




| | |
|---|-----------|
| HUMANARZNEIMITTEL - EMA | 2 |
| <i>Allgemeines – General</i> | 2 |
| <i>Pharmakovigilanz – PRAC</i> | 3 |
| <i>Zulassung – Regulatory Affairs</i> | 4 |
| <i>Orphan Drugs und neuartige Therapierichtungen (ATMP)</i> | 5 |
| <i>Qualität – Quality</i> | 5 |
| <i>(Prä-) Klinische Forschung – Research and Development</i> | 5 |
| <i>Kinderarzneimittel – Paediatrics</i> | 6 |
| <i>Pflanzliche Arzneimittel – Herbal medicines</i> | 6 |
| EUROPEAN COMMISSION | 8 |
| EDQM | 9 |
| MEDIZINPRODUKTE | 10 |
| CMDH | 11 |
| HUMANARZNEIMITTEL - DEUTSCHLAND | 12 |
| HUMANARZNEIMITTEL - ÖSTERREICH | 15 |
| HUMANARZNEIMITTEL - SCHWEIZ | 16 |
|  FRAGEN AN DAS NETZWERK | 17 |
| VERANSTALTUNGEN / EVENTS – BEHÖRDEN UND ANDERE VERANSTALTER | 18 |
| DEUTSCHLAND | 18 |
| ÖSTERREICH | 18 |
| SCHWEIZ | 18 |
| EUROPA | 18 |
| <u>Urheberrechtshinweis:</u> | |

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Allgemeines – General

Minutes of the 122nd meeting of the Management Board, 13-14 December 2023

Published on: 12 - April - 2024

For more information, please refer to:

https://www.ema.europa.eu/en/documents/minutes/minutes-122nd-meeting-management-board-13-14-december-2023_en.pdf

Big data

Published on: 12 - April - 2024

For more information, please refer to:

<https://www.ema.europa.eu/en/about-us/how-we-work/big-data>

Meeting of the Medicine Shortages Single Point of Contact (SPOC) Working Party, Event - 14 - February - 2024 (updated)

Meeting summary available

Published on: 15 - April - 2024

For more information, please refer to:

<https://www.ema.europa.eu/en/events/meeting-medicine-shortages-single-point-contact-spoc-working-party-12>

European Shortages Monitoring Platform

Published on: 18 - April - 2024

For more information, please refer to:

<https://www.ema.europa.eu/en/human-regulatory-overview/post-authorisation/medicine-shortages-availability-issues/european-shortages-monitoring-platform>

Meeting of the Executive Steering Group on Shortages and Safety of Medicinal Products (MSSG), Event - 19 - March - 2024 (updated)

Minutes available

Published on: 18 - April - 2024

For more information, please refer to:

<https://www.ema.europa.eu/en/events/meeting-executive-steering-group-shortages-safety-medicinal-products-mssg-march-2024>

Organisation chart: Stakeholders and Communication

Published on: 16 - April - 2024

For more information, please refer to:

https://www.ema.europa.eu/en/documents/other/organisation-chart-stakeholders-communication_en.pdf

Organisation chart: Advisory functions

Published on: 16 - April - 2024

For more information, please refer to:

https://www.ema.europa.eu/en/documents/other/organisation-chart-advisory-functions_en.pdf

Humanarzneimittel - EMA

Medicinal products for human use: monthly figures - March 2024

Published on: 17 - April - 2024

For more information, please refer to:

https://www.ema.europa.eu/en/documents/report/medicinal-products-human-use-monthly-figures-march-2024_en.pdf

EMA multi-stakeholder workshop on psychedelics – Towards an EU regulatory framework, Event – 16 to 17 - April - 2024 (updated)

Presentations available

Published on: 19 - April - 2024

For more information, please refer to:

<https://www.ema.europa.eu/en/events/ema-multi-stakeholder-workshop-psychedelics-towards-eu-regulatory-framework>

Executive Steering Group on Shortages and Safety of Medicinal Products (updated)

Published on: 23 - April - 2024

For more information, please refer to:

