



EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

Support and assistance offered by EMA to veterinary academic stakeholders

EMA Veterinary Awareness Day

Presented by Valentin Nicorescu on 13 September 2023
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An agency of the European Union





Agency tools available for veterinary academic stakeholders

Support and assistance

1. Innovation Task Force (ITF) briefing meetings
2. Scientific guidance for veterinary medicines
3. Scientific Advice procedures
4. Support to small and medium-sized enterprises (SMEs)

Information and communication

1. Newsletters
2. EMA website
3. Social media



Support and assistance



Fostering research and innovation

A strategic goal

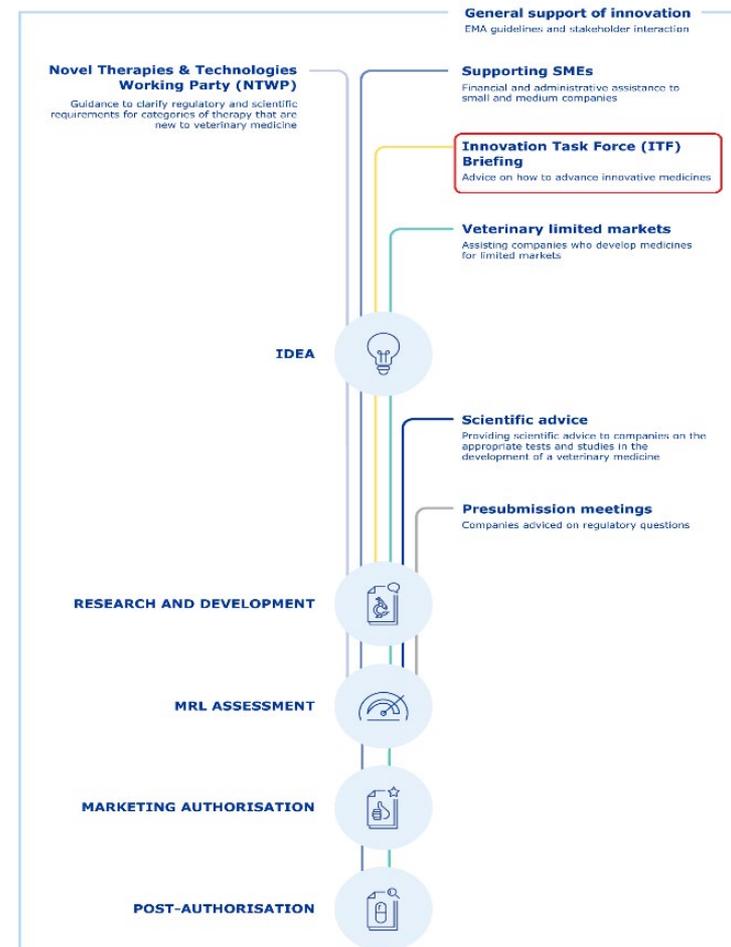
- One of EMA's strategic goals is ***to foster research and the uptake of innovative methods in the development of human and veterinary medicines.***
- The Agency supports the development of innovative methodologies by fostering collaboration across the regulatory network and with academia.





ITF briefing meetings

- Forum for *early* dialogue on medicines innovation, **very early** in the product development.
- Cover issues arising from innovative medicines development, new technologies and borderline products.
- Informal exchange of information and guidance in the development process.
- Attended by experts from the Agency network, scientific committees (CVMP) and working parties (e.g. NTWP).
- Free of charge.





ITF briefings

How to request a meeting

Request

- A [request form](#) to be completed and sent via ITFvet@ema.europa.eu

Review

- EMA reviews the form and advises applicants on how to proceed

Submission

- Applicants prepare submission via a dedicated platform

Early interaction

- A meeting takes place





2. Scientific guidance for veterinary medicines

- Designed to help applicants **prepare their marketing authorisation applications** for veterinary medicines (VMPs).
- Applicants and marketing authorisation holders are strongly encouraged to **follow scientific guidelines** while developing VMPs.
- Scientific guidelines for veterinary medicines available for different sections: quality, safety and residues, efficacy, antimicrobials, immunologicals, novel therapies.
- Stakeholders are encouraged to comment on the draft guidelines during the **public consultation**, as it ensures that their opinion and scientific input are captured and good quality documents can be made available.



