

		LABORATORIUM MIKROBIOLOGI FARMASI UNIVERSITAS MUHAMMADIYAH RIAU		Kode:	
Disiapkan:	Laboran	<i>STANDAR OPERATING PROCEDURE</i>		Hal.:	
Disetujui:	Ka. Lab	Pengoperasian Alat <i>Laminar Air Flow</i>		Terbitan:	2019
				Revisi:	
				Tanggal:	

1. LINGKUP

Standar Operating Procedures (SOP) ini melingkupi prosedur pengoperasian alat *Laminar Air Flow*

2. TUJUAN

Standar oprating Procedures (SOP) ini adalah untuk pengoperasian alat *Laminar Air Flow*

3. PRINSIP

Alat Mikroskop *Laminar Air Flow* dioperasikan merujuk kepada petunjuk pengoperasian (*operating instruction*) dan refrensi dari sumber lain seperti panduan laboratorium.

4. BAHAN KIMIA

Tidak ada penggunaan bahan kimia dalam proses pengoperasian.

5. PERALATAN

Laminar Air Flow (LAF)

6. PROSEDUR

- Nyalakan lampu UV, minimum selama 30 menit, sebelum laminar air flow digunakan. Hindarkan sinarnya dan mata.
- Siapkan semua alat-alat steril yang akan dipergunakan. Alat-alat yang dimasukkan ke dalam laminar air flow cabinet, disemprot terlebih dahulu dengan alkohol 70% atau spiritus.
- Meja dan dinding dalam LAF disemprot dengan alkohol 70% atau dengan spiritus untuk mensterilkan LAF.
- Blower pada LAF dihidupkan untuk menjalankan air flow.
- Nyalakan lampu dalam LAF.
- LAF sudah siap untuk digunakan

7. REFRENSI

- Prosedur Penggunaan Laminar Air Flow dari <https://www.microns.co.id/2020/06/cara-menggunakan-laminar-air-flow.html>
- Laminar Air Flow (LAF) : Pengertian, Fungsi dan Penggunaannya dari <https://enderupn.co.id/laminar-air-flow/>

How To Apply For Laf Sop

Missouri Valley College, Marshall, Mo

How To Apply For Laf Sop:

Standard Operating Procedures and Regulatory Guidelines GP Saluja, GP Singal, 2023-08-25 *Lafayette* Olivier Bernier, 2017-10-24 Historian Olivier Bernier draws an indelible portrait of the man who represented more than anyone else the idea of French nobility to all Americans of the early Republic and who represented to the French the idea of freedom and its American expression Lafayette was indeed the hero of two worlds Bernier's Lafayette much of it based on previously inaccessible documents is a man who lived the liberal ideal as few others have In the war for American independence this twenty year old was a stubborn tenacious and ultimately victorious commander the favorite of George Washington with whom he developed a unique father son relationship Returning to Paris with yearnings for a liberalized government he was soon caught up in the 1789 revolution first as its champion then as the guardian of the king finally as the only man capable of maintaining order in 1790 and 1791 Once the king fled the capital however Lafayette's position became untenable and he was forced to escape to Belgium But there the right wing emigres considered him a traitor and he was arrested and sent to Austria where he languished in prison for years Finally the diplomatic efforts of George Washington and other Americans led to his release and return to France Now Napoleon feared him as a potential rival a fear heightened when Lafayette went into self imposed exile to protest Napoleon's abuse of power During the revolution that followed Napoleon's downfall Lafayette maintained his liberal principles as few others bothered to and his position was vindicated by the uprising that installed the July monarchy and France's first middle class constitution Enriching this chronicle of a man and his age are the stories of young Gilbert's many loves as well as the steadfast relationship with his adoring wife And never far from the marquis's heart was his love for his adopted home He maintained it through a forty year correspondence with the Founding Fathers and an unrelenting if often quixotic defense of liberal ideals For its part the young American republic knew no grander celebrations than those thrown in honor of his return in 1824 *Validation Standard Operating Procedures* Syed Imtiaz Haider, 2006-05-30 Spanning every critical element of validation for any pharmaceutical diagnostic medical device or equipment and biotech product this Second Edition guides readers through each step in the correct execution of validating processes required for non aseptic and aseptic pharmaceutical production With 14 exclusive environmental performance evaluati **Federal Register** , 1990-03 **Annual Catalog** Missouri Valley College, Marshall, Mo, 1904 , Social Security Bulletin , 1992 Tournament of Appeals Roy B. Flemming, 2004 Canada's Supreme Court decides cases with far reaching effects on Canadian politics and public policies When the Supreme Court sets cases on its agenda it exercises nearly unrestrained discretion and considerable public authority But how does the Court choose these cases in the first place From the several hundred requests for judicial review filed every year how and why do the justices pick some cases but not others for review Tournament of Appeals investigates the leave to appeal process in Canada and explores how and why certain cases win a place on the Court's agenda and others do not Taking the approach that the process mimics a sports tournament

