JULY
Training Plan

CENTER FOR CONTINUING PROFESSIONAL DEVELOPMENT
مركز التطوير المهني المستمر

EGYPTIAN DRUG AUTHORITY
هيئة الدواء المصرية

Training plan for July 2021

Central Administration of Pharmaceutical Products

- Overview On Submission And Evaluation Of Module 3 Of The CTD File And Overview On Submission And Evaluation Of Bioequivalence And Comparative In-vitro Dissolution Studies
  10,11 JULY | 12 HOURS / 2 DAYS | 2000 EGP
- Validation, Verification & Methods transfer of analytical procedure according to ICH Q2 & Pharmacopeial requirements.
  25,26 JULY | 8 HOURS / 2 DAYS | 3000 EGP

Central Administration of Pharmaceutical Care

- Marketing Materials & Media Monitoring (Process Of Files Submission)
  8 JULY | 4 HOURS / 1 DAY | 1000 EGP
- Marketing Materials & Media Monitoring (PROMAT)
  28 JULY | 4 HOURS / 1 DAY | 1000 EGP
- PV Condensed Course For Pharmaceutical Companies
  START FROM 25 JULY | 25 HOURS / 5 DAYS | 4000 EGP

Central Administration of Medical Devices

- Best Practice For Importation Of Dental Products
  14 JULY | 3 HOURS / 1 DAY | 1000 EGP
- Best Practice For Importation Of IVD Products
  13 JULY | 3 HOURS / 1 DAY | 1000 EGP

Central Administration of Operations

- Training Program For Licensing Bioequivalence Centers
  14JULY | 4 HOURS / 1 DAY | 1000 EGP
- Training Program For Licensing Stability Centers
  15JULY | 4 HOURS / 1 DAY | 1000 EGP
- GMP For Medical Devices Factories
  27-29JULY | 12 HOURS / 3 DAYS | 2000 EGP

Central Administration of Pharmaceutical Policies & Market Access

- General Rules Regulating Prices Process
  11,14 JULY | 8 HOURS / 2 DAYS | 3000 EGP

Contact us at :

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Egyptian Drug Authority

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AUGUST
Training Plan

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Training plan for August 2021

Central Administration of Pharmaceutical Products

- Antiseptics And Disinfectants Products`File Submission workshop
  11 AUGUST — 5 HOURS / 1 DAY — 1500 EGP
- Pesticides File Submission workshop
  18 AUGUST — 5 HOURS / 1 DAY — 1500 EGP
- Biocides Variation Guidelines
  25 AUGUST — 4 HOURS / 1 DAY — 1500 EGP
- Workshop On Registration Of Herbal Medicine
  19 AUGUST — 5 HOURS / 1 DAY — 2000 EGP
- Dissolution Method Development, comparative study & Quality control testing according to Pharmacopeial & International regulatory requirements for successful CTD Submission
  30-31 AUGUST — 12 HOURS / 2 DAYS — 4000 EGP

Central Administration of Pharmaceutical Care

- Medical Devices Vigilance Training Program
  29-31 AUGUST — 12 HOURS / 3 DAYS — 3000 EGP

Central Administration of Operations

- Advanced training program for pharmaceutical stores
  15-17,22-23 AUGUST — 20 HOURS / 5 DAYS — 4000 EGP
- Essential requirements should be provided by MD Importers for Inspection & (GSDP)
  11-12 AUGUST — 8 HOURS / 2 DAYS — 1500 EGP
- Risk assessment & Risk management Program
  17 AUGUST — 4 HOURS / 1 DAY — 1000 EGP
- Basic GMP
  8 ,10,12,16,18,22,24,26 AUGUST — 32 HOURS / 8 DAYS — 5000 EGP
  3000 for Students

Central Administration of Medical Devices

- Best Practice For Variation Of Registered Medical Devices
  1 AUGUST — 3 HOURS / 1 DAY — 1000 EGP
- Best Practice For Approval Of Scientific Committee
  15 AUGUST — 3 HOURS / 1 DAY — 1000 EGP
- Best Practice For registration of imported medical devices
  9 AUGUST — 6 HOURS / 1 DAY — 1000 EGP

Central Administration of Pharmaceutical Policies & Market Access

- Guidelines and regulation for Medical Device & Kits export
  26 AUGUST — 5 HOURS / 1 DAY — 1200 EGP
- Pharmacoeconomics unit training program (INTERMEDIATE- LEVEL)
  STARTS 28 AUGUST — 30 HOURS / 5 DAYS — 7500 EGP
- General Rules Regulating Prices Process
  15,18 AUGUST — 8 HOURS / 2 DAYS — 3000 EGP

Central Administration of Biological &Innovative Products &Clinical Studies

- Biological Registration Ministerial Decree 297/2009 Updates Workshop
  29 AUGUST — 6 HOURS / 1 DAY — 1500 EGP

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SEPTEMBER
Training Plan

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Training plan for September 2021

Central Administration of Pharmaceutical Products

- Successful filing of DMF of Active pharmaceutical ingredient (API) for CTDaccording to ICH &WHO guidelines
  26-27 SEPT. 12HOURS / 2 DAYS 3000 EGP

Central Administration of Pharmaceutical Care

- Marketing Materials & Media Monitoring (PROMAT)
  16SEPT. 4 HOURS / 1 DAY 1000 EGP
- Pharmacovigilance Awareness and Practices
  Starting from 22 SEPT. 35 HOURS / 7 DAYS 5000EGP

Central Administration of Operations

- Licensing pharmaceutical products factories
  5-8 SEPT. 16 HOURS / 4 DAYS 3000 EGP
- Advanced training program for scientific office
  21-23 SEPT. 12 HOURS / 3 DAYS 2500 EGP
- Process performance life cycle from statistical & practical perspective
  14 SEPT. 4 HOURS / 1 DAY 1000 EGP
- GMP requirements for medical devices plant design
  13 SEPT. 4 HOURS / 1 DAY 1000 EGP

Central Administration of Medical Devices

- Best practice for Stability & Biocompatibility Committee
  15 SEPT. 3 HOURS / 1 DAY 1000 EGP
- Best practice for registration of locally manufactured medical devices
  1 SEPT. 3 HOURS / 1 DAY 1000 EGP

Central Administration of Pharmaceutical Policies & Market Access

- Export Automation Program
  12 SEPT. 6 HOURS / 1 DAY 1200 EGP
- General Rules Regulating Prices Process
  19,22 SEPT. 8 HOURS / 2 DAYS 3000 EGP

Central Administration of Biological &Innovative Products &Clinical Studies

- Emergency Use Authorization Guideline In Arab Republic Of Egypt Workshop 1500 EGP
  28 SEPT. 4 HOURS / 1 DAY

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