# 2017 EMA Info-day

## First announcement - save this date in your diary!

## The latest developments

**Scientific review, regulation and marketing authorisation procedures**

EMA (London), 16-17 March 2017

- Schedule: 16 March 2017 13:30 PM to 17 March 2017 13:00 PM.
- Added value: An opportunity for professionals to get first-hand information on current developments in the regulatory scene with good networking opportunities.
- Registration: Please book this date in your diary! Registration is now open (see attached form).

# Outline Programme

The programme will include sessions covering scientific developments, procedural developments and regulatory policy. The topics included in the draft programme are illustrated below.

### Session I: Scientific Developments

- CVMP work plan for 2017
- Anthelmintic resistance
- AMR update
- PBT substances

### Session II: Availability of VMPs

- Vaccine availability action plan
- Impact of future legislation on availability

### Session III: Innovation

- Horizon scanning
- Innovation – what is industry's view
- Update on ADVENT
- Recent experiences with centralised procedure

### Session IV: EU Procedural Updates

- Transparency/communication policy
- Approach to standard terms
- EMA new VDiv organisation

### Session V: Regulatory Policy

- Future Legislation: harmonisation of SPCs
- Future Legislation: pharmacovigilance