EUROPEAN COMMISSION
ENTERPRISE AND INDUSTRY DIRECTORATE-GENERAL

Resources Based, Manufacturing and Consumer Goods Industries
Chemicals Industry

DRAFT AGENDA

MEETING OF THE WORKING GROUP ON GOOD LABORATORY PRACTICE

BORSCHETTE CONFERENCE CENTRE
FROISSARTSTRAAT 36, 1040 BRUSSELS

19 MARCH 2014, 10:00-17:00

Meeting chair: Maurits-Jan Prinz,
DG Enterprise and Industry

#	Topic & comment	Time
1.	Welcome	10:00 – 10:05
2.	Approval of the agenda	10:05 – 10:10
3.	Approval of the report of the last meeting	10:10 – 10:15
4.	Update on the GLP monitoring programmes of EU Member States not participating in the OECD GLP working group	10:15 – 10:25
5.	Report of the outcome of the last OECD working group meeting.	10:25 – 10:40
6.	Preparation of the next OECD working group meeting	10:40 – 11:00
7.	Update on OECD document on the differences between GLP and ISO 17025	11:00 – 11:10
8.	Draft guidance on the GLP Requirements for Peer Review of Histopathology	11:10 – 11:30
9.	Introduction to the EU Network of Laboratories for the Validation of Alternative Methods and the draft guidance on Good In vitro Method Practice (GIVIMP)	11:30 – 12:10
Lunch break		12:10 – 13:40
10.	Update on EU legislation with GLP requirements	13:40 – 13:50
11.	Collaboration with receiving authorities: EFSA / ECHA / EMA / competent authorities for clinical trials	13:50 – 14:55
Coffee break		14:55 – 15:05
12.	Guest speaker from industry: president of the Society for Quality Assurance (SQA)	15:05 – 16:10
13.	Non-domestic GLP inspections, including interaction with Taiwan	16:10 – 16:20
14.	Report from the meeting on technical issues	16:20 – 16:45
15.	AOB	16:45 – 17:00

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