22 December 2011
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Committee for Medicinal Products for Veterinary Use (CVMP)

CVMP Monthly report of application procedures, guidelines and related documents

December 2011

The CVMP monthly report includes statistical data for the current and previous two years on scientific advice, initial evaluations, variations, line extensions, renewals, MRLs initial evaluations and MRLs extensions/modifications and arbitration and referral procedures.

In addition, the report includes a summary table of the opinions issued by the CVMP in the current year and a list of adopted guidelines and other public documents.

A pre-notification checklist is being published with this report to assist marketing authorisation holders in their submissions of Type IA variations, as well as a template letter to cover all veterinary procedures.

Applications for medicinal products for veterinary use and maximum residue limits (MRLs)

Scientific advice requests					
	95-08	2009	2010	2011	Total
Submitted	69	11	21	26	127
Advice given	65	8	18	24	115

Initial evaluation					
	95-08	2009	2010	2011	Total
Full (Submitted)	110	14	16	8	148
Abridged/ generics (Submitted)	10	1	2	3	16
Withdrawals	12	0	1	0	13
Positive opinions	91	13	14	19	137
Negative opinions	1	0	0	0	1

Marketing authorisations					
	95-08	2009	2010	2011	Total
Granted	88	12	9	22	131
Withdrawals	2	0	4	1	7
Not renewed	2	0	0	0	2

Extensions					
	95-08	2009	2010	2011	Total
Submitted	60	12	3	7	82
Withdrawals	2	1	1	0	4
Positive opinions	40	7	8	4	59
Negative opinions	0	0	0	0	0

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Variations – applications submitted					
	95-08	2009	2010	2011	Total
Type IA	339	32	76	125	763
Type IB		41	63	87	
Type II	210	40	26	45	321
Transfers	11	3	8	3	25

Renewals					
	95-08	2009	2010	2011	Total
Submitted	50	18	7	14	89
Positive opinions	48	17	8	12	85
Negative opinions	0	0	0	0	0

Arbitrations and Community referrals					
	95-08	2009	2010	2011	Total
Referrals submitted	38	9	12	12	71
Opinions reached[1]	20	15 (5)	11 (1)	10	56 (6)

[1] Re-examination of opinions in brackets

Substances considered as not falling within the scope of Regulation (EC) No 470/2009		
	2011	Total
Submitted	7	7
Agreed	9	9
Scientific advice recommended	0	0

MUMS/ Limited market classification		
	2011	Total
Positive with financial incentives	8	8
Positive without financial incentives	12	12
Negative	1	1

Establishment of MRLs for new substances					
	95-08	2009	2010	2011	Total
Submitted	66	4	3	1	74
Withdrawals	5	0	0	0	5
Positive opinions[2]	54	2	2	4	62
Negative opinions[3]	7	0	0	0	7

Extensions / modifications/extrapolations of MRLs					
	95-08	2009	2010	2011	Total
Submitted	98	2	10	13	123
Withdrawals	4	0	0	2	6
Positive opinions[2]	113	3	3	12	131
Negative opinions	6	0	0	0	6

[2] Including opinions recommending the extension of the expiry date for provisional MRLS or definitive MRLs for substances with previously provisional maximum residue limits

[3] Including one opinion concluding that final MRL could not be established for a substance with provisional maximum residue limits previously established

