

Parma, 23 November 2016

EFSA Info session - technical meeting with stakeholders on supplementary guidance for allergenicity assessment of GM plants

23 November 2016

Location: EFSA, Meeting rooms (07/08), Parma (Italy)

Agenda

23 November 2016		
Time	Subject	Speaker
8:00-8:30	Registration	
8:30-8:45	Welcome and scope of the meeting	E. Waigmann , Head, EFSA GMO Unit
8:45-9:00	Introduction to the EFSA guidance document on allergenicity assessment of GM plants	J.M. Wal , GMO Panel, Allergenicity WG/ INRA, France
9:00-9:20	<i>In vitro</i> protein digestibility test for allergenicity assessment – EFSA proposal	C. Mills , Allergenicity WG/ University of Manchester, UK
9:20-09:35	<i>In vitro</i> digestibility testing – Infogest	A. Mackie , University of Leeds, UK
9:35-9:45 9:45-9:55	<i>In vitro</i> protein digestibility testing – feedback from stakeholders after public consultation	R. van Ree , University of Amsterdam/ representative: ILSI-HESI; F. Jacquemart , GIET (Groupe International d'Études Transdisciplinaires)
9:55-10:15	<i>Coffee break</i>	
10:15-11:10	<i>In vitro</i> protein digestibility testing – plenary discussion	
11:10-11:30	Endogenous allergenicity – EFSA proposal	R. Selb , EFSA GMO Unit

11:30–11:45	Endogenous allergenicity – clinical aspects	P. Eigenmann , Allergenicity WG/ University of Geneva, Switzerland
11:45–11:55 11:55–12:05	Endogenous allergenicity – feedback from stakeholders after public consultation	R. Crevel , Unilever/ Focus Group representative: FoodDrink Europe; S. Schnadt , Focus Group patient's representative: DAAB (German Allergy and Asthma association)
12:05–13:05	<i>Lunch break</i>	
13:05–14:00	Endogenous allergenicity – plenary discussion	
14:00–14:20	Non – IgE adverse immune reactions (celiac disease)– EFSA proposal	F. Koning , Allergenicity WG/ Leiden University, The Netherlands
14:20–14:35	Non – IgE adverse immune reactions (celiac disease) – <i>in silico</i> tools	P. Rougé , Université Paul Sabatier, France/ Focus Group representative France
14:35–14:55	<i>Coffee break</i>	
14:55–15:05 15:05–15:15	Non – IgE adverse immune reactions (celiac disease) – feedback from stakeholders after public consultation	Kathryn Miller , Celiac UK/ patients' representative; Andre Silvanovich , Monsanto/ industry representative
15:15–16:10	Non -IgE adverse immune reactions (celiac disease) – plenary discussion	
16:10–16:30	Concluding remarks	A. Fernandez Dumont , EFSA GMO Unit