

2019 Seminar on African Regulatory Authorities - Emerging markets

Presentations to be conducted by representatives from the Regulatory Authorities:
 Ghana FDA, Kenya PPB, Nigeria NAFDAC, Uganda NDA & Eastern African Community Medicines Regulatory Harmonization (EAC MRH) Programme

Prospective dates: 25 - 26 February 2019 |  Altron Bytes Conference Centre, Midrand

A First for South Africa: Open to All Individuals



Mrs Delese Afia Amoakoa Mimi Darko nee Osei-Bonsu
 Chief Executive Ghana Food and Drugs Board



Mr Dominic Munyoroku Kariuki
 Directorate of products evaluation & registration at Pharmacy and Poisons Board in Kenya



Dr Mojisola Christianah Adeyeye
 Director-General at National Agency for Food and Drug Administration and Control in Nigeria



Ms Donna Asimwe Kusemererwa
 Secretary to the Authority of National Drug Authority in Uganda



Mr Mwesigye John Patrick
 Senior Health Officer at East African Community responsible for Medicines Regulatory Harmonization

Content to be discussed at this seminar:

- Requirements for product registration: Medicine/ medical device/ complementary medicines dossier requirements
- Relevant guidelines
- Fees required
- Samples
- Labelling requirements and languages
- Post importation: Laboratory testing of product/batches - where, when and how often?
- Pharmacovigilance
- GXP inspections
- Pre- & post-registration variations
- Harmonisation initiatives and processes to register products under joint procedures
- Tender market/import/export - Regulatory dossier requirements
- Counterfeit management
- **Includes: Interactive question and answer sessions**



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Why you should attend?

- Queries regarding product entrance into these emerging markets will be addressed.
- Brush shoulders with esteemed experienced Regulatory Authority professionals who are directly involved in product evaluation for registration.
- Capitalise on the opportunity to address the speakers directly with our interactive Q&A sessions.
- Each delegate will receive a course pack containing comprehensive documentation which will be a valuable source of reference.

Who should attend?

This seminar will be of interest to pharmaceutical, medical device and complementary medicine companies and individuals who need to learn about successful product applications and registration in Ghana, Kenya, Nigeria, Uganda & understand the EAC MRH Programme.

Date and Venue:

Prospective dates: 25 - 26 February 2019
Day 1 & day 2: 7:30 to 17:00

Altron Bytes Conference Centre
Block C, 241 3rd Rd, Halfway Gardens
Midrand, South Africa, 1686

Each delegate will receive a certificate of attendance

All interested parties to complete registration forms only, by 14 December 2018, in order for Twinz to secure relevant speakers from the various Regulatory Authorities. Visit www.twinztraining.com to register for seminar and more information.

**BOOK NOW AND SAVE 10 %
LIMITED SEATS AVAILABLE**

Cost of the seminar:

Price per delegate for both days:
R 9999,00 (incl. VAT)

Early bird special:
Less 10 % of the total fee (book & pay before 07 December 2018)

Price includes lunch and refreshments for both days

Steps to register:

- 1) Log onto the website: www.twinztraining.com
- 2) Go to the Events Booking page.
- 3) Click the "register" button, choose the number of seats you want (delegates can only book more than 1 seat if from the same company), and then click the "checkout" button.
- 4) Enter personal information and then click the "confirm & proceed" button.
- 5) You will then be directed to a page reflecting the Twinz Foundation banking details (note that payments can only be processed through EFT and once you receive an invoice).
- 6) Click the "buy now" button to **finalise registration**.
- 7) Once you have completed steps 1 to 6, you will receive, via email, a booking confirmation ticket and shortly thereafter an invoice.
- 8) Once you make payment, you will receive a valid ticket attendance.

