Scientific Colloquium Series

13th EFSA Scientific Colloquium: *What's new on Novel Foods*

19-20 November 2009, Amsterdam

Programme

Overall chairs:	Bevan Moseley *NDA Panel member & Chair EFSA Working Group Novel Foods, UK*	Henk van Loveren *National Institute for Public Health and the Environment (RIVM), NL*
Overall rapporteurs:	Sandy Lawrie *Food Standards Agency, UK*	Wolfgang Gelbmann *European Food Safety Authority (EFSA)*

Thursday 19 November 2009

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**
Chair: Henk van Loveren

09.00-09.10 Welcome and introduction to EFSA
Juliane Kleiner, European Food Safety Authority (EFSA)

09.10-09.20 Objectives of the Colloquium
Henk van Loveren, National Institute for Public Health and the Environment (RIVM), NL

09.20-09.40 EFSA's current role on Novel Foods
Bevan Moseley, NDA Panel member & Chair EFSA Working Group Novel Foods, UK

09.40-09.50 Questions

09.50-10.10 Status of the revision of the Novel Food Regulation
Jean-Francois Roche, European Commission

10.10- 10.20 Questions

10.20-11.00 **COFFEE/TEA BREAK**

11.00-11.20 Current Scientific Committee on Food (SCF) guidance on Novel Foods
Karl-Heinz Engel, Technical University Munich, DE

11.20-11.30 Questions

11.30-11.50	Industry perspectives *Nigel Baldwin, Cantox (Consultant), UK*
11.50-12.00	Questions
12.00-12.20	Potential 'new area' such as nanotechnology *Rolf Hertel, Federal Institute for Risk Assessment (BfR), DE*
12.20-12.30	*Questions*
12.30-12.50	General discussion *Chairs*
12.50-13.00	Introduction to discussion groups *Stef Bronzwaer, European Food Safety Authority (EFSA)*

13.00-14.00 LUNCH

14.00-18.00 SESSION 2: DISCUSSION GROUPS (DG)
Four parallel discussion groups to address:

DG 1	History of (safe) use and traditional foods from non-EU-countries	Chair: Rapporteur:	*Seppo Salminen, University of Turku, FI* *Clemens van Rossum, Medicines Evaluation Board, NL*
DG 2	Data requirements and approach for anticipated intake	Chair: Rapporteur:	*Karl-Heinz Engel, Technical University Munich, DE* *Judith Buttriss, British Nutrition Foundation, UK*
DG 3	Key issues in absorption, distribution, metabolism, and excretion (ADME) studies, toxicology and allergenicity	Chair: Rapporteur:	*Annette Poeting, Federal Institute for Risk Assessment (BfR), DE* *Valeria Di Giorgi, Health Ministry, IT*
DG 4	Data requirements to demonstrate safety of foods derived by nanotechnology	Chair: Rapporteur:	*Hans Verhagen, National Institute for Public Health and the Environment (RIVM), NL* *Chris Jones, Food Standards Agency, UK*

16.00 COFFEE / TEA BREAK

18.00 End of Discussion Groups

20.00 NETWORKING DINNER

Friday 20 November 2009

09.00-10.00 SESSION 4: CONTINUATION OF DISCUSSION GROUPS

Including discussion on the outcomes of the discussion groups and the finalization of report back (presentation) to the plenary session

10.00-10.30 COFFEE/TEA BREAK
Rapporteurs to hand-in powerpoint presentation to registration desk for photocopying

10.30-13.30 SESSION 5: FINAL PLENARY SESSION
Chair: Bevan Moseley

10:30-10:50	Report back from discussion group 1	*Clemens van Rossum, Medicines Evaluation Board, NL*
10:50-11:05	*Discussion*	
11:05-11:25	Report back from discussion group 2	*Judith Buttriss, British Nutrition Foundation, UK*
11:25-11:40	*Discussion*	
11:40-12:00	Report back from discussion group 3	*Valeria Di Giorgi, Health Ministry, IT*
12:00-12:15	*Discussion*	
12:15-12:35	Report back from discussion group 4	*Chris Jones, Food Standards Agency, UK*
12:35-12:50	*Discussion*	
12:50-13:30	General discussion	
	Conclusions and take-home messages	*Chairs*

13.30 COLLOQUIUM ADJOURNS

14.30-15.00 Debriefing meeting with overall chairs and overall rapporteurs