CVMP opinions in 2011 on medicinal products for veterinary use

Positive opinions

Product • **Invented name** • **INN**	• **Marketing authorisation holder**	**Therapeutic area** • **Target species** • **Summary of indication**	**EMA/CVMP** • **Validation** • **Opinion** • **Active time** • **Clock stop**	**European Commission** • **Opinion received** • **Date of decision** • **Notification** • **Official Journal**
• CaniLeish	• Virbac S.A.	• Dogs • Vaccine against Leishmania infection	• 17/03/2010 • 12/01/2011 • 210 • 91	• 13/01/2011 • 14/03/2011 • 17/03/2011 • OJ C 184/15
• ZULVAC 1 + 8 Ovis	• Pfizer Limited	• Sheep • Vaccine for prevention of viraemia caused by Bluetongue Virus serotypes 1 and 8	• 18/03/2010 • 12/01/2011 • 180 • 119	• 13/01/2011 • 14/03/2011 • 17/03/2011 • OJ C 184/15
• BLUEVAC BTV8	• CZ Veterinaria S.A	• Cattle, sheep • Vaccine for active immunisation against bluetongue disease	• 17/01/2009 • 09/02/2011 • 210 • 543	• 10/02/2011 • 14/04/2011 • 18/04/2011 • OJ C 184/15
• Procox • Emodepside and toltrazuril	• Bayer Animal Health GmbH	• Dogs • Treatment of dogs when mixed parasitic infections, caused by certain specific roundworms and coccidia are suspected or demonstrated	• 16/02/2010 • 09/02/2011 • 210 • 148	• 11/02/2011 • 20/04/2011 • 28/04/2011 • OJ C 184/15
• Veraflox • Pradofloxacin	• Bayer Animal Health GmbH	• Dogs, cats • Treatment for dogs and cats with particular infections caused by certain specific and susceptible pathogens	• 19/05/2009 • 14/07/2010 • 205 • 217 • 09/02/2011 (re-consideration)	• 11/02/2011 • 12/04/2011 • 14/04/2011 • OJ C 184/15
• Zuprevo • Tildipirosin	• Intervet International BV	• Pigs, cattle • Treatment of bacterial infections in the respiratory tract in pigs and cattle	• 16/02/2010 • 08/03/2011 • 210 • 177	• 10/03/2011 • 06/05/2011 • 06/05/2011 • OJ C 250/16
• CERTIFECT • Fipronil, (S)-	• MERIAL SAS	• Dogs • Treatment and	• 16/03/2010 • 09/03/2011	• 10/03/2011 • 06/05/2011

Product • Invented name • INN	• Marketing authorisation holder	Therapeutic area • Target species • Summary of indication	EMA/CVMP • Validation • Opinion • Active time • Clock stop	European Commission • Opinion received • Date of decision • Notification • Official Journal
methoprene, amitraz		prevention of infestations with ticks, alone or in association with fleas and/or chewing lice	• 210 • 148	• 06/05/2011 • OJ C 250/16
• MS-H Vaccine • *Mycoplasma synoviae* strain MS-H	• Pharmsure Ltd	• Chickens • Vaccine to reduce air sac lesions and reduce the number of eggs with abnormal shell formation caused by *Mycoplasma synoviae*	• 15/12/2009 • 07/04/2011 • 206 • 271	• 08/04/2011 • 14/06/2011 • 14/06/2011 • OJ C 250/16
• Recuvyra • Fentanyl	• Nexcyon Pharmaceuticals Ltd	• Dogs • Control of post-operative pain associated with major orthopaedic and soft tissue surgery	• 16/12/2009 • 04/05/2011 • 210 • 294	• 05/05/2011
• Emdocam • Meloxicam	• Emdoka bvba	• Cattle, pigs, horses • For treatment in respiratory infections, diarrhoea and mastitis in cattle. For treatment in non-infectious locomotor disorders and in puerperal septicaemia and toxaemia in pigs. In horses for treatment in musculo-skeletal disorders as well for the relief of pain in equine colic.	• 18/05/2010 • 09/06/2011 • 175 • 211	• 09/06/2011 • 18/08/2011 • 22/08/2011 • OJ C 316/15
• Proteq West Nile • West Nile recombinant	• MERIAL	• Horses • Vaccine for the active immunisation of horses against	• 18/05/2010 • 09/06/2011 • 196 • 190	• 09/06/2011 • 05/08/2011 • 10/08/2011 • OJ C 316/15

