EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

2017 EMA Veterinary Medicines Info Day
Organised in collaboration with IFAH-Europe

'The latest developments' in scientific review, regulation and marketing authorisation procedures

16-17 March 2017, European Medicines Agency (EMA), London

Final Programme

Thursday 16th March 2017

13:30	**Registration opens**
14:00	**Introduction and welcome**

Session I: Scientific developments — Chair: **Rick Clayton** (IFAH-Europe)

14:10	**CVMP achievements 2016**: Highlights and reflections (30') **David Murphy** (Chair of CVMP)
14:40	**Anthelmintic resistance**: Status including feedback from the focus group meeting 13 June 2016 and presentation on the CVMP Reflection paper (20') **Gesine Hahn** (EU expert, Bundesamt für Verbraucherschutz und Lebensmittelsicherheit)
15:00	**Antimicrobial resistance (AMR)**: Presentation on the joint EMA/EFSA RONAFA opinion; harmonisation of SPCs of antimicrobial veterinary medicines under current legislation (20') **Helen Jukes** (Chair of CVMP Antimicrobials working party)
15:20	**Recent developments in environmental assessment** (20') **Chris van den Eede** (Zoetis/IFAH-Europe)
15:40	Questions (20')
16:00	***Coffee break (30') – EMA info points***

Session II: Availability of veterinary medicines — Chair: **Fia Westerholm** (EMA)

16:30	**Vaccine availability**: - Action plan (20') **Faye Ioannou** (EMA) - Industry perspective (action plan and progress following 2015 workshop) (20') **Frédéric Descamps** (Zoetis/IFAH-Europe)
17:10	**Impact of future legislation on availability** (20') **Pablo Hervas** (Veterindustria/IFAH-Europe)
17:50	Questions (20')

18:00	***Cocktail reception at The Pearson Room Restaurant, followed by walking buffet from 18:45***

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Friday 17th March 2017

Session III: Innovation	Chair: **Roxane Feller** (IFAH-Europe)
09:00	**Horizon scanning**: General outcome from meetings with industry; Early support for novel types of products (15') **Fia Westerholm** (EMA)
09:15	**Innovation**: How to make legislation more supportive for innovation (15') **Peter Oostenbach** (MSD/IFAH-Europe)
09:30	**Update on ADVENT** and progress on Q&A documents: experience with pre-drafting consultations (10') **Minna Leppänen** (EMA)
09:40	Questions (15')
Session IV: EU Procedural updates	Chair: **Melanie Leivers** (EMA)
09:55	**Recent experiences with centralised procedure**: opportunities for development (10') **Teresa Potter** (EMA)
10:05	**Best practice guide on adherence to timetables**: status update (15') **Piotr Kozarewicz** (EMA)
10:20	Questions (15')
10.35	**EMA procedural updates** (15') **Emily Drury** (EMA)
10:50	Questions (15')
11:05	***Coffee break (30') – EMA info points***
Session V: Regulatory policy	Chair: **David Mackay** (EMA)
11:35	**Future Legislation: harmonisation of SPCs:** - Industry viewpoint (15') **Emmanuelle Royer (Caroline Toulemonde)** (Merial/IFAH-Europe)
11:50	Questions (10')
12:00	**Future Legislation: pharmacovigilance:** - Pharmacovigilance and product data (20') **Jos Olaerts** (EMA) - Industry Vision (20') **Tony Simon** (Zoetis/IFAH-Europe)
12:40	Questions (10')
Close of the meeting (5')	**Fia Westerholm** (EMA); **Rick Clayton** (IFAH-Europe)