<https://www.ema.europa.eu/en/about-us/what-we-do/crisis-preparedness-management/executive-steering-group-shortages-medicinal-products>

List of eligible industry stakeholder organisations (updated)

Published on: 24 - April - 2024

For more information, please refer to:

https://www.ema.europa.eu/en/documents/other/list-eligible-industry-stakeholder-organisations_en.pdf

Guide on access to unpublished documents (updated)

Published on: 25 - April - 2024

For more information, please refer to:

https://www.ema.europa.eu/en/documents/other/guide-access-unpublished-documents_en.pdf

SME Regulation and reports

Published on: 26 - April - 2024

For more information, please refer to:

<https://www.ema.europa.eu/en/about-us/support-smes/sme-regulation-reports>

Pharmakovigilanz – PRAC

List of medicines under additional monitoring

Published on: 22 - April - 2024

For more information, please refer to:

<https://www.ema.europa.eu/en/human-regulatory-overview/post-authorisation/pharmacovigilance-post-authorisation/medicines-under-additional-monitoring/list-medicines-under-additional-monitoring>

Guidance on real-world evidence provided by EMA: support for regulatory decision-making

Published on: 12 - April - 2024

For more information, please refer to:

https://www.ema.europa.eu/en/documents/other/guidance-real-world-evidence-provided-ema-support-regulatory-decision-making_en.pdf

International Coalition of Medicines Regulatory Authorities (ICMRA)

Published on: 17 - April - 2024

For more information, please refer to:

<https://www.ema.europa.eu/en/partners-networks/international-activities/multilateral-coalitions-initiatives/international-coalition-medicines-regulatory-authorities-icmra>

SPOR Status Update, Event – 10 - April - 2024 (updated)

Video recording available

Published on: 25 - April - 2024

For more information, please refer to:

<https://www.ema.europa.eu/en/events/spor-status-update>

Call for proposals - Grant procedure EMA/GRANT/2024/02/IA “Medicines regulatory systems strengthening in Sub-Saharan Africa”

Published on: 19 - April - 2024

For more information, please refer to:

https://www.ema.europa.eu/en/documents/procurement/call-proposals-grant-procedure-ema-grant-2024-02-ia-medicines-regulatory-systems-strengthening-sub-saharan-africa_en.pdf

Product Management Service (PMS) Info-Day, Event - 16 - April - 2024

Presentation available

Published on: 19 - April - 2024

For more information, please refer to:

<https://www.ema.europa.eu/en/events/product-management-service-pms-info-day>

Member states contact points for review of national versions of the content of mobile scanning and other technologies

Published on: 23 - April - 2024

For more information, please refer to:

https://www.ema.europa.eu/en/documents/other/member-states-contact-points-review-national-versions-content-mobile-scanning-other-technologies_en.pdf

Contact details of national competent authorities for requests to use a sticker to place the Unique Identifier on the outer/immediate packaging of centrally approved products

Published on: 23 - April - 2024

For more information, please refer to:

https://www.ema.europa.eu/en/documents/other/contact-details-national-competent-authorities-requests-use-sticker-place-unique-identifier-outer-immediate-packaging-centrally-approved-products_en.pdf

Humanarzneimittel - EMA

List of centrally authorised products requiring a notification of a change for update of annexes (updated)

Published on: 25 - April - 2024

For more information, please refer to:

https://www.ema.europa.eu/en/documents/regulatory-procedural-guideline/list-centrally-authorized-products-requiring-notification-change-update-annexes_en.pdf

Orphan Drugs und neuartige Therapierichtungen (ATMP)

Committee for Advanced Therapies (CAT) meeting with interested parties, Event - 16 - May - 2023

Minutes available

Published on: 12 - April - 2024

For more information, please refer to:

<https://www.ema.europa.eu/en/events/committee-advanced-therapies-cat-meeting-interested-parties-may-2023>

Qualität – Quality

No news published

(Prä-) Klinische Forschung – Research and Development

ACT EU Clinical Trials Analytics Workshop, Event – 25 to 26 - January - 2024 (updated)

