dcvmn	**Preparation for DCVMN eWorkshops**

Thank you for registering for DCVMN's e-Workshop:

Regulatory Part II – CRP, EUL and CTD
30 September – 1 October 2020

Ahead of the workshop, please complete the following actions:

1. **Pre-reading** https://dcvmn.net/e-Workshop-Regulatory-Pathways-Part-II
   - Please **read the two documents** on the e-workshop site indicated above.

WHO Emergency Use Listing Procedure v8 January 2020
WHO TRS 996 Annex 08 2016 on the collaborative review procedure

2. **Moodle e-Learning** https://moodle.dcvmn.net/
   - Please **complete at least one of the e-learning courses** on the Moodle site indicated above.
   - We recommend one or both of the following courses:

Post Approval Changes in Biologics Manufacturing
Common Technical Documents (CTD)

3. **Proof of e-Learning**
   - Take the online test(s) for the Moodle courses above
   - Email the completed certificate to m.dennehy@dcvmn.net. with the email header "Regulatory Part II e-workshop preparation" before **Monday 28 September**.

We look forward to connecting with you at the event.

Regards

Sonia, Maureen and the rest of the organizing team.
16 September 2020