IVTIP In Vitro Testing Industrial Platform

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# Tentative Agenda

## IVTIP Spring 2013 Meeting

## '2013: State of the art on alternatives from an industrial point of view: ready for regulation?'

### May 15-16, 2013, Southampton, UK

*(May 14: IVTIP Business Meeting for IVTIP members only.*
*Agenda May 14 to be distributed to IVTIP members by email separately)*

**Host: British American Tobacco, Southampton, UK**

**Meeting location:**
**British American Tobacco Group Research & Development**
**Regents Park Road**
**Millbrook, Southampton SO15 8TL, United Kingdom**
**Tel: + 44 23 8077 7155**
**Meeting room: auditorium**

**Deadline for abstract submission/registration has been extended to March 27**

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# IVTIP Spring 2013 Meeting

**Day 1: May 15, 2013**

09:00h: Welcome
David O'Reilly, Science Director, British American Tobacco, Southampton, UK

09.10h: Welcome
Bart De Wever, ALTEXA Development, Monaco
IVTIP Board/Executive Secretary

## Morning session: Current Perspectives: from 3R opinion leaders

09.15 - 10.00 h: Perspectives from ESTIV (30 min and 15 min discussion)
Chantra Eskes, President ESTIV

10.00 - 10.45 h: Perspectives from CAAT (30 min and 15 min discussion)
Thomas Hartung, Director, CAAT, Baltimore, USA

10.45 - 11.15 h: Coffee break

11.15 – 12.00 h: Perspectives from EURL-ECVAM (30 min and 15 min discussion)
Elisabeth Berggren, EURL-ECVAM, Ispra, Italy

12.00 – 12:45 h: Perspectives from IVTIP (30 min and 15 min discussion)
Erwin Roggen, President IVTIP

12.45 – 14:30 h: Lunch

## Afternoon session: Paradigms for product testing, industry perspectives

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|---|---|
| 14.30 - 15.15 h: | Title (30 min and 15 min discussion)<br>Christopher Proctor, British American Tobacco, Southampton, UK |
| 15.15 - 16.00 h: | Title (30 min and 15 min discussion)<br>Shaun White, Food and Environment Agency, UK<br>to be confirmed |
| 16.00 – 16.30 h: | Coffee break |
| 16.30 – 17.15 h: | Assessing consumer product safety without animals: progress and challenges (30 min and 15 min discussion)<br>David Basketter, DABMED Consultancy Ltd, UK |
| 17.15 – 18.00 h: | Title (30 min and 15 min discussion)<br>Kate Willet, Human Society of the United States, USA |
| 18.00 h: | Round-up |
| 18.15 h: | Closure of the meeting/ social event (dinner) |

**Day 2: May 16, 2013**

## Morning session : US and Europe and regulatory hurdles

09.15 – 10.00 h: Regulatory hurdles in the chemical industry (30 min and 15 min discussion)
Speaker from ECHA to be confirmed

10.00 – 10.45 h: Regulatory hurdles in the cosmetic industry (30 min and 15 min discussion)
Speaker from Cosmetics Europe, Brussels, Belgium
to be confirmed

10.45 – 11.15 h: Coffee break

11.15 – 12.00 h: The FDA point of view (30 min and 15 min discussion)
Donald Prater (Deputy Director Europe Office FDA)
to be confirmed

12.00 – 12.45 h: Regulatory hurdles in the food industry (30 min and 15 min discussion)
Daniela Mauritzi (EFSA) (AP) to be confirmed

12.45 – 14:30 h: Lunch

## Afternoon session : Complex toxicological endpoints and education

14.30 – 15.15 h: Human tissue dose-based in vitro testing (30 min and 15 min discussion)
Jos Bessems, RIVM, The Netherlands

**Oral presentations of the selected posters**

15.15 – 15.35 h: xxxx (15 min and 5 min discussion)
15:35 – 15:55 h: xxxx (15 min and 5 min discussion)

16.00 - 16.30 h: Coffee break

16:30 – 16:50 h: xxxx (15 min and 5 min discussion)
16:50 – 17:10 h: xxxx (15 min and 5 min discussion)
17:10 – 17:30 h: xxxx (15 min and 5 min discussion)

17.30 h: Round-up

17.45 h: Closure of the meeting