أجندة ورشة عمل تحديث اليات فحص ملفات تحليل المستحضرات الصيدلية ومراجعة نتائج التحليل بمعامل الإدارة المركزية للرقابة الدوائية.

**Updated guidelines for reviewing pharmaceutical Products analysis files and Out of Specs (OOS) results in Central administration of drug Control (CADC) Laboratories**

| Topic | Lecturer | Date | Time |
|---|---|---|---|
| **Registration** | | Tuesday 30/3/2021 | 9:00-9:30 |
| Welcome Address | Prof. Dr. Tamer Essam<br>Prof. Dr. Aiman Elkhatib<br>Prof. Dr. Medhat Al- Ghobashy | | 9:30-10:00 |
| Guidelines For Preparation and Assessment Of finished Pharmaceutical Products Analysis File | Dr. Hamada Gamal<br>Dr. Wedian Younes | | 10:00-11:00 |
| Panel Discussion (part 1) | Dr. Rania Abdel Bassit<br>Dr. Dalia El Khatib<br>Dr. Loai Mohamed | | 11:00 – 12:00 |
| **Coffee Break** | | | 12:00-12:30 |
| Panel Discussion (part 2) | Dr. Rania Abdel Bassit<br>Dr. Dalia El Khatib<br>Dr. Mohamed Metwaly | | 12:30-2:30 |
| **Coffee Break** | | | 2:30-3:00 |
| Commonly Encountered Problems of OOS (Chemical-Physical-Microbiology) | Dr. Marwa Hassan | | 3:00-3:30 |
| Panel Discussion | Dr. Sara Hashem<br>Dr. Noran Hamed<br>Dr. Heba Yehia | | 3:30- 4:30 |