cannot guarantee that it is completely free from such problems and we do not accept any liability for loss
mail in error, please immediately return it to the sender and delete it from your system. This e-mail,
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The European Parliament and Council confirmed the
(IVDR) in May 2022
Adoption of the EU In Vitro Diagnostic Regulation
reactions as a result of treatment with the corresponding medicinal product.
(a) identify, before and/or during treatment, patients who are most likely to
benefit from the corresponding medicinal product; or (b) identify, before and/or
Notified Body on Companion Diagnostics
medical device (integral, co-packaged or separately-obtained and referenced in
January 2022. This guideline focuses on product-specific quality aspects of a
CHMP in July 2021, the guidance document is now called
Updates from the EU MDCG
Programme Module 3
Sponsor Handbook
EMA published
Forum aims to facilitate collaboration and co-creation of real solutions for
personalised insurance plans for businesses and individuals in the life science,
a tailored range of specialist insurance products for both individuals and
place at the EMWA Spring Conference in May 2022.
Voting details will be sent out electronically in April 2022 and the successful
include a picture of yourself embedded into the Word document, to
If you would like to apply for any of the roles please provide a candidate
challenges and benefits of writing for the public.
Journal Announcement
Website.
EMWA Newsblast - January 2022

יקים: "NEW" או "Rev." עם התאריך המתאים.

On 24 Nov 2021, EMA published their
Devices Shortages Steering Groups. The groups will manage the
medicines and medical devices
agreed to make the EMA more effective in tackling shortages of
marked by "NEW" or "Rev." with the relevant date upon publication.

UK Parliament has been urged
Edinburgh University Guest Blog

Quick Guide – User Access Management – CTIS Training
Module 11 – How to

This perspective paper published in the same journal titled ‘
On 24 Nov 2021, EMA published their
Transform Research Data Sharing One Plenary at a Time

Recent amendments to the Federal Food, Drug, and Cosmetic Act (FD&C
manufacturer — Part 1: General requirements
The use of
symbols helps simplify medical device labels in Europe (considering the
recent Research Data Alliance (RDA) open Plenary on open sharing and

E-mail updates, Research Data Alliance, Medical Device Labeling, Protocol

**Research Data Alliance (RDA)**

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**Research Data Alliance (RDA)**