



## **NOVEMBER 2022**

## **TRAINING PLAN**

CENTRAL ADMINISTRATION OF Biological & Innovative Products	<ul> <li>Workshop on Test Method Validation</li> <li>29 &amp; 30 November 2022</li> </ul>	3000 EGP
CENTRAL ADMINISTRATION OF Pharmaceutical Products	<ul> <li>Workshop on Preparation and Submission of Scientific reference Products or Internally Transferred Files to Ur Data Evaluation &amp; Drug Development 06 November 2022</li> <li>Workshop on submission procedures of antiseptics &amp; d registration files and requirements of variations 20 November 2022</li> </ul>	it of Scientific 1500 EGP
CENTRAL ADMINISTRATION OF Medical Devices	<ul> <li>Workshop on Best Practice for Importation of Medical I 10 November 2022</li> <li>Workshop on Best Practice for Importation of Medical E 13 November 2022</li> <li>Workshop on Best Practice for Registration of Local M Without Quality Certificates 16 November 2022</li> </ul>	1000 EGP Equipment 1000 EGP
CENTRAL ADMINISTRATION OF Pharmaceutical Care	<ul> <li>Workshop on Medical Insert submission guidance 24 November 2022</li> <li>Webinar on Marketing Materials &amp; Media Monitoring (Pu Submission) 24 November 2022</li> <li>Webinar on follow up &amp; Updates of Online Ads 27 November 2022</li> <li>Training Program on Medical Devices Vigilance 27-29 November 2022</li> </ul>	1000 EGP focess of files 1000 EGP 1000 EGP 3000 EGP