Presentations and Video recording available

Published on: 16 - April - 2024

For more information, please refer to:

<https://www.ema.europa.eu/en/events/act-eu-clinical-trials-analytics-workshop-january-2024>

Clinical Trials Information System Webinar: Last Year of Transition, Event - 25 - March - 2024 (updated)

Presentations and Video recording available

Published on: 16 - April - 2024

For more information, please refer to:

<https://www.ema.europa.eu/en/events/clinical-trials-information-system-webinar-last-year-transition>

Clinical Trials Information System: training and support (updated)

Published on: 16 - April - 2024

For more information, please refer to:

<https://www.ema.europa.eu/en/human-regulatory-overview/research-development/clinical-trials-human-medicines/clinical-trials-information-system-training-support>

Humanarzneimittel - EMA

Clinical Trials Information System (CTIS) Bitesize Talk: How to submit a transitional trial in CTIS, Event - 29 - February - 2024 (updated)

Presentation and Video recording available

Published on: 22 - April - 2024

For more information, please refer to:

<https://www.ema.europa.eu/en/events/clinical-trials-information-system-ctis-bitesize-talk-how-submit-transitional-trial-ctis-0>

Non-clinical Working Party

Published on: 25 - April - 2024

For more information, please refer to:

<https://www.ema.europa.eu/en/committees/working-parties-other-groups/chmp/non-clinical-working-party>

Kinderarzneimittel – Paediatrics

No news published

Pflanzliche Arzneimittel – Herbal medicines

Template for a European Union herbal monograph (updated)

Published on: 15 - April - 2024

For more information, please refer to:

<https://www.ema.europa.eu/en/template-european-union-herbal-monograph>

Herbal medicinal product: Eucalypti aetheroleum, F: Assessment finalised (updated)

Published on: 15 - April - 2024

For more information, please refer to:

<https://www.ema.europa.eu/en/medicines/herbal/eucalypti-aetheroleum>

Herbal medicinal product: Pilocellae herba cum radice, F: Assessment finalised (updated)

Published on: 15 - April - 2024

For more information, please refer to:

<https://www.ema.europa.eu/en/medicines/herbal/pilosellae-herba-cum-radice>

Draft guideline on good agricultural and collection practice (GACP) for starting materials of herbal origin - Revision 1

Published on: 18 - April - 2024

For more information, please refer to:

https://www.ema.europa.eu/en/documents/scientific-guideline/draft-guideline-good-agricultural-collection-practice-gacp-starting-materials-herbal-origi-revision-1_en.pdf

Humanarzneimittel - EMA

Draft template for assessment report for the development of European herbal monographs and European Union list entries - Revision 6

Published on: 18 - April - 2024

For more information, please refer to:

https://www.ema.europa.eu/en/documents/template-form/draft-template-assessment-report-development-european-herbal-monographs-european-union-list-entries-revision-6_en.pdf

Technologies for indoor pathogen management in pandemics: insights from the JRC-HERA study

Published on: 17 - April - 2024

For more information, please refer to:

https://health.ec.europa.eu/latest-updates/technologies-indoor-pathogen-management-pandemics-insights-jrc-hera-study-2024-04-17-0_en

Targeted Consultation on EU4Health: have your say on future priorities, orientations and needs

Published on: 22 - April - 2024

For more information, please refer to:

<https://ec.europa.eu/newsroom/sante/newsletter-archives/52540>

Commission welcomes European Parliament's adoption of the regulation on European Health Data Space and the regulation on substances of human origin

Published on: 24 - April - 2024

For more information, please refer to:

https://ec.europa.eu/commission/presscorner/detail/en/IP_24_2250

Critical medicine shortages - immediate workflow and trial phase with two ongoing shortages

Published on: 19 - April - 2024

For more information, please refer to:

<https://www.edqm.eu/en/-/critical-medicine-shortages-immediate-workflow-and-trial-phase-with-two-ongoing-shortages>

