

Colloquium: What's new on Novel Foods

13th EFSA Scientific Colloquium: *What's new on Novel Foods - 1*9-20 November 2009, Amsterdam Henk van Loveren, NDA Panel Vice Chair, Novel Food Working Group member





...food (ingredients) which have not been used for human consumption to a significant degree within the EU before 15 May 1997 - Regulation (EC) N° 258/97)

Data Requirements



- I Specification of the NF
- II Effect of the production process
- III History of the organism used as the source of the NF
- IV Anticipated intake/extent of use of the NF
- V Previous human exposure to the NF or its source
- **VI** Nutritional information
- **VII** Microbiological information
- **VIII** Toxicological information



Objectives of the Colloquium



• To bring together <u>international experts</u> and interested parties from different sectors for an open scientific debate on <u>key issues</u> of the Novel Foods Regulation that will serve as input for the preparation of an updated Guidance for applicants.

• Discussions will focus on <u>various aspects</u> in the safety assessment of Novel Foods such as <u>history of (safe) use</u>, traditional foods from countries outside the EU, <u>intake</u> assessment, <u>toxicological</u> data requirements and emerging sciences such as <u>nanotechnology</u>.

Programme



- EFSA's current Role on Novel Foods
- Status of the Revision of the Novel Food Regulation
- Current Scientific Committee on Food (SCF) Guidance on Novel Foods
- Industry perspectives
- Potential 'new area' such as Nanotechnology

Break out groups



- History of (safe) use and traditional foods from non-EU-countries
- Data requirements and approaches for anticipated intake
- Key issues in absorption, distribution, metabolism, and excretion studies, toxicology and allergenicity
- Data requirements to demonstrate safety of foods derived by nanotechnology



Input for the preparation of an updated Guidance for applicants