

		Date	Responsible
1.	Heads of Medicines Agencies	05 - 06 July 2010	Xavier De Cuyper
		25 - 26 October 2010	Xavier De Cuyper
2.	Conferentie RIZIV - FAGG	23 - 24 September 2010	Els Geeraerts
3.	Competent Authorities for Medical Devices	23 - 24 September 2010	Frédérique Meulders / Anne Van Nerom
4.	Co-ordination Group for Mutual Recognition and Decentralised Procedure-Veterinary	27 - 28 September 2010	Christophe Debruyne
5.	Committee for Veterinary Medicinal Products	27 - 28 September 2010	Bruno Urbain
6.	Committee for Medicinal Products for Human Use	30 September - 01 October 2010	Pieter Neels
7.	Committee for Orphan Medicinal Products	30 September - 01 October 2010	André Lhoir
8.	Pediatric Committee	30 September - 01 October 2010	Jacqueline Carleer / Daniel Brasseur
9.	Scientific Advice Working Party	30 September - 01 October 2010	Bruno Flamion
10.	Committee for Advanced Therapies	30 September - 01 October 2010	Bruno Flamion
11.	Co-ordination Group for Mutual Recognition and Decentralised Procedure-Human	04 - 05 October 2010	Sophie Colyn
12.	HMA EMACOLEX	14 - 15 October 2010	Paul Ballegeer
13.	HMA Working Group of Enforcement Officers	18 - 19 - 20 October 2010	Roy Vancauwenberghe
14.	Pharmacovigilance Working Party	8 - 9 November 2010	Jean-Michel Dogné / Thierry Roisin
15.	Clinical Trials Facilitation Group	18 - 19 November 2010	Greet Musch
16.	HMA Quality Managers	26 November 2010	Josiane Van der Elst
17.	Committee on Herbal Medicinal Products	7 - 8 December 2010	Heidi Neef
18.	HMA Homeopathic Medicinal Products Working Group	9 - 10 December 2010	Marie-Anne Mouyart
19.	Working Group Communication Professionals	16 - 17 December 2010	Ann Eeckhout