European Crop Protection

Crop Protection

European Regulatory Conference

7th & 8th March 2018

Hotel Crowne Plaza

B-1000 Brussels

For hotel booking, see: https://aws.passkey.com/event/49492632/owner/7604429/home

	CONFERENCE TIMINGS
Wednesday:	
0900-1200	4 workshops
1200-1300	Lunch
1300-1830	Sessions 1 & 2
1830-2000	Panel Debate
2000-2200	Conference Dinner
Thursday:	
0830-1330	Sessions 3 & 4
1330 -	Lunch
1400-1530	Brexit workshop

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Day 1: Wednesday 7th March

08h15	***Registration***

Workshops

Human Health (& risk assessment developments	***Product efficacy & precision farming***		***Legal and intellectual property***	***Political and agricultural policy***
Workshop sponsored by: eurofins	Workshop sponsored by: SCC WE CARE FOR YOUR SUCCESS			
Chair: Vincent Dreze, Eurofins	***Chair: Norbert Weissmann, SCC***	*08h50*	*Chair:* ***Spyros Pappas, Pappas & Associates***	*Chair:* ***GraemeTaylor, ECPA***
Human health challenges for industry (e.g. CRA, genotox) – ***Phil Botham, Syngenta***	Precision farming developments ***Vik Vandecaveye, CNH Industrial***	09h00	In the battle for confidentiality – our way through the Courts - ***Gerardine Garcon, BASF***	Political challenges and communication in Withorwithout campaign – ***Philip Weiss, ZN Consulting***
		09h10		
		09h20	Voluntary Disclosure of Studies by Industry – Legal Aspects – ***Niklas Pieper, Bayer***	
EFSA update on HH – ***Andrea Terron***	Precision farming – what does it mean for the PPP uses and the approval process? ***Martin Schaefer, BASF***	09h30		Cumulative Impact Assessment – Update ***– Anne van Drunen-Little, Steward Redqueen***
		09h40	Recent case law on protection of regulatory data – ***Claudio Mereu, FieldFisher***	
		09h50		
Questions and Answers	Questions and Answers	10h00	Questions and Answers	Questions and Answers
Coffee Break	Coffee Break	10h10	Coffee Break	Coffee Break
		10h20		
		10h30		
Exposure assessment for epidemiological studies on PPPs: ***Karen Galea (IOM)***	Simplify efficacy for today and the future ***– Beth Hall, Syngenta***	10h40	Recent EU case law relevant for the PPP sector: ***Irene Antypas, Ashurst***	SUD, IPM and low-risk – Viewpoint of the bio-control industry. – ***David Cary, IBMA***
		10h50		
Worker and bystander/resident exposure models –Industry input - **Alistair Morriss, DowDuPont**	Simplify efficacy, today and the future – a minor use perspective – ***Jeroen Meeussen, MUCF***	11h00	Import tolerances – ***Patrick Reimer, Syngenta***	SMART resistance management project – ***Maria Torne, DowDuPont***
		11h10		
Tox21 / Risk21 – ***Tina Mehta, DowDuPont***	Adapting to the environment in product registrations: ***Dr. Joachim Kranz, SCC***	11h20	SPCs and EU Unitary patents – ***Bettina Wanner, Bayer***	Organic farmer's viewpoint - ***Stephen Parsley, UK***
		11h30		
Panel discussion with CAAT	Panel discussion With Cristina Alfaro, Suterra	11h40	Panel discussion	Panel discussion
		11h50		
Conclusions & close	Conclusions & close	12h00	Conclusions & close	Conclusions & close

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Day 1: Wednesday 7th March		
12h00	***Lunch***	
13h00	***Session 1: Overview of current challenges*** ***Chairman: Coralie van Breukelen-Groeneveld, Bayer***	
13h10	Welcome address - ***Jean-Philippe Azoulay, ECPA Director General***	
13h20	Update on implementation of Regulation 1107/2009 - ***Klaus Berend, DG SANTE***	
13h50	Regulation 1107/2009 - challenges for today and the future - ***Martyn Griffiths, Bayer***	
14h15	Main issues for national authorities in the AS and product evaluations? - ***Barbara Oliveira. Portugal***	
14h40	Opportunities to improve the legislation and implementation? - ***Pavel Minar, Czech Republic***	
15h05	Question and answer	
15h30	***Coffee break***	
	Session 2: Active substance evaluation update	
16h00	AS review programme - ***Wolfgang Reinert, DG SANTE***	
16h25	MS challenges in AS review evaluation: improving cooperation between Member States and EFSA – ***Luuk van Duijn, CTGB***	
16h50	EFSA update on active substances review – ***José Tarazona or Benedicte Vagenende, EFSA***	
17h15	The risk assessment output – what options for risk managers? – ***Martin Streloke, BVL, Germany***	
17h40	Key hurdles from assessment to decision making - ***Jane West, Syngenta***	
18h05	Panel discussion	
18h30	***Pre-debate drinks***	
19h00	***Evening debate: Does science still have a role in EU decision making on food policy issues?*** ***Chair: [tbc]*** ➢ Philippe Lamberts MEP ➢ Hubert Deluyker ➢ Paul Leonard, BASF	
20h00	***Conference dinner***	Sponsored by: GAB

Day 2: Thursday 8th March	
	Session 3: Product evaluation challenges
08h30	Chairman's introduction
08h35	Article 43 - where's the silver bullet? - ***Christian Prohaska, AGES, Austria***
09h00	South zone update – ***J-L Alonso Prados (tbc)***
09h25	Northern zone update - ***Vibeke Moeller, Denmark***
09h50	Industry view on key challenges in product authorisation – ***Jeanne Roederer, ADAMA***
10h10	Challenges and focus of the Post approval issue group – ***Darren Flynn, CRD***
10h25	Panel discussion
10h45	***Coffee break***
	Session 4: Science update & Legislative review
	Science update
11h15	Update on endocrine disruption and other scientific guidance issues - ***Karin Nienstedt, DG SANTE***
11h40	Scientific developments likely to impact on regulatory implementation - ***Gábor Tőkés, Hungary***
12h00	Industry viewpoint on guidance documents under discussion – ***Peter Campbell, Syngenta***
12h20	Panel discussion
	Legislative review
12h40	Introduction to the panel discussion by chairman
12h50	Panel discussion with: ➢ Klaus Berend ➢ José Tarazona ➢ Luuk van Dijn ➢ Industry
13h25	CONCLUSION AND CLOSE
13h30	***Lunch***
	Workshop: Likely implications of Brexit
14h00	Introduction
14h05	Current status, impact on EU process and likely implications in PPP evaluations – ***Dave Bench, CRD***
14h35	The industry viewpoint in the UK – ***Janet Williams, Bayer & Beth Cannell, CPA***
15h00	Panel discussion; Questions and answers
15h30	Close