2013 AAVPT BIENNIAL SYMPOSIUM (TENTATIVE AGENDA)

Location: Bolger Conference Center, Potomac, MD

Time	Title	Speaker
MONDAY, MAY 20		
8:00-9:30 AM	ACVCP Business meeting	
Session I - AAVPT Biennial Symposium Opens		
9:30-10:30 AM	Welcome	Sanja Modric, AAVPT President
	Keynote Speaker	Congressman Ted Yoho
10:30-11:00 AM	Break	
11:00-11:30 AM	USP Activities	Sanja Modric, FDA-CVM
11:30-12:00 PM	USP Activities	Morgan Puderbaugh, USP
12:00-1:00 PM	Lunch	
Sesson II - Current Challenges - Small Molecule Bioanalytics		
1:00-1:30 PM	Current Issues in Bioanalytical Method Validation and Study Operations – Test and Analytical Facility Perspectives	Jonathan Hare, Kingfisher Intl
1:30-2:00 PM	Current Issues in Bioanalytical Method Validation and Sample Analysis – Perspectives from a High Throughput Facility	Cory Nehls, Bioanalytical Services, PPD, INC.
2:00-2:20 PM	Bioanalytical Method Validation Review: Common Problems	Olutosin Idowu, FDA-CVM
2:20-2:50 PM	Break	
2:50-3:10 PM	Test Facility & Regulatory Review: Common Issues in Bioanalytical Method Validation and Incurred Sample Analysis	John Kadavil, CVM-FDA
3:10-4:00 PM	Bioanalytical Method Validation Question & Answer	
Special Session - AAVPT Biennial Abstracts and Poster Presentation		
4:00-5:00 PM	Student Presentations	
5:00-7:00 PM	Poster Session and Wine & Cheese Reception	

TUESDAY, MAY 21

8:00-9:00 AM	AAVPT business meeting	
Session III - Education Session		
9:00-10:30 AM	Learning Outcomes for Veterinary Pharmacology: The 2012 Teaching Workshop and Beyond	Virginia Fajt, Texas A&M University
10:30-11:00 AM	Break	
Session IV - Pharmacometrics		
11:00-12:00 PM	How modeling and Simulation can Guide the Drug Approval Process and Improve Clinical Trial Design	Joga Gobboru, Univeristy of Maryland
12:00-1:00 PM	Lunch	
1:00-1:45 PM	The Application of Modeling and Simulation for Animal Health (Therapeutics): Gaps, Challenges and Opportunities	Jim Riviere, Kansas State University
1:45-2:05 PM	Physiologically-based Modeling to Optimize Oral Drug Delivery and/or Identifying Drug-drug Formulation Interactions	Marilyn Martinez, FDA-CVM
2:05-2:35 PM	Simcyp Demo	Devendra Pade, Simcyp
2:35-3:00 PM	Break	
3:00-3:30 PM	The future of Population Pharmacokinetic Modeling for Veterinary Medicine	Tomas Martin-Jiminez, University of Tennessee
3:30-4:00 PM	The application of Stochastic Modeling to Predict Antimicrobial Efficacy	Virginia Fajt, Texas A&M University
4:00-4:30 PM	Mechanistic Modeling of Antimicrobial Efficacy and Resistance Development	Ronette Gehring, Kansas State University
Evening Session 5:00-7:00 PM	**Awards Banquet and VPRF Auction**	

WEDNESDAY, MAY 22

Session V - Labeling

8:00-8:25 AM	Improving the Understanding of Animal Drug Labels	Dorothy McAdams, FDA-CVM
8:25-8:50 AM	An Overview of Data Mining, Signal Detection, and Causality Assessment	John Baker, FDA-CVM
8:50-9:20 AM	CDER's Experience with Labeling	TBD, Center for Drug Evaluation and Research, FDA
9:20-9:50 AM	Companion Animal Perspective on Labeling	Ed Fallin
9:50-10:10 AM	Break	
10:10-10:40 AM	Food Animal Veterinarians' Perspective on Labeling	Richard Doak, Mid Maryland Dairy Veterinarians
10:40-11:10AM	Equine Practitioner's Perspective on Drug Labeling	Richard R. Fofa
11:10-11:30 AM	Impact of labeling changes on Industry	Ian Williamson
11:30-12:00 AM	Round-table Discussion	
12:00 PM	**Biennial Adjourned**	
12:00-1:00 PM	Lunch for all registrants	

2013 AAVPT/CVM ANIMAL DRUGS IN FEEDS WORKSHOP

Location: Bolber Conference Center, Potomac, MD

WEDNESDAY AFTERNOON, MAY 22

Session I

1:00-1:10 PM	Medicated Feed Workshop Introduction, Logistics	Bernadette Dunham, FDA-CVM
1:10-1:30 PM	Introducing Center for Veterinary Medicine – Organization and Structure	Sanja Modric,FDA-CVM
1:30-1:55 PM	New Animal Drugs for Use in Animal Feeds: Types and Categories	Dragan Momcilovic,FDA-CVM
1:55-2:15 PM	Development and Approval of Labeling for Medicated Feeds	Suzanne Sechen, FDA-CVM
2:15-2:30 PM	Where to Find Approved Labeling for Medicated Feeds	Dragan Momcilovic and Suzanne Sechen, FDA-CVM
2:30-2:55 PM	Veterinary Feed Directive	Jim Delaney, FDA-CVM
2:55-3:15 PM	BREAK	
3:15 - 3:45 PM	Antimicrobial Drugs in Animal Feeds: Update on FDA's Judicious Use Strategy	Craig Lewis, FDA-CVM
3:45- 4:05 PM	Understanding ADAA Combinations	Dan Benz, FDA-CVM
4:05-4:20 PM	Generic Approval Process and Biowaiver Requirements	Charli Long, FDA-CVM
4:20-5:00 PM	Round Table Discussion	

THURSDAY MORNING, MAY 23

Session II	Biomass Drug Substances	Amy Simms,FDA-CVM
8:00-8:30 AM	Type A Medicated Articles and CMC Issues	James Nitao. FDA-CVM
8:30-9:00 AM	CMC Data to Support Medicated Feeds	Heather Longstaff, FDA-CVM
9:00-9:30 AM	Medicated Feed Assay Methods and Limits: Regulatory Perspective	Asif Rasheed, FDA-CVM
9:30-9:45 AM	CMC Question & Answer	
9:45-10:00 AM	BREAK	
10:00-10:15 AM	What Do Feed Mill Inspections Have to do with Drug Approvals?	Shannon Jorde, FDA-CVM
10:15-10:40 PM	Interactions with CVM	Cory Herr, Elanco
10:40-11:10 AM	Round Table Discussion	
11:10-11:40 AM	Wrap Up and Next Steps	
11:40-12:00 PM	**Workshop Adjourned**	
12:00 - 1:00 PM	Lunch for all workshop registrants	