Medizinprodukte

MDCG 2024-4 - Safety reporting in performance studies of in vitro diagnostic medical devices under Regulation (EU) 2017/746

Published on: 15 - April - 2024

For more information, please refer to:

https://health.ec.europa.eu/latest-updates/mdcq-2024-4-safety-reporting-performance-studies-vitro-diagnostic-medical-devices-under-regulation-2024-04-15_en

MDCG 2022-9 rev.1 - Summary of safety and performance template (updated)

Published on: 15 - April - 2024

For more information, please refer to:

https://health.ec.europa.eu/latest-updates/update-mdcq-2022-9-rev1-summary-safety-and-performance-template-april-2024-2024-04-15_en

MDCG 2024-5 - Guidance on the Investigator's Brochure content

Published on: 17 - April - 2024

For more information, please refer to:

https://health.ec.europa.eu/latest-updates/mdcq-2024-5-guidance-investigators-brochure-content-april-2024-2024-04-17_en

CMDH AGENDAS AND MINUTES (updated)

Published on: 22 - April - 2024

For more information, please refer to:

<https://www.hma.eu/human-medicines/cmdh/agendas-and-minutes.html>

Humanarzneimittel - Deutschland

Liste der Veröffentlichungen von Zusammenfassungen von Risikomanagementplänen nach § 34 Abs. 1a AMG

Veröffentlicht am: 12 - April - 2024

Weitere Informationen finden Sie unter:

<https://www.bfarm.de/SharedDocs/Downloads/DE/Arzneimittel/Pharmakovigilanz/Risikoinformationen/RMP/liste-rmp-summary.html?nn=986770>

Organigramm des BfArM

Veröffentlicht am: 12 - April - 2024

Weitere Informationen finden Sie unter:

https://www.bfarm.de/SharedDocs/Downloads/DE/BfArM/Org/bfarm_organigramm.html?nn=986770

Aktuell laufende und bestätigte Arzneimittel-Härtedefallprogramme

Veröffentlicht am: 15 - April - 2024

Weitere Informationen finden Sie unter:

<https://www.bfarm.de/DE/Arzneimittel/Klinische-Pruefung/Compassionate-Use/compUse-tabelle.html?nn=986770>

Liste der versorgungsrelevanten und versorgungskritischen Wirkstoffe nach § 52b Absatz 3c AMG

Veröffentlicht am: 16 - April - 2024

Weitere Informationen finden Sie unter:

https://www.bfarm.de/SharedDocs/Downloads/DE/Arzneimittel/Zulassung/amInformationen/Lieferengpaesse/liste_wirkstoffe_gesamt_excel.html?nn=986770

Informationen und Schulungsunterlagen

Veröffentlicht am: 16 - April - 2024

Weitere Informationen finden Sie unter:

<https://www.bfarm.de/DE/Arzneimittel/Zulassung/Zulassungsrelevante-Themen/e-Submission/Pharmnet-Bund.html?nn=986770>

Informationen zur Verfügbarkeit von salbutamolhaltigen Arzneimitteln in pulmonaler Darreichungsform und Empfehlung zur Abmilderung möglicher Engpässe

Veröffentlicht am: 17 - April - 2024

Weitere Informationen finden Sie unter:

<https://www.bfarm.de/DE/Arzneimittel/Arzneimittelinformationen/Lieferengpaesse/salbutamol.html?nn=986770>

Meldeverpflichtungen

Veröffentlicht am: 17 - April - 2024

Weitere Informationen finden Sie unter:

<https://www.bfarm.de/DE/Arzneimittel/Arzneimittelinformationen/Lieferengpaesse/Meldeverpflichtungen/artikel.html?nn=986770>

Humanarzneimittel - Deutschland

Wissenswertes zu DiGA

Veröffentlicht am: 17 - April - 2024

Weitere Informationen finden Sie unter:

<https://www.bfarm.de/DE/Medizinprodukte/Aufgaben/DiGA-und-DiPA/DiGA/Wissenswertes/artikel.html?nn=986770>

