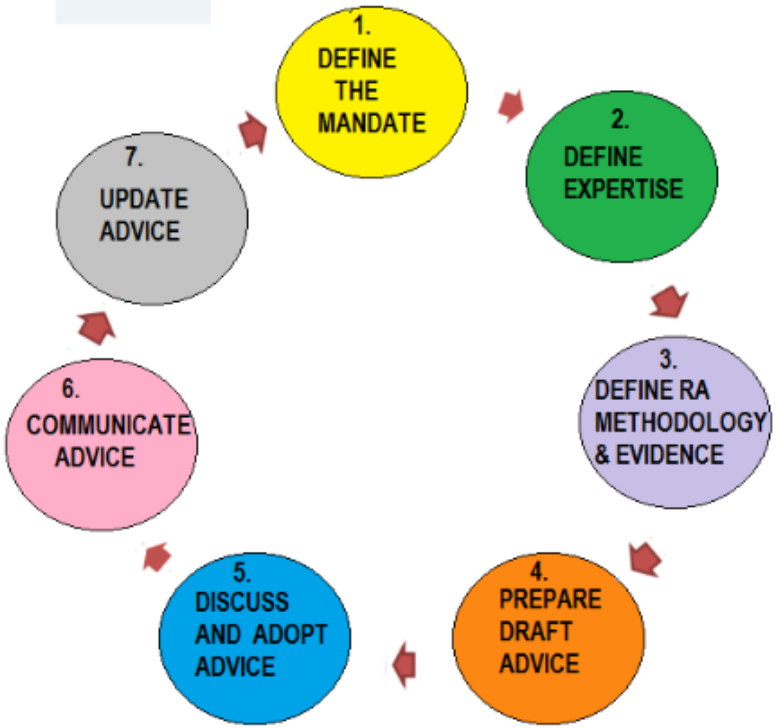
Transparency and Engagement in Risk Assessment

Management Board Meeting
3 December 2015

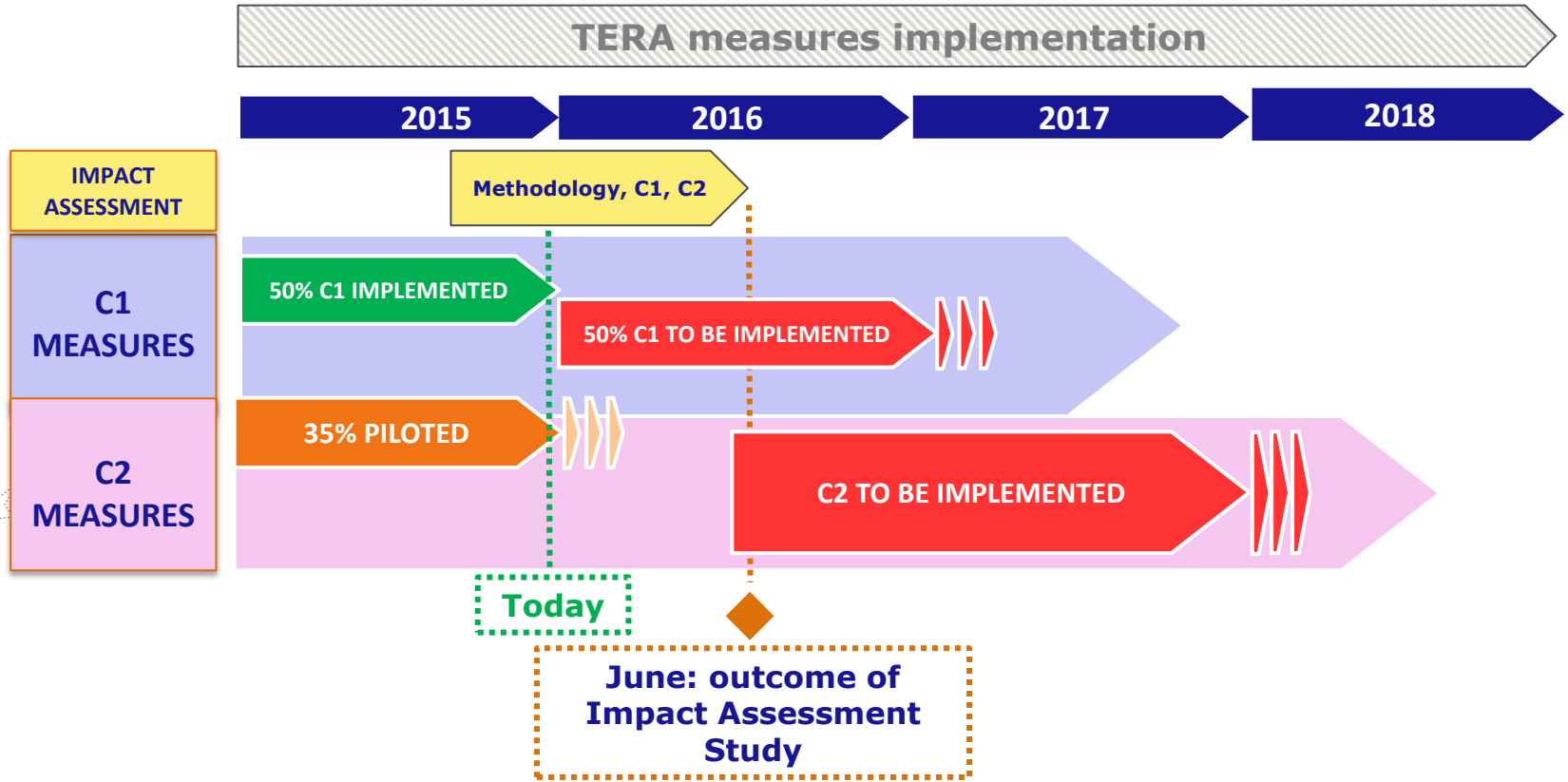
MARCH 2015 MB MEETING IMPLEMENTATION PLAN