Product • **Invented name** • **INN**	• **Marketing authorisation holder**	**Therapeutic area** • **Target species** • **Summary of indication**	**EMA/CVMP** • **Validation** • **Opinion** • **Active time** • **Clock stop**	**European Commission** • **Opinion received** • **Date of decision** • **Notification** • **Official Journal**
canarypox virus (vCP2017 virus)		West Nile disease		
• Zulvac 1 Bovis • Inactivated Bluetongue virus, serotype 1, strain BTV-1	• Pfizer Limited	• Cattle • Active immunisation of cattle for the prevention of viraemia caused by Bluetongue Virus, serotype 1	• 12/08/2010 • 09/06/2011 • 180 • 120	• 06/07/2011 • 05/08/2011 • 10/08/2011 • OJ C 316/15
• Zulvac 1 Ovis • Inactivated Bluetongue Virus, serotype 1, strain BTV-1	• Pfizer Limited	• Sheep • Active immunisation of sheep for the prevention of viraemia caused by Bluetongue Virus, serotype 1	• 15/07/2010 • 09/06/2011 • 179 • 148	• 06/07/2011 • 05/08/2011 • 10/08/2011 • OJ C 316/15
• Nobivac Myxo-RHD • Live myxoma vectored RHD virus strain 009	• Intervet International BV,	• Rabbits • Active immunisation of rabbits to reduce mortality and clinical signs of myxomatosis and to prevent mortality due to rabbit haemorrhagic disease	• 16/02/2010 • 14/07/2011 • 210 • 302	• 15/07/2011 • 07/09/2011
• Recocam • Meloxicam	• CF Pharma	• Cattle, pigs, horses • For treatment in respiratory infections, diarrhoea and mastitis in cattle. For treatment in non-infectious locomotor disorders and in puerperal septicaemia and toxaemia in pigs. In horses for treatment in musculo-skeletal disorders as well for the relief of pain in	• 16/03/2010 • 14/07/2011 • 210 • 274	• 14/07/2011 • 13/09/2011

Product • **Invented name** • **INN**	• **Marketing authorisation holder**	**Therapeutic area** • **Target species** • **Summary of indication**	**EMA/CVMP** • **Validation** • **Opinion** • **Active time** • **Clock stop**	**European Commission** • **Opinion received** • **Date of decision** • **Notification** • **Official Journal**
		equine colic.		
• TruScient • Dibotermin-alfa	• Pfizer Limited	• Dogs • For the treatment of diaphyseal fractures as an adjunct to standard surgical care using open fracture reduction	• 15/06/2010 • 13/10/2011 • 205 • 279	• 14/10/2011 • 14/12/2011
• Panacur AquaSol • Fenbendazole	• Intervet International B.V.	• Pigs • For the treatment and control of gastro-intestinal nematodes in pigs infected with Ascaris suum and Oesophagostomum spp.	• 12/10/2010 • 13/10/2011 • 202 • 163	• 13/10/2011
• Inflacam • Meloxicam	• Chanelle Pharmaceuticals Manufacturing Limited	• Dogs, horses, cattle, pigs • For the alleviation of inflammation and pain in both acute and chronic musculo-skeletal disorders.	• 15/06/2011 • 13/10/2011 • 120 • 0	• 13/10/2011
• Activyl Tick Plus • Indoxacarb, permethrin	• Intervet International B.V.	• Dogs • Treatment of flea and tick infestations	• 07/12/2010 • 10/11/2011 • 210 • 128	• 11/11/2011
• RevitaCAM 5mg/ml	• Abbott Laboratories Limited	• Dogs • For the alleviation of inflammation and pain in acute and chronic musculo-skeletal disorders	• 05/01/2011 • 08/12/2011 • 210 • 128	• 08/12/2011

CVMP opinions in 2011 on establishment of MRLs

Positive opinions

• **Substance** • **INN**	• **Target species**	**EMA/CVMP** • **Validation** • **Opinion** • **Active time** • **Clock stop**	**European Commission** • **Opinion received** • **Date of regulation** • **Official Journal**
• Methylpredni – solone (after provisional MRLs)	• Bovine	• n/a • 12/01/2011 • 90 • 0	• 27/01/2011
• Octenidine dihydrochloride	• All mammalian food producing species	• 11/08/2009 • 08/02/2011 • 210 • 246	• 21/02/2011
• Monepantel (after provisional MRLs)	• Caprine	• n/a • 09/03/2011 • 90 • 0	• 25/03/2011
• Azamethiphos	• Fin fish	• 21/02/2011 • 07/04/2011 • 45 • 0	• 08/04/2011
• Pegylated bovine granulocyte colony stimulating factor	• Bovine	• 16/03/2010 • 05/05/2011 • 210 • 205	• 18/05/2011
• Lasalocid	• Bovine	• 10/08/2010 • 05/05/2011 • 210 • 58	• 18/05/2011
• Ivermectin	• All mammalian food producing species	• n/a • 09/06/2011 • 176 • 0	• 20/06/2011
• Phenoxymethyl-penicillin	• Poultry eggs	• 12/10/2010 • 14/07/2011 • 210 • 65	• 22/07/2011
• Tildipirosin (after provisional MRLs)	• Bovine, porcine and caprine	• n/a • 15/09/2011 • 90 • n/a	• 29/09/2011