this study raises several vital questions For example is there an elite Supreme Court bar that routinely wins the tournament Do the Court s rules affect the tournament s outcomes Or does winning and losing reflect the resources of the parties As players in this tournament how do the judges play the game and how does it affect their votes to grant or deny judicial review Drawing from systematically collected information on the process applications and lawyers that has never before been used in studies of Canada s Supreme Court Roy B Flemming offers both a qualitatively and quantitatively based explanation of how Canada s justices grant judicial review The first of its kind this innovative study will draw the attention of lawyers academics and students in Canada as well as in the Commonwealth and European countries whose high courts share many features of the appeals process in Canada *The Sourcebook of Local Court and County Record Retrievers* Carl R. Ernst, Michael L. Sankey, 1999 **The Official Railway Equipment Register**, 1910 *Quality Operations Procedures for Pharmaceutical, API, and Biotechnology* Syed Imtiaz Haider, Erfan Syed Asif, 2012-06-06 To stay in compliance with regulations pharmaceutical medical and biotech companies must create quality SOPs that build in the regulatory requirements into actions and describe personal flow internal flow flow of information and processing steps Quality Operations Procedures for Pharmaceutical API and Biotechnology and the accompanying CD ROM take into account all major international regulations such as FDA EU GMP cGMP GLP PDA technical monographs PDA technical reports PMA concepts journals of PDA GCP and industry standard ISO 9000 to be in compliance with documentation guidelines No other resource deals exclusively with the key elements of quality control and quality assurance procedures for pharmaceutical operations and provides hands on templates to be tailored to achieve global regulatory compliance The book provides instant answers about what to include in critical quality assurance and quality control SOPs and how to enhance productivity The CD ROM contains nineteen quality control and thirty three quality assurance SOPs designed so that users can input them into their computers and use their Microsoft Word programs to edit and print these documents The book ensures minimization of the number of documents helping to reduce the nightmare like aura that surrounds an FDA audit The SOPs exclusively refer to the documents specially required for compliance however specific formats are not included to ensure that the electronic templates can be easily used by pharmaceutical bulk pharmaceutical medical device and biotechnology industries The combination of text and CD ROM presents a ready to use resource on the quality systems of aseptic pharmaceutical non aseptic production and to provide general information and guidelines They comprise a tool that can be used to develop a set of quality SOPs in order to support the road map established for the on time successful start up of the facility operation in compliance with the GMP requirements **Music News**, 1928 **Good Clinical, Laboratory and Manufacturing Practices** Philip Carson, Nigel Dent, 2007-10-31 Quality assurance and good laboratory practices are becoming essential knowledge for professionals in all sorts of industries This includes internal and external audit procedures for compliance with the requirements of good clinical laboratory and manufacturing practices Spanning chemical cosmetic and manufacturing industries Good Clinical Laboratory

and Manufacturing Practices Techniques for the QA professional is aimed at chemists clinicians ecotoxicologists operation managers pharmaceutical process managers quality assurance officers technicians and toxicologists In addition sections on harmonisation of quality systems will be of value to safety health and environment advisors This comprehensive and high level reference will be an indispensable guide to research laboratories in academia and industry Additional training material is also included

Laboratory Control System Operations in a GMP Environment David M. Bliesner, 2020-06-03
Develop an understanding of FDA and global regulatory agency requirements for Laboratory Control System LCS operations In Laboratory Control System Operations in a GMP Environment readers are given the guidance they need to implement a CGMP compliant Laboratory Control System LCS that fits within Global Regulatory guidelines Using the Quality Systems Approach regulatory agencies like the FDA and the European Medicine Agency have developed a scheme of systems for auditing pharmaceutical manufacturing facilities which includes evaluating the LCS In this guide readers learn the fundamental rules for operating a CGMP compliant Laboratory Control System Designed to help leaders meet regulatory standards and operate more efficiently the text includes chapters that cover Laboratory Equipment Qualification and Calibration Laboratory Facilities Method Validation and Method Transfer Laboratory Computer Systems Laboratory Investigations as well as Data Governance and Data Integrity The text also includes chapters related to Laboratory Managerial and Administrative Systems Laboratory Documentation Practices and Standard Operating Procedures and General Laboratory Compliance Practices Additionally a chapter outlining Stability Program operations is included in the text In addition to these topics it includes LCS information and tools such as End of chapter templates checklists and LCS guidance to help you follow the required standards Electronic versions of each tool so users can use them outside of the text An In depth understanding of what is required by the FDA and other globally significant regulatory authorities for GMP compliant systems For quality assurance professionals working within the pharmaceutical or biopharma industries this text provides the insight and tools necessary to implement government defined regulations

Memories of Lafayette Larroque, Charles, Text and photographs capture the festivals cuisine and history of the city once known as Vermillionville
Hendricks' Commercial Register of the United States ,1925 **The Topsy-turvy Hotel** Victor Roger, 1898 *How to Design and Implement Powder-to-Tablet Continuous Manufacturing Systems* Fernando Muzzio, Sarang Oka, 2022-04 How to Design and Implement Powder to Tablet Continuous Manufacturing Systems provides a comprehensive overview on the considerations necessary for the design of continuous pharmaceutical manufacturing processes The book covers both the theory and design of continuous processing of associated unit operations along with their characterization and control In addition it discusses practical insights and strategies that the editor and chapter authors have learned Chapters cover Process Analytical Technology PAT tools and the application of PAT data to enable distributed process control With numerous case studies throughout this valuable guide is ideal for those engaged in or learning about continuous processing in

pharmaceutical manufacturing Discusses the development of strategy blueprints in the design of continuous processes Shows how to create process flowsheet models from individual unit operation models Includes a chapter on characterization methods for materials the use of statistical methods to analyze material property data and the use of material databases Covers the evolving regulatory expectations for continuous manufacturing Provides readers with ways to more effectively navigate these expectations *Scientific and Technical Aerospace Reports* ,1990 Enhancing compliance to good manufacturing practices and pharmaceutical quality system requirements in vaccine production World Health Organization,2024-06-25

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