MUHAS School of Pharmacy
Dar Es Salaam
CALL FOR TRAINING APPLICATION
Medicines Evaluation and Registration
20th to 24th January 2020

Our Main Goals:

- To ensure that the public accesses medicines that meet acceptable standards in terms of quality, safety and efficacy and that they are manufactured in facilities that meet acceptable standards of Good Manufacturing Practices (GMP).
- To address the problem of scarcity of skilled human resources in regulation of medicines, the MUHAS-TMDA RCORE has in place short training programs to be conducted twice a year, and a long training plan for a Masters course in Regulatory Sciences.

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We can undertake the following assignments:

- 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 other NMRA's to establish Quality Management System (QMS) for medicines evaluation and registration;
- To help the countries in the region to move towards effective implementation of medicine regulation harmonization;
- To help the countries in the region to establish framework for information sharing and move towards mutual recognition and
- To assist the region to move towards establishment of a single African Medicines Agency (AMA).

→Tailor made courses can be offered on special request.

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Our Capacity

Our consortium consists of experienced experts in the field:

Lead Coordinator:

- Prof. Dr. Eliangiringa Kaale, Professor of Quality Assurance with vast experience in the field of medicine Quality Control and Quality Assurance, medicines evaluation and registration and Pharmaceutical product Development and Technology Transfer.

Other staff

- Staff from MUHAS and Senior assessors from TMDA with proven competence on the task at hand.

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UPCOMING TRAINING COURSE

Title: Medicines Evaluation and Registration

Date: 20th to 26th January 2020

Place: Dar-es-Salaam, Tanzania.

Description: The trainees will be taken into 3 days didactic training and 2 days hands on training with experienced assessors.

Scope: Common Technical Document for the Registration of Pharmaceuticals for Human Use - Quality

Participation fee: 600 USD per person

Including: Lunch + teas and training materials. Excluding: Travels and Daily Subsistence Allowance. (selected candidates will be linked to potential sponsor)

Please confirm your participation through: Sunday.kisoma@tfda.go.tz/ ptibalinda@gmail.com

Deadline for confirmation: 10th January 2020

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- To serve as training centre for in-service training of the regulatory staff on Medicine Evaluation and Registration (MER) and Good Manufacturing Practice (GMP);

Please contact us:
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Who we are:

- SoP-MUHAS/TMDA Regional Centre of Regulatory Excellency (RCORE) in Medicines Evaluation and Registration is a consortium made up of the School of Pharmacy, Muhimbili University of Health and Allied Sciences (MUHAS) and Tanzania Medicines and Medical Devices Authority (TMDA).
- This RCORE has been established and received a formal designation by NEPAD African Medicine Registration Harmonization Initiative as an RCORE in medicines Evaluation and registration.

Our Mission:

- To assist National Medicines Regulatory Authorities (NMRAs) in the region to build up national and regional capacity in pre-approval scientific evaluation of medicines

Our Vision:

- To become a Regional fountain of knowledge for medicines evaluation and registration.

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