• Altrenogest	• Porcine, *equidae*	• n/a • 13/10/2011 • 129 • n/a	• 18/10/2011
• Neomycin	• All food producing species	• 14/09/2010 • 10/11/2011 • 210 • 212	• 16/11/2011
• Closantel	• Bovine and ovine milk	• n/a • 10/11/2011 • 83 • 0	• 16/11/2011
• Nitroxinil	• Bovine and ovine milk	• n/a • 10/11/2011 • 72 • 0	• 16/11/2011
• Triclabendazole	• All ruminants	• n/a • 10/11/2011 • 83 • 0	• 16/11/2011
• Fenbendazole	• Chicken and all food producing species except fish.	• 13/07/2011 • 08/12/2011 • 148 • 0	• 14/12/2011
• Clorsulon	• Bovine milk	• n/a • 08/12/2011 • 111 • 0	• 14/12/2011

Arbitrations and Community referrals in 2011

Type of referral	• **Date of clock start** • **CVMP opinion**	• **Product name** • **INN**
Referral under Art. 34 of Directive 2001/82/EC	• 11/11/2009 • 10/11/2011	• Fortekor vet and associated names • Benazepril hydrochloride
Referral under Art. 34 of Directive 2001/82/EC	• 14/04/2010 • 07/06/2011	• Synulox Lactating Cow and associated names • Amoxicillin, clavulanic acid, prednisolone
Referral under Art. 33(4) of Directive 2001/82/EC	• 14/07/2010 • 07/04/2011	• Combimox Lactating Cow • Amoxicillin, clavulanic acid, prednisolone
Referral under Art. 33(4) of Directive 2001/82/EC	• 14/07/2010 • 07/04/2011	• Nisamox Lactating Cow • Amoxicillin, clavulanic acid, prednisolone
Referral under Art. 33(4) of Directive 2001/82/EC	• 14/07/2010 • 07/04/2011	• Combisyn Lactating Cow • Amoxicillin, clavulanic acid, prednisolone

Type of referral	• Date of clock start • CVMP opinion	• Product name • INN
Referral under Art. 34 of Directive 2001/82/EC	• 14/07/2010 • 04/05/2011	• Doxycycline 50% WSP and associated names • Doxycycline hyclate
Referral under Art. 34 of Directive 2001/82/EC	• 14/07/2010 • 04/05/2011	• Doxyfar 50% WSP and associated names • Doxycycline hyclate
Referral under Art. 34 of Directive 2001/82/EC	• 09/11/2010	• Baytril 10% oral solution and associated names • Enrofloxacin
Referral under Art. 33(4) of Directive 2001/82/EC	• 09/02/2011 • 08/06/2011	• Clavudale 50 mg tablet for cats and dogs • Amoxicillin and clavulanic acid
Referral under Art. 35 of Directive 2001/82/EC	• 09/03/2011	• Veterinary medicinal products containing active substances belonging to the class of flukicides for which no MRL has been established in milk
Referral under Art. 35 of Directive 2001/82/EC	• 06/04/2011 • 13/10/2011	• All veterinary medicinal products containing systemically administered (parenteral and oral) 3rd and 4th generation cephalosporins and intended for use in food producing species • Cefquinome and ceftiofur
Referral under Art. 33(4) of Directive 2001/82/EC	• 04/05/2011	• Prontax 5 mg/ml pour-on solution for cattle • Doramectin
Referral under Art. 33(4) of Directive 2001/82/EC	• 04/05/2011	• Prontax 10 mg/ml solution for injection for sheep, cattle and pigs • Doramectin
Referral under Art. 35 of Directive 2001/82/EC	• 04/05/2011	• All pre-mixes for medicated feedingstuffs containing 40, 100 or 200 g tilmicosin per kg pre-mix • Tilmicosin
Referral under Art. 78 of Directive 2001/82/EC	• 04/05/2011 • 14/07/2011	• HIPRABOVIS PNEUMOS Emulsion for injection for cattle and associated names • Inactivated *Mannheimia haemolytica* and *Histophilus somni*
Referral under Art. 34 of Directive 2001/82/EC	• 14/09/2011	• Milaxyn Plus, Strantel Plus, Prazical Plus, Voxical Plus, Exitel Plus, Cazitel Plus and Prazitel Plus and associated names