35 Measures, applicable to the R.A. cycle, categorised according to:

- Readiness for implementation
- Potential impact on EFSA's organisational set up and/or resources

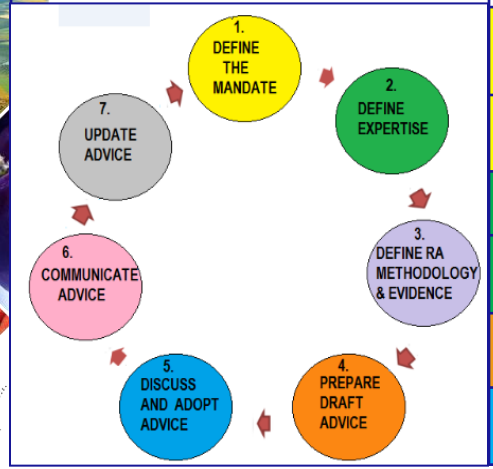


TERA PROJECT OVERALL TIMELINE



C1 MEASURES IMPLEMENTATION

6/12 measures implemented (= 50%) @ 3 DEC. 2015

STEP IN THE R.A. CYCLE	DESCRIPTION
	1.2 Simplification of requirements to take active role in public consultations
	1.3 Pre-notify interested parties of forthcoming public consultation
	2.1 Publish full biographies
	2.2 Documentation of the criteria of selection of WG members
	4.11 Consistent decision making on confidentiality of application dossiers
	5.1 Open Panel plenary meetings extended by half a day/year/panel

Full list of measures: http://www.efsa.europa.eu/sites/default/files/assets/150319-ax5.pdf

EVIDENCE

CURRICULUM VITAE

19 June 20

Title and name
Prof. dr. Jan Alexander

Nationality
Norwegian

Panel / Scientific Committee
Scientific Panel on Contaminants in the Food Chain

Education
EUROTOX Registered Toxicologist reconfirmed 2012; Norwegian Medical Association, Specialist in Occupational Medicine, 1993; University of Oslo, PhD, 1983. Toxicology; Ministry of Health Authorisation as Physician (General Practitioner) 1975; University of Oslo, MD, 1973. Medicine

Work Experience

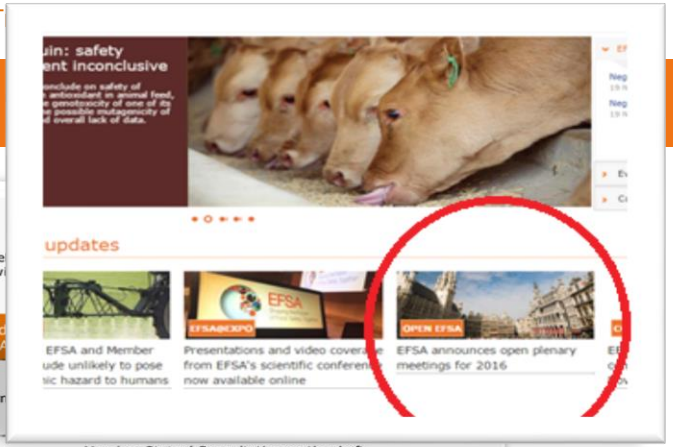
(2009 – present)	Norwegian Institute of Public Health, Oslo.	Deputy Director-General. Environmental Medicine, international relations
(1983 – 2009)	Norwegian Institute of Public Health, Oslo.	Division Director Division of Environmental Medisine. Department Director, Dept. of Food

Public consultation forecast

Latest updated date: 20/11/2015

This draft planner of upcoming consultations aims to facilitate stake
provided in the planner is indicative and subject to change in line wi
launched on the EFSA website.