Scientific guidance for veterinary medicines

Public consultation

- Concept papers, reflection papers, scientific guidelines
- Draft guidance documents are published on EMA website
- Stakeholders are invited to review these documents and provide their general and specific comments in a specified timeframe – **public consultation**
- Stakeholders' comments are considered when the final version of the document is prepared
- An 'Overview of comments' document is also prepared and published, where feedback is given to all comments received

More information on scientific guidelines: [here](#)

Data requirements for veterinary medicinal products intended to reduce the risk of transmission of vector-borne pathogens in dogs and cats - Scientific guideline [← share](#)

Table of contents

- Current version
- Document history - First version

This [guideline](#) provides recommendations for the design and conduct of studies to support the efficacy of veterinary medicinal products (VMPs) intended for the reduction of the risk of transmission of vector-borne pathogens (VBPs) in dogs and cats, which can be transferred by blood-feeding arthropods. The [guideline](#) outlines the requirements for pre-clinical studies and clinical trials.

Keywords: Vector-borne disease (VBD), vector-borne pathogen (VBP), blood-feeding arthropods, reduction of the risk of transmission of VBPs, efficacy, veterinary medicinal product (VMP), ectoparasitocides, repellent effect, dog, cat

Current version

[Guideline on data requirements for veterinary medicinal products intended to reduce the risk of transmission of vector-borne pathogens in dogs and cats \(PDF/422.33 KB\)](#)

Adopted

First published: 24/06/2022
Legal effective date: 01/01/2023
EMA/CVMP/EWP/278031/2015

[Overview of comments received on the 'Guideline on data requirements for veterinary medicinal products for the prevention of transmission of vector-borne diseases in dogs and cats' \(EMA/CVMP/EWP/278031/2015\) \(PDF/364.93 KB\)](#)

First published: 24/06/2022
EMA/CVMP/EWP/552708/2018

Document history - First version

[Draft guideline on data requirements for veterinary medicinal products for the prevention of transmission of vector-borne diseases in dogs and cats \(PDF/181.1 KB\)](#)

Draft: consultation closed

First published: 27/07/2018
Last updated: 27/07/2018
Consultation dates: 27/07/2018 to 31/08/2019
EMA/CVMP/EWP/278031/2015

[Concept paper for a guideline on data requirements regarding veterinary medicinal products for the prevention of transmission of canine and feline vector-borne diseases \(PDF/81.29 KB\)](#)

Draft: consultation closed

First published: 18/11/2014
Last updated: 18/11/2014
Consultation dates: 18/11/2014 to 28/02/2015
EMA/CVMP/EWP/309734/2014



3. Scientific Advice procedures

- EMA can provide scientific advice on the appropriate tests and studies to be conducted in the development of a VMP.
- Available to applicants including commercial enterprises, but **also research organisations and academics**.
- Scientific advice can be requested at any stage.
- For veterinary medicines, scientific advice is given by the CVMP on the recommendation of the Scientific Advice Working Party (SAWP-V).



Scientific Advice procedures

How it works

- The applicant poses specific questions on the development of a VMP.
 - *Questions should be clear as possible and address quality, safety and efficacy issues.*
- The applicant presents the VMP development plan and identifies possible solutions.
- CVMP gives advice on the developer's proposals.



Scientific advice from EMA is not legally binding on EMA or on the medicine developer.



NOT Legally binding

- [Guidance for applicants](#) requesting scientific advice
- Request for scientific advice [form](#)
- More information: [here](#)

4. Small and medium-sized enterprises (SMEs)

- SMEs are considered an important source of innovation and development of new medicines.
- EMA provides incentives and support for SMEs developing medicines for human or veterinary use.



