European Medicines Agency
Veterinary Medicines and Inspections

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COMMITTEE FOR MEDICINAL PRODUCTS FOR VETERINARY USE

Monthly Report of Application Procedures, Guidelines and Related Documents

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

	95-06	2007	2008	2009	Total
Submitted	51	7	5	1	64

Initial Evaluation

	95-06	2007	2008	2009	Total
Full[1]	83	14	13	3	113
Abridged/Generics	6	1	3	0	10
Withdrawals	11	0	1	0	12
Positive Opinions	69	9	13	5	96
Negative Opinions	1	0	0	0	1

Marketing Authorisations

	95-06	2007	2008	2009	Total
Granted	66	9	13	5	93
Withdrawals	1	0	1	0	2
Not renewed	1	1	0	0	2

Extensions - Annex II Applications[2]

	95-06	2007	2008	2009	Total
Submitted	47	9	4	5	65
Withdrawals	1	0	0	1	2
Positive Opinions	32	1	7	1	41
Negative Opinions	0	0	0	0	0

Variations – Applications submitted

	95-06	2007	2008	2009	Total
Type IA	238	29	23	7	358
Type IB		24	25	8	
Type II	111	47	52	13	219
Transfers	7	2	2	0	11

[1] Initial applications submitted and validated: 123 applications in total (full + abridged), comprising 64 immunologicals and 59 pharmaceuticals. Negative opinions: in case of appeals, the opinion will not be counted twice.

[2] Extensions applications submitted and validated: 58 line extensions in total, comprising 11 immunologicals and 47 pharmaceuticals; one opinion can cover a number of extensions

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Renewals					
	95-06	**2007**	**2008**	**2009**	**Total**
Submitted	29	14	7	6	56
Positive Opinions	29	11	8	3	51
Negative Opinions	0	0	0	0	0

Establishment of MRLs for new substances					
	95-06	**2007**	**2008**	**2009**	**Total**
Submitted	63	2	1	1	67
Withdrawals	5	0	0	0	5
Positive Opinions[3]	49	3	2	1	55
Negative Opinions[4]	6	0	1	0	7

Arbitrations and Community Referrals					
	95-06	**2007**	**2008**	**2009**	**Total**
Referrals Submitted	21	6	11	3	41
Opinions Reached	4	10	6	3	23

Extensions / Modifications/Extrapolations of MRLs					
	95-06	**2007**	**2008**	**2009**	**Total**
Submitted	95	1	2	2	100
Withdrawals	4	0	0	0	4
Positive Opinions[3]	107	4	2	2	115
Negative Opinions[4]	6	0	0	0	6
Extrapolations	45	0	5	0	50

[3] Including opinions recommending definitive MRLs for substances with previously provisional maximum residue limits
[4] Including one opinion concluding that final MRL could not be established for a substance with provisional maximum residue limits previously established

CVMP Opinions in 2009 on Medicinal Products for Veterinary Use

Positive Opinions

Product ▪ Brand name ▪ INN	**Marketing authorisation holder**	**Therapeutic area** ▪ Target species ▪ Summary of indication	**EMEA/CVMP** ▪ Validation ▪ Opinion ▪ Active time ▪ Clock stop	**European Commission** ▪ Opinion received ▪ Date of decision ▪ Notification ▪ Official Journal
• Netvax	Schering-Plough, UK	• Chickens • Necrotic enteritis	• 10/02/2007 • 11/02/2009 • 210 • 379	
• BTVPUR Alsap 8 • Inactivated adjuvanted vaccine	Mérial, France	• Sheep, cattle • Prevention of Blue Tongue virus serotype 8	• 25/03/2008 • 11/02/2009 • 175 • 149	
• Improvac • GnRF analogue	• Pfizer, UK	• Male pigs • Control of boar taint	• 14/08/2007 • 11/03/2009 • 210 • 365	•
• Leucofeligen FeLV/RCP	• Virbac, France	• Cats • Immunisation against against feline calicivirosis, viral rhinotracheitis, panleucopenia ad leukaemia	• 18/03/2008 • 11/03/2009 • 210 • 147	•

• Leucogen	• Virbac, France	• Cats • Immunisation against feline leukaemia	• 18/03/2008 • 11/03/2008 • 210 • 147	•

