

|                                                                                   |         |                                                                       |  |           |      |
|-----------------------------------------------------------------------------------|---------|-----------------------------------------------------------------------|--|-----------|------|
|  |         | LABORATORIUM<br>MIKROBIOLOGI FARMASI<br>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

**Daniel S. Smalley**



## How To Apply For Laf Sop:

**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 evaluation     *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     **Polyglott**  
**Lexicon: Being a New Dictionary** ,1848     **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

**Pharmaceutical Quality Assurance** Mr. Manohar A. Potdar,2006 Establishing A CGMP Laboratory Audit System David M. Bliesner,2006-04-28 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

**Parenteral Quality Control** Michael J. Akers,1985 **Enhancing compliance to good manufacturing practices and pharmaceutical quality system requirements in vaccine production** World Health Organization,2024-06-25

**The International Code of Signals for Use of All Nations** Great Britain. Board of Trade,1926 **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.

**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.

**Polarization Dynamics of Mode-Locked Fiber Lasers** Sergey V. Sergeyev, Chengbo Mou, 2023-04-20. This book provides a comprehensive review of the latest research on the science, technology, and applications of mode-locked fiber lasers generating pulse trains with the evolving state of polarization at time scales ranging from a few pulse widths to 10,000 laser cavity round trip times. It supports readers with a timely source of information on the current, novel scientific concepts and cost-effective schematics, in addition to an overview of the feasible applications. The book aims to demonstrate for the nonlinear science community a newly emerging field of nonlinear science and so stimulates the development of new theoretical approaches and opens new horizons for the photonics community by pushing boundaries of the existing laser systems towards new applications. The new classes of optical sources and photonic devices explored in this book will be relevant with applications to other fields, including medicine, bio-photonics, metrology, and environmental safety.

**Key Features:** Provides a cutting-edge review of the latest emerging science, technology, and applications in the field. Tackles a topic with fast-growing interest in the USA, Europe, and China. Explores the simple and cheap design and tests of lasers and outlines the feasible applications.

*The American Phonetic Dictionary of the English Language* Daniel S. Smalley, 1855

**Daily Report, Foreign Radio Broadcasts** United States. Central Intelligence Agency, 1966

**The David Bispham Song Book** David Bispham, 1920

**Practical Handbook of**

**Pharmaceutical Analysis** Phoolsingh Yaduwanshi Reenu Yadav Prashanti Chitrapu, 2023-02-20 This book provides a systematic course of practical in Pharmaceutical analysis is a very sincere attempt to arouse the interest of the students in these fast developing branches of pharmaceutical sciences It gives concise and point wise information requiring during practical in single book and eliminates the need of too many reference book The subject matter has been explained in such a single way that the students should feel no difficulty to understand it The concepts as clear as crystal language simple and subject matter in flow and continuity the students will also discover the real pleasure of extra information All efforts have been made to make the book student friendly

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