Prüfkriterien für die von digitalen Gesundheitsanwendungen (DiGA) und digitalen Pflegeanwendungen (DiPA) nachzuweisenden Anforderungen an den Datenschutz

Veröffentlicht am: 24 - April - 2024

Weitere Informationen finden Sie unter:

<https://www.bfarm.de/SharedDocs/Downloads/DE/Medizinprodukte/diga-dipa-datenschutzkriterien.html?nn=986770>

Europäisches Arzneibuch - Kapitel "Phage therapy medicinal products (5.31)"

Veröffentlicht am: 17 - April - 2024

Weitere Informationen finden Sie unter:

<https://www.bfarm.de/DE/Arzneimittel/Zulassung/Zulassungsrelevante-Themen/Arzneibuch/euopaisches-arzneibuch-bakteriophagen.html?nn=986770>

Rohdaten der Stoffbezeichnungen

Veröffentlicht am: 18 - April - 2024

Weitere Informationen finden Sie unter:

<https://www.bfarm.de/DE/Arzneimittel/Arzneimittelinformationen/Arzneimittel-recherchieren/Stoffbezeichnungen/artikel.html?nn=986770>

Statistiken

Veröffentlicht am: 18 - April - 2024

Weitere Informationen finden Sie unter:

<https://www.bfarm.de/DE/Aktuelles/Statistiken/artikel.html?nn=986770>

Homöopathische und anthroposophische Arzneimittel

Veröffentlicht am: 18 - April - 2024

Weitere Informationen finden Sie unter:

<https://www.bfarm.de/DE/Arzneimittel/Zulassung/Zulassungsarten/Besondere-Therapierichtungen-und-traditionelle-Arzneimittel/Homoeopathische-und-anthroposophische-Arzneimittel/amanthropo-inhalt.html?nn=986770>

Arzneimittel-Lieferengpass-bekämpfungs- und Versorgungs-verbesserungs-gesetz (ALBVVG)

Veröffentlicht am: 19 - April - 2024

Weitere Informationen finden Sie unter:

<https://www.bfarm.de/DE/Arzneimittel/Arzneimittelinformationen/Lieferengpaesse/ALBVVG/artikel.html?nn=986770>

PSUR Single Assessment (PSUSA)

Veröffentlicht am: 19 - April - 2024

Weitere Informationen finden Sie unter:

<https://www.bfarm.de/DE/Arzneimittel/Pharmakovigilanz/Periodic-Safety-Update-Reports/PSURs/PSUR-Single-Assessment/artikel.html?nn=986770>

Humanarzneimittel - Deutschland

Terminologieserver

Veröffentlicht am: 24 - April - 2024

Weitere Informationen finden Sie unter:

https://www.bfarm.de/DE/Kodiersysteme/Services/Terminologieserver/_artikel.html?nn=986770

Sunset Clause

Veröffentlicht am: 24 - April - 2024

Weitere Informationen finden Sie unter:

https://www.bfarm.de/DE/Arzneimittel/Zulassung/Folgeverfahren/Sunset-Clause/_artikel.html?nn=986770

Vorkommnismeldung durch Hersteller und Bevollmächtigte

Veröffentlicht am: 25 - April - 2024

Weitere Informationen finden Sie unter:

https://www.bfarm.de/DE/Medizinprodukte/Antraege-und-Meldungen/Vorkommnis-melden/Hersteller-und-Bevollmaechtigte/_artikel.html?nn=986770

Humanarzneimittel - Österreich

Elektronische Einreichung

Veröffentlicht am: 16 - April - 2024

Weitere Informationen finden Sie unter:

<https://www.basq.gv.at/fuer-unternehmen/zulassung-life-cycle/elektronische-einreichung>

Geschäftsordnung des BASG

Veröffentlicht am: 22 - April - 2024

Weitere Informationen finden Sie unter:

<https://www.basq.gv.at/ueber-uns/basq-veroeffentlichungen/geschaeftsordnung>

FAQ Pharmakovigilanz

Veröffentlicht am: 25 - April - 2024

Weitere Informationen finden Sie unter:

