

A focus on the science of veterinary drugs

→ **Understanding Veterinary Active Pharmaceutical Ingredients (APIs): A Guide to Navigating Regulatory and Pharmacopeial Standards**—July 18–July 19, 2018



Co-sponsored by:
Generic Animal Drug Alliance (GADA)

Final Agenda

DAY ONE: Wednesday, July 18, 2018

- 8:30 – 9:00 a.m.** **Registration & Coffee**
- 9:00 – 9:15 a.m.** **Welcome and Opening Introduction**
Ronald T. Piervincenzi, Ph.D., *CEO, USP*
Courtney Tallman, *Program Planning Chair, GADA*
- 9:15 – 10:00 a.m.** **Using eSubmitter For Type II VMFs**
Scott Fontana, Ph.D., *Center for Veterinary Medicine, FDA*
Elizabeth Pollina Cormier, Ph.D., *Center for Veterinary Medicine, FDA*
Trupti Dhami, Ph.D., *Center for Veterinary Medicine, FDA*
- 10:00 – 10:45 a.m.** **Common Type II VMF Deficiencies**
Jason Dreabit, M.A., *Center for Veterinary Medicine, FDA*
Greg Hunter, Ph.D., *Center for Veterinary Medicine, FDA*
Renée Pietsch, Ph.D., *Center for Veterinary Medicine, FDA*
- 10:45 – 11:00 a.m.** **Morning Break**
- 11:00 – 11:15 a.m.** **CVM's VMF Website**
Greg Hunter, Ph.D., *Center for Veterinary Medicine, FDA*
- 11:15 – 11:45 a.m.** **Foreign Inspections – Compliance and Expectations**
Alonza Cruse, *Office of Regulatory Affairs, FDA*
- 11:45 – 12:00 p.m.** **Q&A/ Expert Panel Discussion**
Moderated by Michael Kerrigan, Ph.D., *CVM, FDA*
- 12:00 – 1:00 p.m.** **Lunch**
- 1:00 – 1:45 p.m.** **Selection of Active Pharmaceutical Ingredient (API) Starting Materials**
Michael Kerrigan, Ph.D., *Center for Veterinary Medicine, FDA*
Kevin Cheng, Ph.D., *Center for Veterinary Medicine, FDA*
Anna Kooser, Ph.D., *Center for Veterinary Medicine, FDA*
John Stanko, Ph.D., *Center for Veterinary Medicine, FDA*

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- 1:45 – 2:30 p.m.** **Suitability Assessment of Regulatory Starting Materials (RSMs)**
Dimitrios Zarkadas, Ph.D, *Director, Engineering, API Technology & Portfolio Management, Merck*
- 2:30 – 2:45 p.m.** **Q&A/ Expert Panel Discussion**
Moderated by Brian Wachter, MBA, *Boehringer-Ingelheim*
- 2:45 – 3:00 p.m.** **Afternoon Break**
- 3:00 – 3:45 p.m.** **Drug Shortages and Medically Necessary Veterinary Products**
Susan Homire, DVM, *Center for Veterinary Medicine, FDA*
- 3:45 – 4:45 p.m.** **Supply Chain and Regulatory Requirements – An API Manufacturer’s Point of View**
Frank Jellen, Ph.D., *Regulatory Affairs, Excella*
- Drug Shortage / Economic Perspective**
Frank Amorese, *Senior Vice President, Animal Health, Flavine North America Inc.*
- 4:45 – 5:00 p.m.** **Q&A/ Expert Panel Discussion**
Moderated by Brian Wachter, MBA, *Boehringer-Ingelheim*
- 5:00 p.m.** **Day One Workshop Adjournment**
- 5:00 – 6:00 p.m.** **Networking Reception**
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DAY TWO: Thursday, July 19, 2018

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|--------------------------------|---|
| 8:30 – 9:00 a.m. | Registration & Coffee |
| | Opening Introduction
Courtney Tallman, <i>Program Chair, GADA</i> |
| 9:00 – 9:45 a.m. | Fundamental Principles of Developing and Maintaining Veterinary Master Files within cGMP Compliance for API Manufacturers
Herschel Gaddy, Ph.D., <i>President and CEO, Gaddy & Associates</i> |
| 9:45 – 10:30 a.m. | Import Alerts: An Overview of the Import Process
Dillard Woody, <i>Center for Veterinary Medicine, FDA</i>
Nawab Siddiqui, MS, MPA, <i>Center for Veterinary Medicine, FDA</i> |
| 10:30 – 10:45 a.m. | Q&A/ Expert Panel Discussion
Moderated by Stephanie Batliner, <i>Bimeda Inc.</i> |
| 10:45 – 11:00 a.m. | Morning Break |
| 11:00 – 11:45 a.m. | Quality and Safety of Inactive Ingredients Critical for Animal Health Drug Products
George B. Collins, Jr., <i>Vice President, Manufacturing, Vanderbilt Chemicals LLC</i> |
| 11:45 a.m. – 12:00 p.m. | Q&A/ Expert Panel Discussion
Moderated by Stephanie Batliner, <i>Bimeda Inc.</i> |
| 12:00 – 1:00 p.m. | Lunch |
| 1:00 – 1:30p.m. | USP to Present
Jennifer Devine, J.D., <i>Vice President, Global Legal Affairs, USP</i> |
| 1:30 – 2:15 p.m. | CVM Interaction with USP
Sohail Mosaddegh, <i>Senior U.S. Regulatory Affairs Manager, USP</i>
Sarai Obando, Ph.D., <i>Center for Veterinary Medicine, FDA</i> |
| 2:15 – 2:45 p.m. | USP/NF Monographs
Morgan Puderbaugh, <i>Senior Scientific Liaison-Chemical Medicines, USP</i> |
| 2:45 – 3:00 p.m. | Q&A/ Expert Panel Discussion
Moderated by Morgan Puderbaugh, <i>USP</i> |

USP Workshops

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3:00 – 3:15 p.m.

Afternoon Break

3:15 – 4:00 p.m.

USP Reference Standards

Doreen McDonald, *Director, Reference Standards Planning and Management, USP*

4:00 – 4:45 p.m.

USP-NF New Platform/Subscription Models

Frank (Trey) White, III, Ph.D., *Senior Director, Strategic Marketing & Program Operations-Documentary Standards, USP*

4:45 – 5:00 p.m.

Q&A/ Expert Panel Discussion

Moderated by Morgan Puderbaugh, *USP*

5:00 p.m.

Day Two Workshop Adjournment