- The **EMA's SME office** provides advice, guidance and assistance to SMEs to help to find their way through the centralised authorisation procedure.
- SME briefing meetings possible; questions on regulations, administrative requirements or procedures can be addressed.



Agency's support to SMEs

- Enterprises should have **SME status assigned** by EMA.
- Once the SME status has been granted, the company is included in **EMA's public SME register**.
- The support for SMEs includes:
 - **fee exemptions** and **reductions** for regulatory procedures.
 - **translation assistance**.

More information can be found in the EMA's [SME user guide](#).
Guidance on [applying for SME status](#) and [financial advantages](#) for SMEs is also available.



Information and communication



1. Newsletters

Veterinary Medicines Highlights:

- published quarterly;
- focuses on news, activities and interviews from the Veterinary Medicines Division of the EMA.

SME Office Newsletter:

- news and guidance of relevance to SMEs and their stakeholders;
- developments in EU regulations and regulatory environment;
- events and training opportunities.



Welcome to the 13th edition of the Veterinary Medicines highlights and to its new format! As previously announced, we are expanding the content to cover news and updates on all activities of the Veterinary Medicines Division.



“ We asked your opinion, and your feedback was fundamental in re-shaping the newsletter to a format that better suits your interests. ”

2. EMA's corporate website - www.ema.europa.eu



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Academia

On this page, you will find information on the European Medicines Agency's (EMA) activities that are most relevant to academia, including news and events. Learn more about the Agency's resources to support medicine development:

[Human regulatory: Research and development](#)
[Veterinary regulatory: Research and development](#)

Learn more about [how EMA interacts with academia](#).

Featured information

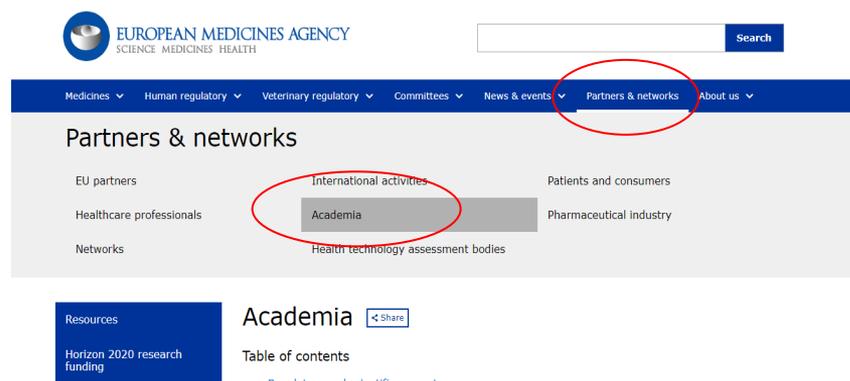
Live interview with EMA's Head of veterinary medicines: 7 September, 13:15 to 13:45 CEST

EMA is inviting interested stakeholders to follow and ask questions during our live interview with the Agency's Head of veterinary medicines, Ivo Claassen. The interview is broadcasted on EMA's corporate LinkedIn account. The conversation focuses on the interlinks between human and animal health, and the need for innovation in veterinary medicines.

Join the EMA Veterinary Awareness Day

EMA invites stakeholders in the veterinary field to the first edition of its Veterinary Awareness Day, held on 12 and 13 September. This online event will provide insight in the Agency's work on topics such as animal health and welfare, antimicrobial resistance, support for innovation and environmental risk assessment. Register by 11 September.