<https://www.basq.gv.at/fuer-unternehmen/pharmakovigilanz/faq-pharmakovigilanz>

Zur Stellungnahme (Monographieentwürfe)

Veröffentlicht am: 26 - April - 2024

Weitere Informationen finden Sie unter:

<https://www.basq.gv.at/gesundheitsberufe/oesterreichisches-arzneibuch/zur-stellungnahme>

Humanarzneimittel - Schweiz

Paclitaxel-beschichtete Ballone und Paclitaxel-eluierende Stents

Veröffentlicht am: 15 - April - 2024

Weitere Informationen finden Sie unter:

<https://www.swissmedic.ch/swissmedic/de/home/medizinprodukte/uebersicht-medicinprodukte/infos-zu-bestimmten-medicinprodukten/update-paclitaxel-beschichteten-ballonen-und-paclitaxel-eluierenden-stents.html>

Teilnahme von Swissmedic am Swiss Biotech Day in Basel

Veröffentlicht am: 24 - April - 2024

Weitere Informationen finden Sie unter:

<https://www.swissmedic.ch/swissmedic/de/home/news/mitteilungen/teilnahme-smc-am-swiss-biotech-day-basel.html>

Aktualisierte Vorgabedokumente

Veröffentlicht am: 24 - April - 2024

Weitere Informationen finden Sie unter:

https://www.swissmedic.ch/swissmedic/de/home/news/updates/updated_documents/apr-2024.html



Fragen an das Netzwerk

Falls Sie eine Frage haben, die Sie gerne in unserem Netzwerk diskutieren würden, senden Sie uns einfach eine E-Mail an info-as@megra.org zur anonymen Publikation im nächsten Newsletter.*

*Bei der Beantwortung der Fragen handelt es sich um eine Zusammenfassung von persönlichen Meinungen und Erfahrungswerten der MEGRA Mitglieder mit keinem Anspruch auf Rechtssicherheit. Wir empfehlen zur Absicherung die Konsultation entsprechender zugrunde liegender Regularien.

Veranstaltungen / Events – Behörden und andere Veranstalter

Deutschland

Klinische Prüfung: BfArM trifft...!

Ort: online

Termin: 10 - Mai - 2024

Weitere Informationen finden Sie unter:

<https://www.bfarm.de/DE/Aktuelles/Veranstaltungen/Termine/2024-05-10-klinische-pruefung-austausch.html?nn=986770>

BfArM im Dialog „4. Anwenderforum SNOMED CT“

Ort: online

Termin: 14 - Mai - 2024

Weitere Informationen finden Sie unter:

<https://www.bfarm.de/DE/Aktuelles/Veranstaltungen/Termine/2024-05-14-snomed-anwenderforum.html>

8. Nationale Impfkonzferenz

Ort: World Conference Center, ehemaliger Plenarsaal Bundestag, Bonn

Termin: 13 bis 14 - Juni - 2024

Weitere Informationen finden Sie unter:

<https://www.pei.de/SharedDocs/veranstaltungen-events/DE/2024/2024-06-13-nationale-impfkonzferenz.html?nn=170994>

Österreich

Keine Veranstaltungen veröffentlicht

Schweiz

Keine Veranstaltungen veröffentlicht

Europa

Information and Q&A session on updated CAPs in web-based eAF

Where: online and European Medicines Agency, Amsterdam, the Netherlands

Date: 07 - May - 2024

For more information, please refer to:

<https://www.ema.europa.eu/en/events/information-qa-session-updated-caps-web-based-eaf>

Veranstaltungen / Events – Behörden und andere Veranstalter

Clinical Trials Information System (CTIS): Walk-in clinic

Where: online and European Medicines Agency, Amsterdam, the Netherlands

Date: 15 - May - 2024

For more information, please refer to:

<https://www.ema.europa.eu/en/events/clinical-trials-information-system-ctis-walk-clinic-may-2024>

