MUHAS School of Pharmacy, Dar es Salaam, Tanzania
Call for cGMP Training course Application
20-24th September 2021

Training at Pharmaceutical Analysis laboratory (PAL)

Our Capacity

Our consortium consists of experienced experts in the field:

Lead Coordinator:

Prof. Dr. Eliangiringa Kaale

Other Trainers

Staff from MUHAS, cGMP technical experts from Germany and Senior assessors and cGMP experts from TMDA with proven competence on GMP inspections.

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Our Vision:

- To become a Regional Centre of Excellence for practical training of pharmaceutical processionals and offering Medicine's valuation and registration, Good Manufacturing Practice (GMP), Formulation Development, Manufacturing Procedures, Scale-up Processes, Stability Testing, Analytical testing and Qualification and Validation services

Some activities in our scope include:

- To provide hands on training on dossier evaluation through joint reviews;
- To establish mechanisms for the countries in the region to move towards effective implementation of mutual policy framework and guidelines;
- To train and assist NMRA's and Pharmaceutical manufacturers to establish Quality Management System (QMS) for medicines regulatory affairs;

Tailor made courses can be offered onspecial request

Please contact us:

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CALL FOR cGMP TRAINING COURSE-1 APPLICATION

Who we are:

- MUHAS Pharm R&D laboratory was established in 2009 as a neutral ground area to support local pharmaceutical industries in terms of Good Manufacturing Practice and formulation development of Pharmaceuticals.
- We are also in a consortium for Regional Centre of Regulatory Excellency (RCORE) in Medicines Evaluation and Registration. The consortium is made up of the School of Pharmacy, Muhimbili University of Health and Allied Sciences (MUHAS) and Tanzania Medicines and Medical Devices Authority (TMDA).
- We are also an African proficiency testing providers for Pharmaceuticals. Hence, we prepare PT matrices and test performance of peer laboratories in the region.
- A proficiency test is a tool for continuous improvement in laboratory operations, staff training and orientation. The PT results provide information on the validation of analytical methods and an assessment of measurement uncertainties.
- We are currently the only ISO 17043:2010 accredited laboratory providing PT schemes for pharmaceutical testing labs in the East African Community (EAC).

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UPCOMING TRAINING COURSE

Title: cGMP Training

Date: 20-24th September 2021

Place: Dar-es-Salaam, Tanzania.

Description: The trainees will be taken into 3 days didactic training and 2 days hands on training with experienced current GMP experts from Germany and Tanzania.

Scope: On site cGMP training with hands on and field tour activities

Participation: 1000 USD per person including: breakfast, lunch, evening tea and training materials.
Excluding: Travels and Daily Subsistence Allowance.
Please confirm your participation before 03.09.2021 through: ptibalinda@gmail.com

Participants can apply to action medeor, for a **scholarship** by sending a resume and a motivation letter through: pharmazie@medeor.de before 03.09.2021.
Please note that the number of scholarships is limited and will cover the participation fee.

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Why should you take our cGMP training course?

- Possibility of hands-on training from Pharmaceutical Technology lab and Pharmaceutical Analysis laboratory.
- International collaborative trainings. The upcoming training will have experts from Germany.
- Multidisciplinary trainers i.e., MUHAS-academia, TMDA-regulator and Action-medeor-International experts from industrial field.
- Field and tours to the pharmaceutical industries for consolidation of theoretical knowledge.
- We offer accredited courses which can be used as an evidence for continuing professional development (CPD).
- Our training can be tailormade based on needs.
- We have a long experience in providing short courses.
- Our training materials have been developed in line with the WHO cGMP requirements.

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Our full cGMP course includes the following topics

1. Pharmaceutical Quality System (Risk Management) *
2. cGMP for Pharmaceutical Product*
3. Sanitation and hygiene
4. Qualification and Validation*
5. Handling of Complaints
6. Product recalls
7. Contract production, analysis and other activities
8. Self-inspection, Quality Audits, suppliers audit and approval
9. Personnel
10. Training
11. Personal hygiene
12. Premise
13. Equipment*
14. Materials for cGMP*
15. Documentation*
16. Good practices in production
17. Good practices in quality control

* Topics will be addressed during the upcoming training session (20-24th September 2021)