Negative Opinions

Product ▪ Brand name ▪ INN	**Marketing authorisation holder**	**Therapeutic area** ▪ Target species ▪ Summary of indication	**EMEA/CVMP** ▪ Validation ▪ Opinion ▪ Active time ▪ Clock stop	**European Commission** ▪ Opinion received ▪ Date of decision ▪ Notification ▪ Official Journal

Withdrawals prior to opinion

Product ▪ Brand name ▪ INN	**Marketing authorisation holder**	**Therapeutic area** ▪ Target species ▪ Summary of indication	**EMEA/CVMP** ▪ Validation ▪ Opinion ▪ Active time ▪ Clock stop	**European Commission** ▪ Opinion received ▪ Date of decision ▪ Notification ▪ Official Journal

CVMP Opinions in 2009 on establishment of MRLs for new substances

Positive Opinions

Substance INN	**Therapeutic area** ▪ Target species	**EMEA/CVMP** ▪ Validation ▪ Opinion ▪ Active time ▪ Clock stop	**European Commission** ▪ Opinion received ▪ Date of regulation ▪ Official Journal

Negative Opinions (Recommendation for inclusion in Annex IV or inability to recommend inclusion in any of the Annexes to Regulation 2377/90)

Substance INN	**Therapeutic area** ▪ Target species	**EMEA/CVMP** ▪ Validation ▪ Opinion ▪ Active time ▪ Clock stop	**European Commission** ▪ Opinion received ▪ Date of regulation ▪ Official Journal

Arbitrations and Community Referrals in 2009

Type of referral	**Date of clock start / CVMP opinion**	▪ **Product name** ▪ **INN**
Referral for arbitration – Art. 33(4) of Directive 2001/82/EC	13/05/2008 15/01/2009	▪ ENRO-K 10% oral solution ▪ Enrofloxacin

Referral for arbitration – Art. 33(4) of Directive 2001/82/EC	13/05/2008 15/01/2009	▪ Unisol (avifox) 10% oral solution ▪ Enrofloxacin
Referral for arbitration – Art. 33(4) of Directive 2001/82/EC	11/03/2009 (clock start)	▪ Pharmasin 100% w/w water soluble granules ▪ Tylosine tartrate
Referral under Art. 35 of Directive 2001/82/EC	16/01/2008 12/02/2009)	▪ Injectable veterinary medicinal products containing ivermectin indicated for use in cattle ▪ Ivermectin
Referral under Art. 35 of Directive 2001/82/EC	11/02/2009 (clock start)	▪ All strengths of water soluble powders and oral solutions containing doxycycline hyclate ▪ Doxycycline hyclate
Referral under Art. 35 of Directive 2001/82/EC	16/04/2009 (clock start)	▪ Veterinary medicinal formulations containing colistin at 2 MIU/ml and intended for administration in drinking water to any food producing species ▪ Colistin sulfate

Urgent procedures

Type of procedure	CVMP opinion	Product name

Guidelines and Working Documents in 2009

CVMP Efficacy

Reference number	Document title	Status
EMEA/CVMP/016/00-Rev.1-CONSULTATION	Guideline on the conduct of bioequivalence studies for veterinary medicinal products	Adopted for consultation, March 2009 (End of consultation: September 2009)
EMEA/CVMP/EWP/82829/2009	Question and Answer document in relation to 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, March 2009
EMEA/CVMP/28510/2008	Guideline on dossier requirements for anticancer medicinal products for dogs and cats	Adopted, April 2009

CVMP Environmental Risk Assessment (ERA)

Reference number	Document title	Status
EMEA/CVMP/ERA/10043/2009-CONSULTATION	Concept paper on the fate of veterinary medicinal products in manure	Adopted, April 2009

CVMP Immunologicals

Reference number	Document title	Status
EMEA/CVMP/IWP/105506/2007-CONSULTATION	Guideline on data requirements for multi-strain dossiers for inactivated vaccines against avian influenza, bluetongue and foot-and-mouth disease	Adopted for consultation, March 2009 (End of consultation: September 2009)
EMEA/CVMP/IWP/439467/2007-CONSULTATION	Reflection paper on the demonstration of a possible impact of maternally derived antibodies on vaccine efficacy in young animals	Adopted for consultation, March 2009 (End of consultation: September 2009)
EMEA/CVMP/IWP/250147/2008-CONSULTATION	Guideline on data requirements to support in-use stability claims for veterinary vaccines	Adopted for consultation, March 2009 (End of consultation: September 2009)
EMEA/CVMP/IWP/123243/2006-Rev.1-CONSULTATION	Guideline on data requirements for immunological veterinary medicinal products intended for Minor Use or Minor Species/ Limited markets	Adopted for consultation, March 2009 (End of consultation: June 2009)