Type of referral	• Date of clock start • CVMP opinion	• Product name • INN
		• Praziquantel, pyrantel and febantel
Referral under Art. 35 of Directive 2001/82/EC	• 15/09/2011	• All long acting formulations for injection containing barium selenate for all food producing species • barium selenate
Procedure under Art. 30(3) of Regulation (EC) No 726/2004	• 15/09/2011	• N/a • Dapsone
Procedure under Article 33(4) of Directive 2001/82/EC	• 12/10/2011	• Nuflor 300 mg/ml solution for injection for cattle and sheep • Florfenicol
Procedure under Article 35 of Directive 2001/82/EC	• 12/10/2011	• Hipralona Enro-S and its generics • Enrofloxacin

Guidelines and working documents in 2011

CVMP Quality

Reference number	Document title	Status
EMA/CVMP/VICH/502/1999-Rev.1	VICH GL18(R) on residual solvents in new veterinary medicinal products, active substances and excipients	Adopted September 2011
EMA/CVMP/814/00-Rev.2	HMPC Guideline on quality of herbal medicinal products/traditional herbal medicinal products	Adopted September 2011
EMA/CVMP/815/00-Rev.2	Guideline on specifications: test procedures and acceptance criteria for herbal substances, herbal preparations and herbal medicinal products/traditional herbal medicinal products	Adopted September 2011
EMA/CVMP/VICH/858875/2011	VICH GL51 on Quality: Statistical evaluation of stability data	Adopted for consultation, December 2011 (End of consultation 12 June 2012)

CVMP Efficacy

Reference number	Document title	Status
EMA/CVMP/016/00-Rev.3	Guideline on the conduct of bioequivalence studies for veterinary medicinal products	Adopted April 2011
EMA/CVMP/760764/2010	Concept paper on the revision of the CVMP Guideline for the demonstration of efficacy for veterinary medicinal products containing antimicrobial substances	Adopted for consultation, April 2011 (End of consultation 31 July 2011)
EMA/CVMP/EWP/459868/2008	Guideline on demonstration of target animal safety and efficacy of veterinary medicinal products intended for use in farmed finfish	Adopted May 2011
EMA/CVMP/EWP/325284/2011	Questions and Answers document in relation to the CVMP Guideline on pharmaceutical fixed combination products (EMEA/CVMP/83804/05)	Adopted October 2011
EMA/CVMP/EWP/82829/2009-Rev.1	Question and Answer document in relation to the CVMP guideline on testing and evaluation of the efficacy of antiparasitic substances for the treatment and prevention of tick and flea infestations in dogs and cats.	Adopted November 2011

CVMP Safety

Reference number	Document title	Status
EMEA/CVMP/VICH/463072/2009	VCHI GL46: Metabolism study to determine the quantity and identify the nature of residues	Adopted March 2011
EMEA/CVMP/VICH/463104/2009	VCHI GL47: Laboratory animals comparative metabolism studies	Adopted March 2011
EMEA/CVMP/VICH/463199/2009	VCHI GL48: Marker residue depletion studies to establish product withdrawal periods	Adopted March 2011
EMEA/CVMP/VICH/463202/2009	VCHI GL49: Validation of analytical methods used in residue depletion studies	Adopted March 2011
EMA/CVMP/VICH/467/2003	VCHI GL36: General approach to establish a microbiological ADI	Adopted for consultation, March 2011 (End of consultation 14 September 2011)
EMA/CVMP/90250/2010	Draft Guideline on risk characterisation an assessment of maximum residue limits (MRLs) for	Adopted for consultation, September 2011 (End of consultation 30 June

Reference number	Document title	Status
	biocides.	2012)

CVMP Environmental Risk Assessment

Reference number	Document title	Status
EMA/CVMP/ERA/147844/2011	Reflection paper on the testing strategy and risk assessment for plants	Adopted December 2011
EMA/CVMP/ERA/430327/2009	Guideline on determining the fate of veterinary medicinal products in manure	Adopted March 2011
EMA/CVMP/ERAWP/409328/2010	Reflection paper on risk mitigation measures related to the environmental risk assessment of veterinary medicinal products	Adopted for consultation, May 2011 (End of consultation 31 August 2011)
EMA/CVMP/ERA/172074/2008-Rev.3	Questions and answers document on implementation of ERA Guideline in support of VICH guidelines (GL 6 and GL 38)	Adopted July 2011

