Before opening any attachments, a copy of the EMWA Privacy policy can be found. It cannot be guaranteed that it is completely free from such problems and we do not accept any liability for loss.

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Photo Contest
Get inspired! Bring your EMWA join to Vienna and capture the essence!

Show It, Share IT Session
We are looking for volunteers to participate in the Show It, Share IT session. If you want to share IT you can do your daily work, send the images to us.

You can also voluntary to join the session to update you staff with new IT.

5th EMWA Expert Seminar Series
Don’t miss the PV ESS on Wednesday 08 May - 13:00 - 16:00
This is a fantastic opportunity for medical writers interested in PV. Dr. David Lewis (Novartis) will explain the challenges in writing RMPs and PBRERs and Doctors Steph Millican and Janet Nooney (expert reviewers from the UK regulator) will describe the common mistakes they see, what they expect from the medical writers and their thoughts on how we can improve our performance. There will also be an opportunity for the audience to ask questions. Bring your questions and learn how to write these important documents and how to master them to a regulator’s dream!

Veterinary Medical Writing Special Interest Group Meeting
There will be a non-official dinner on Wednesday evening in case anybody is interested.

Please send an email to: non-workshop@emwa.org to book your place. You will be responsible for paying the restaurant for your own meal and drinks.

EMWA Newsblast - April 2019

EMA updated Q&A on article 20 non-pharmacovigilance procedures
EMA updated Q&A on article 31 pharmacovigilance referrals
New safety features for medicines sold in the EU against falsified medicines
PSMF” for no deal Brexit scenario
UK Health Authority MHRA published guidance on “UK QPPV” and “UK Public Registry Posting Procedures” for medicinal products

PhUSE White Paper
Clinical trials
No-deal Brexit Scenario: Guidance on UK public registry postings for medicinal products
Health Canada’s Final Guidance on “Public release of clinical trials information” 12 March 2019

NEWS OF INTEREST

Regulatory Public Disclosure (RPD) Regulatory News
· Health Canada Final Guidance on “Public release of clinical information” 12 March 2019
· Nonclinical Brand Summary Guidance on UFM public registry for clinical trials
· PICS Newsletter

Regulatory Update on Regulatory News
· UK medicines authority MHRA published guidance on UK (GMP) and UK (GDP) for non-clinical stress scenarios
· EMA guidance for monitors involved in the 4010a (GCP) inspection process
· EMA updated Q&A on subject 23 non-pharmacovigilance procedures

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