

		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

Roman Wölfel



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Standard Operating Procedures and Regulatory Guidelines GP Saluja, GP Singal, 2023-08-25 *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 evaluations, *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 **Social Security Bulletin**, 1992 **Signal**, 2012 *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 **In Vitro Fertilization** Zsolt Peter Nagy, Alex C. Varghese, Ashok Agarwal, 2019-07-03 Now in its revised and expanded second edition including over 20 new chapters this comprehensive textbook remains a unique and accessible description of the current and developing diagnostic and treatment techniques and technologies comprising in vitro fertilization IVF Arranged thematically in sections each chapter covers a key topic in IVF in a sensible presentation Parts one and two describe the planning design and organization of an ART unit and IVF laboratory and equipment and systems respectively The sections that follow provide detailed descriptions of IVF techniques embryo culture methods sperm processing and selection insemination procedures micromanipulation embryo evaluation cryopreservation and embryo transfer Concluding sections

address issues of management and regulation of ART labs across the globe as well as special topics and emerging techniques and devices Chapter authors all experts in the field contribute their expertise from around the world With the addition of learning key points and review questions at the beginning and end of each chapter this new edition of In Vitro Fertilization is a readily accessible high quality instructional resource for reproductive medicine trainees at all levels Practicing reproductive endocrinologists urologists and embryologists also will find value in the book as will infertility researchers

The American Phonetic Dictionary of the English Language Daniel S. Smalley,1855 **QUANUM 3.0: An Updated Tool for Nuclear Medicine Audits** IAEA,2021-08-25 The Quality Management Audits in Nuclear Medicine QUANUM programme has proven to be applicable to many nuclear medicine services across a variety of economic circumstances It considers the diversity of nuclear medicine practices around the world and covers multidisciplinary contributions The present revision QUANUM 3 0 follows the principle of continuous quality improvement and reflects new scientific developments It draws on valuable lessons learned from more than a decade of global implementation of QUANUM with the assistance of experienced nuclear medicine professionals **Parenteral Quality Control** Michael J. Akers,Dana Morton Guazzo,1994

Pharmaceutical Quality Assurance Mr. Manohar A. Potdar,2006 *The David Bispham Song Book* David Bispham,1920
Daily Report, Foreign Radio Broadcasts United States. Central Intelligence Agency,1966 **Impact of Airline Deregulation** United States. Congress. Senate. Committee on Commerce, Science, and Transportation. Subcommittee on Aviation,1979 **Morbidity and Mortality Weekly Report** ,2004 **Establishing A CGMP Laboratory Audit System** David M. Bliesner,2006-04-20 The first systematic hands on auditing guide for today s pharmaceutical laboratories In today s litigious environment pharmaceutical laboratories are subject to ever stricter operational guidelines as mandated by the FDA and must be able to establish and demonstrate sustainable operational practices that ensure compliance with the current good manufacturing practice CGMP regulations David Bliesner s Establishing a CGMP Laboratory Audit System A Practical Guide is designed to provide laboratory supervisors and personnel with a step by step hands on audit system that they can rely on to ensure their facility remains compliant with all current and future requirements Focusing on a team approach the author uses detailed flowcharts checklists and descriptions of the auditing process to help readers develop a new audit system or upgrade their current system in order to Improve current compliance Demonstrate sustainable compliance Produce data for federal inspections Avoid regulatory action Enhanced with detailed checklists and a wealth of practical and flexible auditing tools on CD ROM this book provides an ideal resource for new and future laboratory personnel and an excellent means for keeping existing industry practitioners up to date on the nuances of operating a consistently compliant pharmaceutical laboratory **Pharmaceutical Microbiological Quality Assurance and Control** David Roesti,Marcel Goverde,2020-01-02 Relying on practical examples from the authors experience this book provides a thorough and modern approach to controlling and monitoring microbial contaminations during the manufacturing of non sterile pharmaceuticals

Offers a comprehensive guidance for non sterile pharmaceuticals microbiological QA QC Presents the latest developments in both regulatory expectations and technical advancements Provides guidance on statistical tools for risk assessment and trending of microbiological data Describes strategy and practical examples from the authors experience in globalized pharmaceutical companies and expert networks *Harper's Magazine* ,1890 Important American periodical dating back to 1850 **Enhancing compliance to good manufacturing practices and pharmaceutical quality system requirements in vaccine production** World Health Organization,2024-06-25

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