



23 March 2011
EMA/CVMP/37837/2011
Veterinary Medicines and Product Data Management

Committee for Medicinal Products for Veterinary Use

Monthly report of application procedures, guidelines and related documents

March 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.

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	2	103
Advice given	65	8	18	5	96

Marketing authorisations					
	95-08	2009	2010	2011	Total
Granted	88	12	9	8	117
Withdrawals	2	0	4	0	6
Not renewed	2	0	0	0	2

Initial evaluation					
	95-08	2009	2010	2011	Total
Full (Submitted)	110	14	16	2	142
Abridged/ generics (Submitted)	10	1	2	0	13
Withdrawals	12	0	1	0	13
Positive opinions	91	13	14	6	124
Negative opinions	1	0	0	0	1

Extensions					
	95-08	2009	2010	2011	Total
Submitted	60	12	3	1	76
Withdrawals	2	1	1	0	4
Positive opinions	40	7	8	1	56
Negative opinions	0	0	0	0	0



Variations – applications submitted					
	95-08	2009	2010	2011	Total
Type IA	339	32	76	18	579
Type IB		41	63	10	
Type II	210	40	26	4	280
Transfers	11	3	8	2	24

Renewals					
	95-08	2009	2010	2011	Total
Submitted	50	18	7	4	79
Positive opinions	48	17	8	1	74
Negative opinions	0	0	0	0	0

Arbitrations and Community referrals					
	95-08	2009	2010	2011	Total
Referrals submitted	38	9	12	2	61
Opinions reached ¹	20	15 (5)	11 (1)	0	46 (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	1	1
Agreed	6	6
Scientific advice recommended	0	0

MUMS/ Limited market classification		
	2011	Total
Positive with financial incentives	0	0
Positive without financial incentives	6	6
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	2	60
Negative opinions ³	7	0	0	0	7

Extensions / modifications/extrapolations of MRLs					
	95-08	2009	2010	2011	Total
Submitted	98	2	10	1	111
Withdrawals	4	0	0	0	4
Positive opinions ³	113	3	3	1	120
Negative opinions	6	0	0	0	6
Extrapolations	50	0	0	0	50

² Including opinions recommending 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	Marketing authorisation holder	Therapeutic area	EMA/CVMP	European Commission
<ul style="list-style-type: none"> Invented name INN 		<ul style="list-style-type: none"> Target species Summary of indication 	<ul style="list-style-type: none"> Validation Opinion Active time Clock stop 	<ul style="list-style-type: none"> Opinion received Date of decision Notification Official Journal
<ul style="list-style-type: none"> CaniLeish 	<ul style="list-style-type: none"> Virbac S.A. 	<ul style="list-style-type: none"> Dogs Vaccine against Leishmania infection 	<ul style="list-style-type: none"> 17/03/2010 12/01/2011 210 91 	<ul style="list-style-type: none"> 13/01/2011 14/03/2011
<ul style="list-style-type: none"> ZULVAC 1 + 8 Ovis 	<ul style="list-style-type: none"> Pfizer Limited 	<ul style="list-style-type: none"> Sheep Vaccine for prevention of viraemia caused by Bluetongue Virus serotypes 1 and 8 	<ul style="list-style-type: none"> 18/03/2010 12/01/2011 180 119 	<ul style="list-style-type: none"> 13/01/2011 14/03/2011
<ul style="list-style-type: none"> BLUEVAC BTv8 	<ul style="list-style-type: none"> CZ Veterinaria S.A 	<ul style="list-style-type: none"> Cattle, sheep Vaccine for active immunisation against bluetongue disease 	<ul style="list-style-type: none"> 17/01/2009 09/02/2011 180 511 	<ul style="list-style-type: none"> 10/02/2011
<ul style="list-style-type: none"> Procox Emodepside and toltrazuril 	<ul style="list-style-type: none"> Bayer Animal Health GmbH 	<ul style="list-style-type: none"> Dogs Treatment of dogs when mixed parasitic infections, caused by certain specific roundworms and coccidia are suspected or demonstrated 	<ul style="list-style-type: none"> 16/02/2010 09/02/2011 210 148 	<ul style="list-style-type: none"> 11/02/2011
<ul style="list-style-type: none"> Veraflox Pradofloxacin 	<ul style="list-style-type: none"> Bayer Animal Health GmbH 	<ul style="list-style-type: none"> Dogs, cats Treatment for dogs and cats with particular infections caused by certain specific and susceptible pathogens 	<ul style="list-style-type: none"> 19/05/2009 14/07/2010 205 217 09/02/2011 (re-consideration) 	<ul style="list-style-type: none"> 11/02/2011

