

2012 EMA/IFAH-Europe Info-day

First announcement - please book this date in your diary!



The latest developments

Scientific review, marketing authorisation
procedures and legislation

EMA (London), 8-9 March 2012

- **Schedule:** 8 March 2012 13:30 to 9 March 2012 13:00
- **Added value:** An opportunity for professionals to get first-hand information on current developments in the regulatory scene with good networking opportunities.
- **Fee:** €300 (same as every year)
- **Registration:** Please book this date in your diary! Registration forms will be circulated in January 2012.

Outline programme

The programme will follow the favoured 3 session format similar to recent years. The final topics have not been fixed, but examples of those topics being considered are illustrated below.

Session I: Scientific developments

- Antimicrobial resistance – latest activities
- Immunological topics – latest updates
- New technologies and innovative medicines – latest thinking on how to manage these

Session II: EMA procedural updates

- Experience in electronic submissions
- Implementation of the EMA Roadmap
- Latest activities and future plans at VICH
- Pharmacovigilance, databases, and signal detection

Session III: Future of veterinary medicinal product legislation

- Update from DG SANCO
- SPC harmonisation (joint CVMP/CMDv group)
- Progress on ideas for rationalisation of packaging
- Holistic approach to legislation influencing the control of antimicrobial resistance