Unit	Mandate Number	Question Number	Food		
Food Ingredients and Packaging	M-2015-0173	EFSA-Q-2015-00515	Enzyr		
GMO	M-2015-0155	EFSA-Q-2015-00654	GMO generic	Member States' Consultation on the draft Guidelines on possible derogation of existing requirements for applications of genetically modified (GM) food and feed at low levels submitted under Regulation (EC) No 1829/2003 on GM food and feed	21/06/2016
2015-0155		EFSA-Q-2015-00668	GMO generic	Public Consultation on the draft Guidelines on possible derogation of existing requirements for applications of genetically modified (GM) food and feed at low levels submitted under Regulation (EC) No 1829/2003 on GM food and feed GMO	21/12/2016



Public consultation on the draft general scientific guidance for stakeholders on health claim applications

Deadline: 11 September 2015

Document (755.87 KB)

Privacy statement (158.67 KB)

...

Interested parties are invited to submit written comments **by 11 September 2015**. Please use the electronic template provided to submit comments and refer to the line and page numbers. Kindly note that after 2 hours of non-activity your working session will expire and comments submitted after that time will not be recorded and transmitted. Therefore, if the page is left inactive for more than 2 hours, please re-open it from the link before restarting to comment. If you would like to submit additional data to support your comments or files send an email to:
NDA.PublicConsult.58@efsa.europa.eu. Please note that comments will not be considered if they:

SCIENTIFIC EVALUATION OF REGULATED PRODUCTS DEPARTMENT

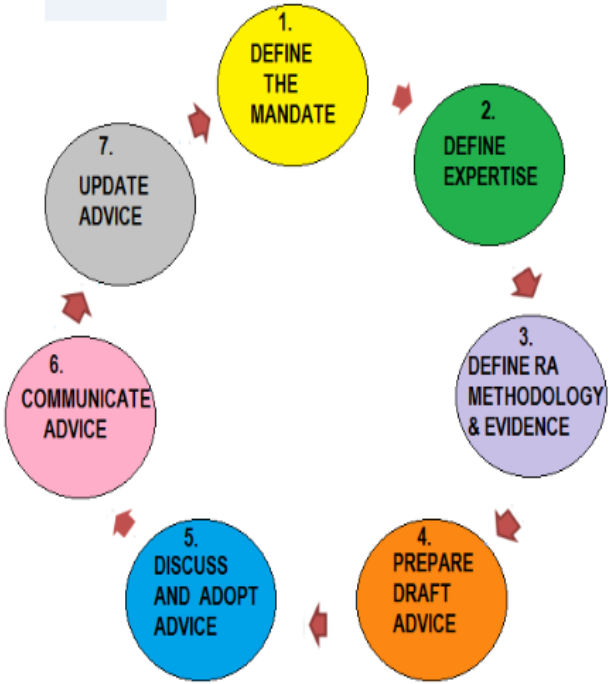
Parma, 25/09/2015

Note on the establishment of a Standing Working Group on Claims of the Scientific Panel on Dietetic Products, Nutrition and Allergies (2015-2018)

The Nutrition Unit is requested to manage the administrative and scientific activities of the Standing Working Group on Claims. The Nutrition Unit is also requested to organise the meetings of the Standing Working Group and assist in

MEASURES USED AD HOC OR PILOTED (C1 + C2)

10/29 measures already tested by EFSA (= 35%)

STEP IN THE R.A. CYCLE	DESCRIPTION
	3.2 Open and/or targeted call for data/information
	3.3 Consultation on the call for data/information format
	4.2 Proactive release of data/information in a readable/reusable format
	4.3 Increased accessibility to key data packages of MSs
	4.4 More feedback on the extent and on the reasons why certain data were/were not used
	4.6 Public consultation on draft opinions
	4.10 Increase transparency of the weight of evidence approach
	4.12 Transparency on the identification of key studies and detailed reasons to discard studies which document harmful effects
	6.2 Publication of applied assessment methodologies
	6.5 Structured process allowing post comments on opinions

UPDATE ON IMPACT ASSESSMENT

OUTPUTS

- **Methodology,** developed together with EFSA, to be applied to the 35 measures
- **Preliminary** impact assessment for all 35 measures
- **Fully-fledged** impact assessment for the 10 prioritised measures

TIMELINE

- December-January: **interviews** with stakeholders
- January-March: **survey** (on-line)
- Mid-May: **findings and conclusions**

NEXT STEPS

- **March 2016**:
  - EFSA presents the new Stakeholder Engagement Approach to the Management Board
- **May 2016**:
  - Impact Assessment study outcome
- **June 2016**:
  - Update to the Management Board on progress with C1 measures implementation
  - Proposal to the Management Board of prioritisation of C2 measures implementation