Organización Mundial de Sanidad Animal
Fundada como OIE

camevet
Comité de Las Américas de Medicamentos Veterinarios

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

**Buenos Aires, Argentina**

**14–16 October 2026**

# PRELIMINARY AGENDA

| **Day 1: 14 October 2026** |
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| **Topic** |
| Registration of participants |
| Opening Session – Opening Addresses at the Seminar |
| Assumption of the Presidency of CAMEVET |
| **Section I: History and Future – 30 years of CAMEVET** |
| - What is CAMEVET and how does it work?<br>- Progress of the harmonisation process in the Americas.<br>- Key milestones achieved.<br>- Impact of the harmonised guidelines.<br>- Public-private partnership as a working model.<br>- Regulatory challenges for the coming decade.<br>- The future of CAMEVET – relationship with WOAH |
| New regulatory challenges |
| **Section II: Working documents** |
| **Progress on the working documents** |
| Documents currently status IV – Submission of the final draft |
| Documents currently status III – Submission of the draft with comments |
| Documents currently status II – Submission of the first draft |
| Documents currently status I – Submission of a Concep Note |
| Launch of new guides |

| **Day 2: 15 October 2026** | |
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| **Section III: Quality of veterinary products** | |
| **Topic** | |
| **Turning harmonised guidelines into practical tools**<br>*Panel comprising health authorities, industry representatives and technical experts; real-life implementation case studies.* | |
| - **Good manufacturing practice**<br>- **Pharmacovigilance**<br>- **Stability**<br>- **Labelling** | - Status of implementation in different countries.<br>- Regulatory experiences.<br>- Industry challenges.<br>- Common inspection criteria.<br>- Benefits of harmonisation. |
| **Antimicrobials and the One Health approach** | |
| Vaccination strategies as a driver for reducing antimicrobial use (AMU) in animal production | |

| Section IV: Use of veterinary products |
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| **Substandard and counterfeit veterinary products:** |
| **Regional experiences, challenges and cooperation in combating the problem**<br>**Panel of regulatory authorities and industry representatives**<br>- Countries' experiences in detecting and managing substandard and counterfeit products<br>- Relevant cases and lessons learnt.<br>- Industry involvement in prevention and reporting.<br>- Information exchange and regional cooperation.<br>- Needs and opportunities for strengthening surveillance in the Americas.<br>**Strengthening regional tools for the monitoring of unsafe veterinary products**<br>- The evolution from VSAFE to TRUVET.<br>- Objectives and scope of the platform.<br>- Benefits for authorities and industry. |
| **Human medicines in veterinary medicine: experiences, challenges and regulatory approaches in the region** |
| **Panel of regulatory authorities and industry representatives**<br>- Situations leading to the use of human medicines in veterinary practice.<br>- Experiences from different countries.<br>- Health and regulatory risks.<br>- Impact on food-producing animals.<br>- The veterinary industry's perspective.<br>- Possible regulatory strategies and the need for harmonisation. |
| Section V: Training |

| Day 3: 16 October 2026 |
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| Section VI: Review of outstanding issues from the public sector and industry |
| Conclusions of the Plenary Meeting of the Public and Private Sectors |
| Round-table discussion between government representatives and industry representatives at the |
| Section VII: Future vision |
| CAMEVET's Future |
| Approval of the new executive committee |
| CAMEVET's budget and resources |
| Approval of the proposed venues for the seminars 2027 |
| ***Closing of the Seminar*** |