CVMP Pharmacovigilance

Reference number	Document title	Status
SOP-EMEA/599270/2007	SOP on Handling of pharmacovigilance Rapid Alerts (RAs) and Non Urgent Informaion (NUI)for veterinary use	Endorsed, January 2009
EMEA/CVMP/10418/2009	Combined VeDDRA list of clinical terms for reporting suspected adverse reactions in animals and humans to veterinary medicinal products	Adopted, February 2009
SOP/V/4023-Rev.1	Management of Period Safety Update Reports (PSURs) for Centrally Authorised Products (CAPs) and Annex I – Contact details of national competent authorities for PSUR submission	Adopted, April 2009
EMEA/CVMP/PhVWP/133883/2004-Rev.2	Mandate, Objectives and Rules of Procedure For The CVMP Pharmacovigilance Working Party (PhVWP-V)	Adopted, April 2009
EMEA/INS/PhV/85061/2008	Procedure for Reporting of Pharmacovigilance Inspections Requested by the CVMP	Adopted, April 2009

Joint CHMP/CVMP Quality

Reference number	Document title	Status
EMEA/CVMP/QWP/544461/2007	Guideline on the quality aspects of single-dose veterinary spot-on products	Adopted, January 2009

EMEA/CHMP/CVMP/QWP/663093/2008	Question and Answer document on Plastic Immediate Packaging Materials	Adopted, January 2009
EMEA/CHMP/CVMP/QWP/17760/2009-Rev.1-CONSULTATION	Revised Guideline on the use of near infrared spectroscopy by the pharmaceutical industry and the data requirements for new submissions and variations	Adopted for consultation, February 2009 (End of consultation: August 2009)
EMEA/555991/2007	New Question and Answers which aim to clarify several issues associated with the use of Process Analytical Technology (PAT),	Adopted, February 2009
EMEA/CHMP/CVMP/QWP/160263/2009	Question and Answer documents on endotoxin/sterility testing during and at the end of shelf-life	Adopted, April 2009
EMEA/CHMP/CVMP/QWP/450653/2006	Recommendation on the Assessment of the quality of medicinal products containing existing/ known active substances	Adopted, April 2009

CVMP Safety

Reference number	Document title	Status
EMEA/CVMP/SWP/322484/2008-Rev.1-CONSULTATION	Guideline on user safety for pharmaceutical veterinary medicinal products	Adopted for consultation, April 2009 (End of consultation, August 2009)
EMEA/CVMP/VICH/486/02-Rev.2	VICH Guideline on Studies to Evaluate the Safety of Residues of Veterinary Drugs in Human Food: General Approach to Testing	

CVMP Scientific Advisory Group on Antimicrobials

Reference number	Document title	Status
EMEA/CVMP/SAGAM/81730/2006	Revised Reflection Paper on the use of 3rd and 4th generation cephalosporins in food producing animals in the European Union: development of resistance and impact on human and animal health, including recomendations	Adopted, March 2009
EMEA/CVMP/SAGAM/68290/2009	Reflection paper on MRSA in food producing and companion animals in the European Union: epidemiology and control options for human and animal health	Adopted, March 2009

CVMP General

Reference number	Document title	Status
EMEA/INS/GCP/390778/2008	Procedure for the preparation of a risk-based programme for routine PhV Inspections of MAHs connected with Veterinary Centrally Authorised Products (CAPs)	Adopted, January 2009
EMEA/INS/GCP/85059/2008	Procedure for coordination of pharmacovigilance inspections requests by the CVMP	Adopted, January 2009
EMEA/INS/S&T/75010/2009	Sampling and Testing of Centrally Authorised products	Adopted, April 2009
EMEA/CVMP/248499/2007-Rev.1	Recommendation on the evaluation of the benefit-risk balance of veterinary medicinal products	Adopted, April 2009