Academia news page



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Partners & networks

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- Networks
- International activities
- Academia
- Health technology assessment bodies
- Patients and consumers
- Pharmaceutical industry

Resources

Horizon 2020 research funding

Academia

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- Regulatory and scientific support
- Cancer Medicines Forum
- Enabling oncology scientists' participation in medicine regulation (pilot project) (new)
- Exchange of expertise
- Support for consortia in externally funded projects
- Framework for collaboration

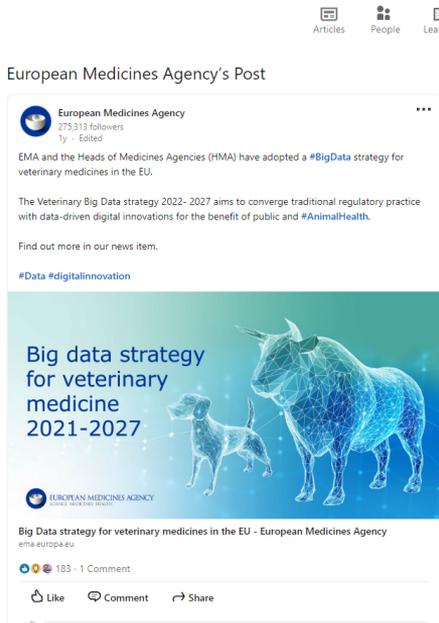
The European Medicines Agency (EMA) is committed to maintaining a strong working relationship with European academics and researchers. Collaboration between the Agency and academia is necessary for the Agency to be prepared for future challenges and opportunities offered by advances in science and technology. The Agency has targeted engagement with academia, learned societies and research groups in a range of areas.

Academia dedicated landing page

3. LinkedIn, X, YouTube

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EMA and the Heads of Medicines Agencies (HMA) have adopted a #BigData strategy for veterinary medicines in the EU.

The Veterinary Big Data strategy 2022- 2027 aims to converge traditional regulatory practice with data-driven digital innovations for the benefit of public and #AnimalHealth.

Find out more in our news item.
#Data #digitalinnovation



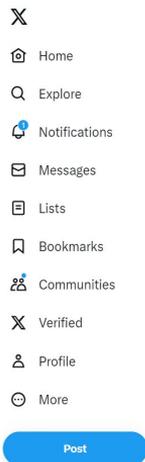
Big data strategy for veterinary medicine 2021-2027

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Big Data strategy for veterinary medicines in the EU - European Medicines Agency
ema.europa.eu

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Mark your calendar for the 1st-ever EMA Veterinary Awareness Day!

Join us on 12 & 13 Sept to learn more about the Agency's work in #veterinary medicines and the following topics

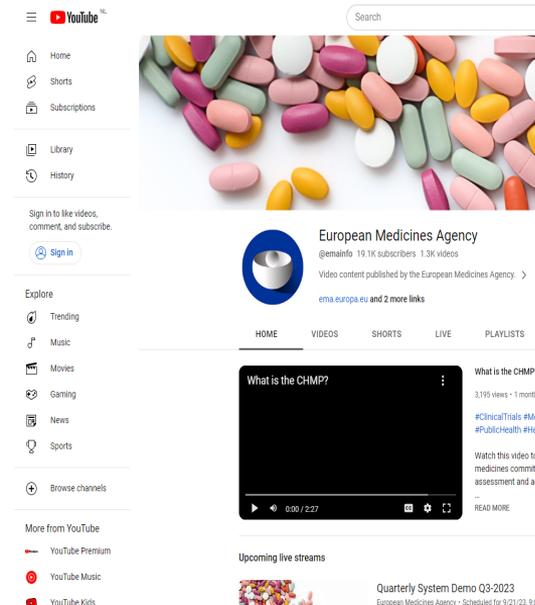
- animal welfare
- animals in society
- antimicrobial resistance
- environmental risk assessment



Register now!
Veterinary Awareness Day
12 & 13 September 2023

EMA
europa.eu
EMA Veterinary Awareness Day

4:38 pm · 2 Aug 2023 · 9,250 Views



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Upcoming live streams

Quarterly System Demo Q3-2023
European Medicines Agency · Scheduled for 9/12/23, 9:00 AM



Any questions?

Further information

Send a question to the European Medicines Agency

Official address Domenico Scarlattilaan 6 • 1083 HS Amsterdam • The Netherlands

Telephone +31 (0)88 781 6000

Send us a question Go to www.ema.europa.eu/contact

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