European Medicines Agency
Post-authorisation Evaluation of Medicines for Human Use

London, *18 September 2007*
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BRAINSTORMING MEETING ON THE PROVISION OF INFORMATION BY THE EMEA TO ITS STAKEHOLDERS

20 September 2007 - 09.45-16.00

EMEA conference room 2A, 2nd floor

AGENDA

Chairperson: N. Wathion

09.45- 10.00 Coffee and reimbursement arrangements

10.00 –10.30

I. INTRODUCTION

I.1 Welcome and objectives of the meeting
N. Wathion, Head of EMEA Unit for Post-authorisation Evaluation

I.2 Presentation of participants
All participants

10.30 - 12.30 (with coffee break)

II. INITIATIVES TO BE UNDERTAKEN IN THE CONTEXT OF THE FURTHER IMPLEMENTATION OF THE EMEA ROAD MAP

II.1 Presentation: Reflection paper on the further implementation of the EMEA Road Map in the field of provision of information to the EMEA's stakeholders
I. Moulon, Head of EMEA Medical Information Sector

II.2 Feedback from participants on experience with the provision of information by the EMEA
All participants

II.3 General discussion on activities to be undertaken by the EMEA in context of the further implementation of the EMEA Road Map
All participants

12.30 – 13.30 Lunch break

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13.30 – 16.00 (with coffee break)

III. REVIEWING CURRENT INFORMATION PRACTICES AND COMMUNICATION TOOLS

III.1 Presentation: Overview of current information practices and communication tools
A. Blaedel Lassen, EMEA Medical Information Sector

III.2 Presentation: Points to consider for restructuring of the EMEA website
J. García Burgos, EMEA Medical Information Sector

III.3 Feedback (incl. on current experience & expectations) and general discussion
All participants

IV. CONCLUSIONS AND NEXT STEPS
N. Wathion, Head of EMEA Unit for Post-authorisation Evaluation