The Ambassador’s program group will be having a lunch time roundtable talk on the 27th of February at the Imperial College London during their Focus skills” and the role of EMWA training as part of the Health and Life Sciences candidates, many of whom had never heard of the field of medical this program are interested in finding out more about this program. The Ambassador’s program is an example of how to interact with candidates, many of whom had never heard of the field of medical

Alison Rapley will be holding an interactive course on Improving your writing communication before.

Pharmacology in Buch, Berlin. This was well received by over 30 PhD EMWA at the Winter School career event at the Leibniz Institute of Molecular EMWA President Abe Shevack has given a talk on medical communication and regulations and guidelines, the new requirements and documents, and the impact of parallel regulatory-HTA scientific advice on clinical development. Assessing the uptake of regulatory and HTA recommendations.

Strengthening standards, transparency, and collaboration to support medicine prevention of diabetes mellitus - Revision 2. 2) The European Medicines Agency has released for public consultation a PRAC

3) The EMA has revised the Annex I to the GMP Annex 20, adding a new section on the prevention of diabetes mellitus, Revision 2. 4) Following the safety recommendation and the public hearing on valproate, the EMA has issued a draft guideline on the use of valproate in pregnant women and children. 5) The Ukraine now accepts RMPs written according the EU format as per GVP

6) The EMA has published a question-and-answer document addressing the requirements for the preparation of RMPs. For more information about orphan designation please click here. Are you interested in contributing an article, short text, or news item related to one of these themes or any other topic relevant to medical writing? Please contact our editorial office at mew@emwa.org. If you are interested in contributing an article, short text, or news item related to one of these themes or any other topic relevant to medical writing, please contact our editorial office at mew@emwa.org.

Upcoming issues of Medical Writing

June 2019: Generics and biosimilars (deadline March 11, 2018)
March 2019: Careers in medical writing (deadline December 10, 2018)
January 2019: Training Standards in the World of Medical Writing (deadline October 10, 2018)

Regulatory updates

•    Medical journalism
•    General Data Protection Regulation and Regulatory update: Clinical Trial
•    Orphan drugs and rare disorders
•    Pharmacovigilance

This year, EMWA’s ESS comprises a range of topics led by international experts. The topics are:

EMWA Expert Seminar Series

•    Medical journalism
•    General Data Protection Regulation and Regulatory update: Clinical Trial
•    Orphan drugs and rare disorders
•    Pharmacovigilance

In this environment, the role of medical communicators in the medical device industry has gained in importance. In this context, the regulatory aspects of medical device development and the role of medical writers have become critical. The role of medical communicators in the medical device industry has gained in importance. In this context, the regulatory aspects of medical device development and the role of medical writers have become critical.

This webinar provides regulatory medical writers a short introduction to the key aspects of orphan medicines. The Q&A document is available online and can be downloaded from the EMWA website.

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The webinar will cover the following topics:

1) The European Medicines Agency has raised several issues associated with the preparation of RMPs, including the prevention of diabetes mellitus, Revision 2. 2) The European Medicines Agency has released for public consultation a PRAC

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