Product <ul style="list-style-type: none"> Invented name INN 	<ul style="list-style-type: none"> Marketing authorisation holder 	Therapeutic area <ul style="list-style-type: none"> Target species Summary of indication 	EMA/CVMP <ul style="list-style-type: none"> Validation Opinion Active time Clock stop 	European Commission <ul style="list-style-type: none"> Opinion received Date of decision Notification Official Journal
<ul style="list-style-type: none"> Zuprevo Tildipirosin 	<ul style="list-style-type: none"> Intervet International BV 	<ul style="list-style-type: none"> Pigs, cattle Treatment of bacterial infections in the respiratory tract in pigs and cattle 	<ul style="list-style-type: none"> 16/02/2010 08/03/2011 210 177 	<ul style="list-style-type: none"> 10/03/2011
<ul style="list-style-type: none"> CERTIFECT Fipronil, (S)-methoprene, amitraz 	<ul style="list-style-type: none"> MERIAL SAS 	<ul style="list-style-type: none"> Dogs Treatment and prevention of infestations with ticks, alone or in association with fleas and/or chewing lice 	<ul style="list-style-type: none"> 16/03/2010 09/03/2011 210 148 	<ul style="list-style-type: none"> 10/03/2011

CVMP opinions in 2011 on establishment of MRLs for new substances

Positive opinions

<ul style="list-style-type: none"> Substance INN 	<ul style="list-style-type: none"> Therapeutic area Target species 	EMA/CVMP <ul style="list-style-type: none"> Validation Opinion Active time Clock stop 	European Commission <ul style="list-style-type: none"> Opinion received Date of regulation Official Journal
<ul style="list-style-type: none"> Methylpredni – solone (after provisional MRLs)	<ul style="list-style-type: none"> Bovine 	<ul style="list-style-type: none"> n/a 12/01/2011 90 0 	<ul style="list-style-type: none"> 27/01/2011
<ul style="list-style-type: none"> Octenidine dihydrochloride 	<ul style="list-style-type: none"> All mammalian food producing species 	<ul style="list-style-type: none"> 11/08/2009 08/02/2011 210 246 	<ul style="list-style-type: none"> 21/02/2011
<ul style="list-style-type: none"> Monepantel (after provisional MRLs)	<ul style="list-style-type: none"> Caprine 	<ul style="list-style-type: none"> n/a 09/03/2011 90 0 	<ul style="list-style-type: none">

Arbitrations and Community referrals in 2011

Type of referral	<ul style="list-style-type: none"> Date of clock start CVMP opinion 	<ul style="list-style-type: none"> Product name INN
Referral under Art. 34 of Directive 2001/82/EC	<ul style="list-style-type: none"> 11/11/2009 	<ul style="list-style-type: none"> Fortekor vet and associated names Benazepril hydrochloride
Referral under Art. 34 of Directive 2001/82/EC	<ul style="list-style-type: none"> 14/04/2010 	<ul style="list-style-type: none"> Synulox Lactating Cow and associated names Amoxicillin, clavulanic acid, prednisolone
Referral under Art. 33(4) of Directive 2001/82/EC	<ul style="list-style-type: none"> 14/07/2010 	<ul style="list-style-type: none"> Combimox Lactating Cow Amoxicillin, clavulanic acid, prednisolone
Referral under Art. 33(4) of Directive 2001/82/EC	<ul style="list-style-type: none"> 14/07/2010 	<ul style="list-style-type: none"> Nisamox Lactating Cow Amoxicillin, clavulanic acid, prednisolone
Referral under Art. 33(4) of Directive 2001/82/EC	<ul style="list-style-type: none"> 14/07/2010 	<ul style="list-style-type: none"> Combisyn Lactating Cow Amoxicillin, clavulanic acid, prednisolone
Referral under Art. 34 of Directive 2001/82/EC	<ul style="list-style-type: none"> 14/07/2010 	<ul style="list-style-type: none"> Doxycycline 50% WSP and associated names Doxycycline hyclate
Referral under Art. 34 of Directive 2001/82/EC	<ul style="list-style-type: none"> 14/07/2010 	<ul style="list-style-type: none"> Doxyfar 50% WSP and associated names Doxycycline hyclate

Type of referral	<ul style="list-style-type: none"> • Date of clock start • CVMP opinion 	<ul style="list-style-type: none"> • Product name • INN
Referral under Art. 34 of Directive 2001/82/EC	<ul style="list-style-type: none"> • 09/11/2010 	<ul style="list-style-type: none"> • Baytril 10% oral solution and associated names • Enrofloxacin
Referral under Art. 33(4) of Directive 2001/82/EC	<ul style="list-style-type: none"> • 09/02/2011 	<ul style="list-style-type: none"> • Clavudale 50 mg tablet for cats and dogs • Amoxicillin and clavulanic acid
Referral under Art. 35 of Directive 2001/82/EC	<ul style="list-style-type: none"> • 09/03/2011 	<ul style="list-style-type: none"> • Veterinary medicinal products containing active substances belonging to the class of flukicides for which no MRL has been established in milk

Guidelines and working documents in 2011

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 for consultation, March 2011 (End of consultation 30 June 2011)
EMA/CVMP/ERA/430327/2009	Guideline on determining the fate of veterinary medicinal products in manure	Adopted March 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 for consultation, March 2011 (End of consultation 30 September 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

CVMP Scientific Advisory Group on Antimicrobials

Reference number	Document title	Status
EMA/CVMP/SAGAM/736964/2009	Reflection paper on meticillin-resistant <i>Staphylococcus pseudintermedius</i> (MRSP)	Adopted January 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