Curriculum Vitae

Personal information

JOSEPH MURINZI

Work experience

Regulatory Officer: National Drug Authority-Uganda (Jan 2018 to date)
DUTIES AND RESPONSIBILITIES
- Assessing CTD dossiers for registration
- Assessing Medical devices applications for registration
- Assessing post regulation amendments (variations)
- Participate in drafting of SOPS and other guidelines in drugs and medical devices registration
- Participate in cGMP/IS0 13485 inspections of pharmaceutical and medical devices manufacturing facilities.

Production Officer: Cipla Quality Chemical Industries Limited-Uganda (June 2015 to Dec 2017)
DUTIES AND RESPONSIBILITIES
- Raising internal and purchase requisitions for critical supplies and equipment to stores and Procurement Departments
- Monitoring inventory of machine tools and accessories to mitigate mechanical breakdowns
- Providing monthly section Reports of overall Equipment Efficiency to management
- Manpower planning and allocation to ensure maximum output
- Ensuring appropriate validations, calibrations and qualifications are taken and recorded
- Maintaining up to date documentation including S.O.Ps, change control systems and procedures
- Conducting training on different aspects of cGMP
- Production planning and execution to meet delivery targets
- Participating in Self inspection audits

Technical Director/Supervising Pharmacist
- Bacton Pharmacy and Horris Pharmacy-Uganda (Jan 2016 to Dec 2017)
- M-Medius Pharmacy and Elite Pharmacy-Uganda (Jan 2015 to Dec 2015)
- A &S Pharmacy-Uganda (Jan 2014 to Dec 2014)

DUTIES AND RESPONSIBILITIES
- Pre-qualifying all suppliers of medicines to the pharmacy
- Approving the list of all medicines and supplies that can be purchased by the pharmacy.
- Promoting and Implementing Pharmacovigilance in the pharmacy
- Ensuring good storage and warehousing practices are followed
- Putting in place and monitoring systems to ensure adequate inventory stock and timely replenishment
- Developing S.O.Ps for various pharmacy operations
- Recruitment and training of Pharmacy auxiliary STAFF
- Handling all regulatory affairs with NDA and PSU
- Dispensing of medicine and providing Pharmaceutical Care to clients

Assistant Lecturer: Mbarara University of Science and Technology-Uganda (Jan 2014 to June 2015)
DUTIES AND RESPONSIBILITIES
- Conducting Lectures, Practical sessions, Presentations and tutorials
- Supervising Student research and projects
- Administering and grading students' coursework, tests and exams.
- Participating in planning and review meetings for the pharmacy department

Education and training

SEPT 2024-PRESENT POST BACCALAUREATE CERTIFICATE IN BIOTECHNOLOGY INNOVATION AND REGULATORY SCIENCES-PURDUE UNIVERSITY (USA)

AUG 2017-OCT 2024 MASTERS IN BUSINESS ADMINISTRATION AND MANAGEMENT-UGANDA MARTYRS UNIVERSITY (UGANDA)

APR 2022-OCT 2022 ADVANCED POST-GRADUATE DIPLOMA IN REGULATORY AFFAIRS-INSTITUTE OF PHARMACEUTICAL MANAGEMENT (INDIA)

2008-2012: BACHELOR OF PHARMACY-MBARARA UNIVERSITY OF SCIENCE AND TECHNOLOGY (UGANDA)

2006-2007: UGANDA ADVANCED CERTIFICATE OF EDUCATION (UACE)-NTARE SCHOOL (UGANDA)

2002-2005: UGANDA CERTIFICATE OF EDUCATION (UCE)-NTARE SCHOOL (UGANDA)

PROFESSIONAL TRAINING & CERTIFICATES
- ISO 13485 (April 2023)

Certificate of ISO 13485 Foundation
- USP (April 2023)

Certificate of completion of online training in Medical Devices Standards and Classifications: Focus on Maternal Neonatal and Child Health
- Pharmaceuticals and Medical Devices Agency (PMDA) (November 2022)

Certificate of participation in the PMDA-ATC Medical Devices Workshop 2022- Explanation of/insight into the IMDRF Documents
- Pharmaceuticals and Medical Devices Agency (PMDA) (August 2022)

Certificate of completion in the PMDA-ATC E-learning Training Course: Medical Devices Review
- African Vaccines Regulatory Forum (AVAREF) (July 2022)

Clinical Trials Inspection Training Course certificate
- USP (July 2022)

Certificate of completion of virtual training in Foundations of GMP
- Institute of Pharmaceutical Management-India (Jan 2019)

Certificate Training in Review of DMF' in reference to EMA guideline: 'Chemistry of active substances' and other relevant guidelines
- ISO 9001:2015

QMS Implementation course certificate
- European Medicines Agency (August 2018)

Online EU GCP Inspectors Basic Training Course certificate
- National Drug Authority (NDA) (July 2018)

Certificate of Good Manufacturing Practices (GMP) Training.
- Mulago National Referral Hospital Pharmacy department – (August 2012–August 2013)

Internship training in pharmaceutical care, drug handling, clinical pharmacy, dispensing, stock control and drug procurement.
- National Drug Authority (August 2012–November 2012)

3 months Internship training in the departments of, Drug assessment and registration and Drug Information where I gained hands on experience in the different aspects of drug regulation
- Mavid Pharmaceuticals limited – June 2011

Industrial training placement that focused on the different aspects of drug manufacture especially current good manufacturing processes
- Mbarara Regional Referral Hospital – June 2010

Community placement that provided training in inventory management, drug dispensing and pharmaceutical care.

Additional information

Publications

Projects

Memberships

Member of the Pharmaceutical Society of Uganda

Other Relevant Information