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Video on ICH Guideline Development Process

EICF Implementation Guide

Transparency and Vaccine Hesitancy

Call for Publication of Unredacted CSRs

Support economic operators in implementing the obligations and requirements of the Medical Devices Regulation EU 2017/745 and in vitro diagnostic medical devices (IVDR), which become applicable from 26 May 2022.

The UDI system enhances the identification of medical devices and makes it easier for health care professionals and patients to trace the origin and history of medical devices in order to improve safety and quality.

Regulations (EU) 2017/745 on medical devices (MDR) and (EU) 2017/746 on in vitro diagnostic medical devices (IVDR), which become applicable from 26 May 2022.

Principles for clinical trials' estimands and sensitivity analysis in clinical trials to the guideline on statistical principles for clinical trials.'

Important conversations around topics that will impact our industry in the future with the EMWA Special Interest Groups.

This is achieved through the EMWA journal, EMWA website, and events at a career level. It can also be achieved through mentorships, where experts provide guidance to emerging professionals.

Meet&Share Session with the MedComms-SIG

Support the public about data integrity and can stimulate research and development.

Provide information on T&D requirements.

Medical writers can better support economic operators in implementing the obligations and requirements of the Medical Devices Regulation EU 2017/745 and in vitro diagnostic medical devices (IVDR), which become applicable from 26 May 2022.

Participation of international networks provides a unique opportunity for medical writers to share their experiences and best practices with colleagues from around the world. This sharing not only helps to improve the quality of medical writing but also fosters a sense of community and collaboration.

The IMWEDSIG proposal needs 5 putative sub-committee members:

- 50+% of Industry executives anticipate large-scale AI use in healthcare by 2025

- Global biotechnology sector market size was $449.06 billion in 2019

- Molecular diagnostics market size is likely to exceed $25.2 billion by 2025

- Robotics/AI experience

- In vitro/molecular diagnostic regulatory writing and medical communication experience

- ATMP regulatory writing and medical communication experience

- Additional experience in the area of medical writing

- Educate and inform EMWA members about this important and expanding area of medical writing

- Work with medical writers from organisations focused on engagement with the public to help to highlight the importance and value of trained medical writers to do this work

- Support the need and value-add that medical writers can bring to this area of medical writing

- Network with others in the field

- PR team is looking for some volunteers to help with posting in social media, and preparing this Newsblast. If you want to be part of the team, email info@emwa.org and we will add it to the agenda. Please also email any other suggestions or requests for future publications. Any other challenges you would like to discuss? Please email info@emwa.org to receive the Zoom link.

- Let's Meet Up in Switzerland

- Call for volunteers

- EMWA Newsletters - June 2021

- Member Newsletters - June 2021

- Call for EMWA Fellows

- Nick Thompson Fellowship Award

- EMWA Newsblast - June 2021