12th European Conference on Rare Diseases & Orphan Products (ECRD)

Where: online and at The Square in Brussels

Date: 15 to 16 - May - 2024

For more information, please refer to:

<https://www.rare-diseases.eu/>

Product Management Service (PMS) Product UI and API training (access & navigation)

Where: online and European Medicines Agency, Amsterdam, the Netherlands

Date: 03 - June - 2024

For more information, please refer to:

<https://www.ema.europa.eu/en/events/product-management-service-pms-product-ui-api-training-access-navigation>

Mandatory use of ISO/ICH E2B(R3) individual case safety reporting in the EU: hands-on training course using the EudraVigilance system

Where: online

Date: 03 to 07 - June - 2024

For more information, please refer to:

<https://www.ema.europa.eu/en/events/mandatory-use-iso-ich-e2br3-individual-case-safety-reporting-eu-hands-training-course-using-eudravigilance-system-40>

Save the date - “TODAY, TOMORROW, TOGETHER FOR PUBLIC HEALTH”

Where: Palais de l'Europe, Strasbourg (France)

Date: 11 to 12 - June - 2024

For more information, please refer to:

<https://www.edqm.eu/en/celebrating-60-years-of-excellence-in-public-health>

Clinical Trials Information System (CTIS) sponsor end user training programme

Where: European Medicines Agency, Amsterdam, the Netherlands

Date: 10 to 13 - June - 2024

For more information, please refer to:

<https://www.ema.europa.eu/en/events/clinical-trials-information-system-ctis-sponsor-end-user-training-programme-june-2024>

Veranstaltungen / Events – Behörden und andere Veranstalter

Extended EudraVigilance medicinal product dictionary (XEVMPPD) training course (for sponsors)

Where: online

Date: 12 to 13 - June - 2024 and 18 to 20 - June - 2024

For more information, please refer to:

<https://www.ema.europa.eu/en/events/extended-eudravigilance-medicinal-product-dictionary-xevmpd-training-course-sponsors-0>

<https://www.ema.europa.eu/en/events/extended-eudravigilance-medicinal-product-dictionary-xevmpd-training-course-1>

Industry Update webinar on Regulatory Procedure Management for Product Lifecycle Management on IRIS

Where: online and European Medicines Agency, Amsterdam, the Netherlands

Date: 13 - June - 2024

For more information, please refer to:

<https://www.ema.europa.eu/en/events/industry-update-webinar-regulatory-procedure-management-product-lifecycle-management-iri>

Joint HMA/EMA Big Data Steering Group workshop on RWE methods

Where: online and European Medicines Agency, Amsterdam, the Netherlands

Date: 14 - June - 2024

For more information, please refer to:

<https://www.ema.europa.eu/en/events/joint-hma-ema-big-data-steering-group-workshop-rwe-methods>

Network Update webinar on Regulatory Procedure Management for Product Lifecycle Management on IRIS

Where: online and European Medicines Agency, Amsterdam, the Netherlands

Date: 18 - June - 2024

For more information, please refer to:

<https://www.ema.europa.eu/en/events/network-update-webinar-regulatory-procedure-management-product-lifecycle-management-iris>

38th ISBT International Congress

Where: Barcelona, Spain

Date: 23 to 27 - June - 2024

For more information, please refer to:

<https://www.edqm.eu/en/38th-isbt-international-congress>

EMA workshop on the challenges in drug development, regulation and clinical practice in hemoglobinopathies

Where: online and European Medicines Agency, Amsterdam, the Netherlands

Date: 01 - July - 2024

For more information, please refer to:

<https://www.ema.europa.eu/en/events/ema-workshop-challenges-drug-development-regulation-clinical-practice-hemoglobinopathies>

Veranstaltungen / Events – Behörden und andere Veranstalter

SPOR Status Update

Where: online and European Medicines Agency, Amsterdam, the Netherlands

Date: 10 - July - 2024

For more information, please refer to:

<https://www.ema.europa.eu/en/events/spor-status-update-0>