronautical and aerospace engineering this handbook seeks to filter mechanical engineering applications to specifically
 address aircraft and spacecraft science and military engineering

Validation of Aseptic Pharmaceutical Processes
 Frederick J. Carleton, James P. Agalloco, 1986

IEEE/CHMT International Electronic Manufacturing Technology Symposium, 1989

The Massachusetts register, 1995

Good Manufacturing Practice Guide for Active Pharmaceutical Ingredients, 2001

Guide to the Preparation, Use and Quality Assurance of Blood Components Council of
 Europe, 2007 In the absence of substitutes the use of blood components remains essential in therapy This guide contains a
 compendium of measures designed to ensure the safety efficacy and quality of blood components and is particularly intended
 for all those working in blood transfusion services In accordance with the approach recommended by the Council of Europe
 in this field it is based on the premise of voluntary non remunerated blood donation It describes the different blood
 components and gives information on their clinical indications and possible side effects

Technical Report Series, 1950

A Laboratory Quality Handbook of Best Practices Donald C. Singer, 2001

Quality Assurance Integrated Training Packages, 1992 The competence of the personnel to establish and implement an effective quality assurance programme is
 essential for attaining and maintaining the safety reliability and economic performance of a nuclear installation This manual

offers training packages covering quality assurance principles and practices which can be adjusted to suit different levels of management and working personnel and can be adapted to national variables and differing needs in the field of quality assurance

Validation Guidelines for Automated Control of Food Processing Systems Used for the Processing and Packaging of Preserved Foods ,2002 **ESA Bulletin** European Space Agency,1989 Commonwealth Of Australia Gazette Australia,1971 **Florida Administrative Weekly** ,1990 **Chemical Engineering Progress** ,2007

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