CVMP Immunologicals

Reference number	Document title	Status
EMA/CVMP/IWP/206555/2010	Guideline on requirements for the production and control of immunological veterinary medicinal products	Adopted for consultation, March 2011 (End of consultation 30 September 2011)
EMA/CVMP/IWP/314550/2010	Guideline on the design of studies to evaluate the safety and efficacy of fish vaccines	Adopted November 2011
EMA/CVMP/IWP/785621/2011	Concept paper on the need of revision of the position paper on indications for veterinary vaccines	Adopted for consultation, October 2011 (End of consultation 15 January 2012)
EMA/CVMP/IWP/314550/2010	Guideline on the design of studies to evaluate the safety and efficacy of fish vaccines	Adopted November 2011
EMA/CVMP/IWP/594618/2010	Guideline on the requirements for combined vaccines and associations of immunological veterinary medicinal products (IVMPs)	Adopted for consultation, November 2011 (End of consultation 30 April 2012)
EMA/CVMP/VICH/463/2002	VICH GL34 on Biologicals: Mycoplasma - Test for the detection	Adopted for consultation, December 2011

Reference number	Document title	Status
	of Mycoplasma contamination	(End of consultation 12 March 2012)
EMA/CVMP/VICH/582610/2009	VICH GL50 on Biologicals: Testing harmonisation of criteria to waive target animal batch safety testing for inactivated vaccines for veterinary use	Adopted for consultation, December 2011 (End of consultation 12 June 2012)
EMA/675371/2011	EMA report on the implementation of the possibility for waiving the target animal batch safety test for immunological veterinary medicinal products in the European Union	Endorsed December 2011

CVMP Pharmacovigilance

Reference number	Document title	Status
EMA/CVMP/PhVWP/471721/2006	Recommendation on the basic surveillance of EudraVigilance Veterinary (EVVet) data	Adopted February 2011
EMA/CVMP/PhVWP/44873/2011	Public bulletin - Veterinary pharmacovigilance for 2010	Adopted February 2011
EMA/CVMP/10418/2009-Rev.3	CVMP combined VeDDRA list of clinical terms for reporting suspected adverse reactions in animals and humans to veterinary medicinal products	Adopted June 2011
EMA/CVMP/PhVWP/377827/2011	List of species and breeds for electronic reporting of suspected adverse reactions in veterinary pharmacovigilance	Adopted June 2011
EMA/CVMP/PhVWP/288284/2007-Rev.4	Guidance notes on the use of VeDDRA terminology for reporting suspected adverse reactions in animals and humans	Adopted June 2011
SOP/V/4019	Standard operating procedure - Annual review of standard lists to be used in EudraVigilance Veterinary	Adopted June 2011
SOP/V/4032	Standard operating procedure - Safety monitoring of centrally authorised products	Adopted October 2011

CVMP Scientific Advisory Group on Antimicrobials

Reference number	Document title	Status
EMA/CVMP/SAGAM/736964/2009	Reflection paper on meticillin-resistant *Staphylococcus pseudintermedius* (MRSP)	Adopted January 2011
EMA/CVMP/287420/2010	CVMP Strategy on antimicrobials 2011-2015	Adopted July 2011
EMA/CVMP/SAGAM/435644/2011	Concept paper on Use of pleuromutilins in food-producing animals in the European Union: development of resistance and impact on human and animal health	Adopted for consultation, October 2011 (End of consultation 31 January 2012)
EMA/CVMP/SAGAM/741087/2009	Reflection paper on the use of macrolides, lincosamides and streptogramins (MLS) in food-producing animals in the European Union: development of resistance and impact on human and animal health	Adopted for consultation, October 2011

General

Reference number	Document title	Status
EMA/347137/2010	Summary of procedures for consultation by CVMP of Scientific Advisory Groups (SAGs) and ad-hoc expert groups functioning as SAGs in relation to applications for authorisation for veterinary medicinal products	Adopted February 2011
EMA/CVMP/287420/2010	CVMP Strategy on antimicrobials 2011-2015	Adopted July 2011
EMA/CVMP/414812/2011	Question and answer document on the CVMP guideline on the SPC for antimicrobial products	Adopted July 2011