

Annual Seminar on the Harmonisation of the Registration and Control of Veterinary Medicines Committee of the Americas on Veterinary Medicines (CAMEVET)

14 to 16 October 2026
Buenos Aires, Argentina

AGENDA

Day 1: 14 October 2026			
8:00-12:00	<u>MANDATORY - Participant registration -</u>		
Time	Topic	Duration	Speaker
12:30-12:45	Opening Session – Opening Speeches at the Seminar - WOAHA Authorities - SENASA Argentina Authorities - CAMEVET Authorities	15 min	To be confirmed
12:45-12:50	Assumption of the Presidency of CAMEVET	5 min	To be confirmed
Module I: History and Future – 30 years of CAMEVET			
12:50-13:10	- What is CAMEVET and how does it work? - The evolution of the harmonisation process in the Americas. - Key milestones achieved. - The impact of harmonised guidelines. - Public-private partnership as a working model. - Regulatory challenges for the coming decade.	30 min	Ana M. Sgammini CAMEVET Administrative Secretary
13:10-14:40	Lunch		
Module II: Working documents			
14:40-14:50	Introduction to CAMEVET working group procedures	10 min	Ana M. Sgammini CAMEVET Administrative Secretary
Progress on the working documents			
Documents currently being processed IV – Submission of the final draft			
14:50-15:05	Supplementary studies to the stability guide	15 min	Andrea Fraga CAPROVE Argentina
15:05-15:20	Update to free sale certificate templates and exclusive export authorisation	15 min	Luiz Monteiro SINDAN Brazil
15:20-15:35	Annexes – Good Manufacturing Practice for Veterinary Products	15 min	Natalia Cardozo Uruguay
15:35-15:45	Guidance on the Classification and Inspection of Veterinary Products without a Therapeutic Indication	10 min	To be confirmed
Documents currently being processed II – Submission of the first draft			
15:45-16:00	Guide to registering fixed combinations	15 min	Marién Gutiérrez ALFA El Salvador
16:00-16:15	Register of cannabis-based medicines	15 min	Angélica Barbosa APROVET Colombia
16:15-16:45	Coffee Break		
16:45-17:00	Good practice in the use of veterinary products	15 min	Aida Rojas Colombia
17:00-17:15	Efficacy trials for the registration of internal and external antiparasitic products for small animals	15 min	Ximena Melón CAPROVE Argentina
17:15-17:30	Guidance on establishing criteria for the use of computational models as alternative methods for determining the efficacy of veterinary antimicrobials)	15 min	Leonardo Viana Brazil



17:30-17:45	Guidance on establishing the criteria for requesting residue depletion testing in the registration and renewal of veterinary medicinal product authorisations containing known active substances in non-innovative formulations	15 min	Carlos Rufrano CLAMEVET Argentina
17:45-18:00	Guide to Critical Veterinary Products	15 min	To be confirmed
Documents currently being processed I – Submission of a memorandum			
18:00-18:15	Inspection guide for the manufacture of pharmaceutical, biological and ectoparasiticide products	15 min	To be confirmed
18:15	News guides		

Day 2: 15 October 2026			
Time	Topic	Duration	Speaker
Module III: Quality of veterinary products			
Turning harmonised guidelines into practical tools			
<i>Panel comprising health authorities, industry representatives and technical experts; real-life implementation examples.</i>			
9:00-9:45	Pharmacovigilance: A Global Perspective - An overview of the main global pharmacovigilance regulations (EMA, VMD and FDA). - Impact of differing requirements amongst CAMEVET member countries. - Review of the CAMEVET guide 'Guide to the Implementation of a Pharmacovigilance System for Veterinary Medicinal Products' Experience in implementing the CAMEVET Guidelines in Mexico	45 min	Martin Tiemann Responsable Global de Políticas e Inteligencia de Farmacovigilância, Boehringer Ingelheim Maria Elena González Ruiz Director of Establishment Regulation; OMSA Focal Point for Veterinary Products
Substandard and counterfeit veterinary products:			
Regional experiences, challenges and cooperation in tackling the issue			
<i>Round Table</i>			
9:45-10:30	SECTION 1. Current situation and scale of the problem SECTION 2. Prevention and detection SECTION 3. Incident response and coordination SECTION 4. National and regional challenges and opportunities Q&A session	45 min	Moderator: Andrés García Campos WOAH Department of Antimicrobials
10:30 - 11:00 Coffee Break			
Strengthening regional tools for the monitoring of unsafe veterinary products			
11:00-11:45	- The evolution from VSAFE to TRUVET. - Objectives and scope of the platform. - Benefits for authorities and industry.	45 min	Andrés García Campos WOAH Department of Antimicrobials
Module IV: Use of veterinary products			
Antimicrobials and the One Health approach Read more			
11:45-12:25	From setting priorities to regulatory intervention: vaccines as a tool for reducing the use of antibiotics	40 min	Gita Mailk-Dahiya WOAH Veterinary Products Focal Point and member of the ad hoc group
The use of human medicines in veterinary medicine: experience, challenges and regulatory approaches in the region			
<i>Panel of regulatory authorities and industry representatives</i>			
12:25-13:10	- Situations leading to the use of human medicines in veterinary practice - Experience in different countries - Health risks - Impact on food-producing animals - The veterinary industry's perspective - Possible regulatory strategies and the need for harmonisation	45 min	Coordination: Executive Board
13:30-15:00 Lunch			
Section V: Training			
Training for regulatory authorities on the use of TRUVET			
<i>During this event, participants from the private sector will be able to use the coffee area for informal meetings</i>			
15:00-16:30	- Initial connection. - Submission of the baseline report. - Practical exercise in incident reporting. - Methods of clarification. - Viewing and using the database.	90 min	Andrés García Campos WOAH Department of Antimicrobials



16:30-17:00	Coffee Break		
Training for the private sector on the use of TRUVET			
17:00-18:00	- Initial contact - Reporting incidents. - Liaising with WOA - Sharing information with the relevant authorities.	60 min	Andrés García Campos WOAH Department of Antimicrobials

Day 3: October 16, 2026			
Time	Topic	Duration	Speaker
Review of priority issues from the public sector and industry			
09:00-09:20	Issue tracking: Public Sector Progress, challenges and proposals	20 min	Members of the Executive Committee
09:20-09:40	Issue tracking: Private sector Progress, challenges and proposals	20 min	Members of the Executive Committee
09:40-10:10	Conclusions of the Plenary Meeting of the Public and Private Sectors	20 min	Coordination: Executive Board
10:10-10:40	Round-table discussion between the public sector and industry 	30 min	Coordination: Executive Board
10:40 - 11:10	Coffee Break		
Vision for the Future			
11:10-11:45	Work plan for analysing the functioning of CAMEVET and its relationship with WOA, and discussion of opportunities for strengthening CAMEVET in the future.	45 min	Moderator: Francisco D'Alessio WOAH Regional Representative for the Americas
11:45-12:00	Approval of the new executive committee and review of the previous executive committee's performance	15 min	
12:00-12:15	CAMEVET's budget and resources	15 min	
12:15-12:25	Approval of the proposed venues for the forthcoming seminars	10 min	
Palabras de cierre seminario			