CONFERENCE ON ENVIRONMENTAL RISK ASSESSMENT FOR HUMAN AND VETERINARY MEDICINAL PRODUCTS

To be held at the European Medicines Agency, London on 27-28 October 2005

Programme

Thursday 27th October 2005

12:00
- Registration at the ground floor
- Coffee and conference materials at meeting room 3A, 3rd floor

13:00 Moderator’s Opening Address
Moderator: Annika Wennberg, FAO, Italy

Session I: Exposure of the environment to veterinary and human medicinal products.
Chair: Kornelia Grein, EMEA

13:15
- Pharmaceuticals in the environment (abstract)
- Questions
- Current ‘state of the art’ and challenges in Exposure Assessment for Pharmaceuticals (ERAPharm) (abstract)
- Questions

14:55
- European Regulatory Situation – Veterinary Medicinal Products (abstract)
- VICH guidelines/ Technical Guidance Documents (abstract)

15:45 Tea and Coffee

16:10
- European Regulatory Situation – Human Medicinal Products
- CHMP Guideline (abstract)

16:50
- Comments from industry – Veterinary Medicinal Products (abstract)
- Questions

17:10
- Comments from industry – Human Medicinal Products (abstract)
- Questions

17:30 Discussion with panel
Friday 28th October 2005

Session III  Risk management / Mitigation – Too much or not enough?

Chair: Hans Hoogland, CVMP
(co) Chair: Martin van der Graaff, EFPIA

09:00  Regulators - National Competent Authority’s perspective

- NCA Germany (abstract)
- NCA UK (abstract)
- NCA The Netherlands
- Questions

Session IV  Future Developments / Impacts

10:00  Environmental risk assessment for pharmaceuticals - challenges from a scientific point of view (abstract)

- Questions

10:30  Tea and coffee


- Questions

11:25  Discussion with panel

Conference conclusions

12:10  Conference Summary

12:25  Conference Conclusions

12:40  Close