



### **Scientific Colloquium Series**

### **EFSA Scientific Colloquium N°17**

on low dose response in toxicology and risk assessment 14 -15 June 2012 | Parma, Italy

## Updated draft programme 12.06.2012

### **Overall chairs**

Robert Luttik, National Institute of Public Health and the Environment, NL Alexandre Feigenbaum, European Food Safety Authority, IT

#### **Overall rapporteurs**

Wim Mennes, National Institute of Public Health and the Environment, NL Anna F. Castoldi, European Food Safety Authority, IT

### Thursday 14 June

08.30-09.00	Registration participants
08.15-08.45	Briefing meeting with all chairs and rapporteurs
09.00-13.00	SESSION 1: INTRODUCTORY PLENARY SESSION
09.00-09.10	Welcome and introduction to EFSA Claudia Heppner, European Food Safety Authority, IT
09.10-09.20	Objectives of the Colloquium Robert Luttik, National Institute of Public Health and the Environment, NL
09.20-09.30	"Report on Pew, Nature and IFT cosponsored workshop on Non-Monotonic Dose Responses: Relevance and Implications for Food".  Maricel Maffini, The Pew Health Group, US
09.30-09.50	Nature of an effect: adverse or non-adverse?  David Bell, European Chemicals Agency, Fl
09.50-10.00	Questions
10.00-10.20	Dose response relationships: biological and modeling aspects  Jason Aungst, U.S. Food and Drug Administration, US
10.20-10.30	Questions
10.30-10.50	Low dose effects: is the lowest the most relevant?  Dieter Schrenk, University of Kaiserslautern, DE
10.50-11.00	Questions
11.00-11.20	COFFEE / TEA BREAK
11.20-11.40	When the dose doesn't make the poison: low dose effects and endocrine disrupting chemicals Laura Vandenberg, Tufts University, US
11.40-11.50	Questions

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11:50-12:10	Low dose effects - impact for risk assessment Iona Pratt, Food Safety Authority of Ireland, IE				
12:10-12:20	Questions				
12.20-12.40	The Hormetic Dose Response  Edward Calabrese (via video-conference), University of Massachusetts, US				
12:40-12:50	Questions				
12.50-13.00	Introduction to Discussion Groups Stef Bronzwaer, European Food Safety Authority,	IT			
13.00-14.00	LUNCH				
14.00-18.00	SESSION 2: DISCUSSION GROUPS (DG)				
DG 1	Nature of an effect: adverse or non-adverse?	Chair:	Susanne Hougaard Bennekou, Danish EPA, DK		
		Rapporteur:	Trine Husøy, Norwegian Institute of Public Health, NO		
DG 2	Dose response relationships	Chair:	George Loizou, Health & Safety Authority, UK		
		Rapporteur:	Ursula Gundert-Remy, Charité, DE		
DG 3	Low dose effects: is there sufficient evidence for non-monotonic dose response curves?	Chair:	Paul Brantom, Brantom Risk Assessment Ltd, UK		
		Rapporteur:	Christophe Rousselle, ANSES, FR		
DG 4	Impact for risk assessment	Chair: Rapporteur:	Anthony Hardy, UK Fernando Aguilar, ANSES, FR		
16.00	COFFEE/TEA BREAK				
20.00	DINNER				

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### Friday 15 June

09.00-10.00	SESSION 3: CONTINUATION OF DISCUSSION GROUPS				
10.00-10.30	COFFEE/TEA BREAK				
10.30-13.30	SESSION 4: FINAL PLENARY SESSION				
	10:30-10:50	Report back from Discussion Group 1	Trine Husøy, Norwegian Institute of Public Health, NO		
	10:50-11:05	Discussion	rieditii, NO		
	11:05-11:25	Report back from Discussion Group 2	Ursula Gundert-Remy, Charité, DE		
	11:25-11:40	Discussion			
	11:40-12:00	Report back from Discussion Group 3	Christophe Rousselle, ANSES, FR		
	12:00-12:15	Discussion			
	12:15-12:35	Report back from Discussion Group 4	Fernando Aguilar, ANSES, FR		
	12:35-12:50	Discussion			
	12:50-13:20	General discussion			
	13:20-13:30	Take-home messages	Wim Mennes, National Institute of Public Health and the Environment, NL		

### 13.30 COLLOQUIUM ADJOURNS

### 13.30-14.30 Lunch

### **Organising Committee**

Stef Bronzwaer European Food Safety Authority, IT
Anna F. Castoldi, European Food Safety Authority, IT
Jean-Lou Dorne, European Food Safety Authority, IT
Alexandre Feigenbaum, European Food Safety Authority, IT
Djien Liem, European Food Safety Authority, IT
Robert Luttik, National Institute of Public Health and the Environment, NL
Luc Mohimont, European Food Safety Authority, IT
Iona Pratt, Food Safety Authority of Ireland, IE