28 November 2013
EMA/456228/2013
Veterinary Medicines Division

List of acronyms and abbreviations used in CVMP agenda and minutes

3Rs	Replacement, Reduction and Refinement
ADI	Acceptable Daily Intake
AHEG	Ad Hoc Expert Group
Alt	Alternate
AMEG	Antimicrobial Advice ad hoc Expert Group
AR	Assessment Report
ARfD	Acute Reference Dose
Art.	Article
ASMF	Active Substance Master File
AWP	Antimicrobials Working Party
BMD	Bone Mineral Density
CAP	Centrally Authorised Product
CCRVDF	Codex Committee on Residues of Veterinary Drugs in Foods
CHMP	Committee for Medicinal Products for Human Use
CKD	Chronic Kidney Disease
CMD(h)	Co-ordination Group for Mutual Recognition and Decentralised Procedures (Human)
CMDv	Co-ordination Group for Mutual Recognition and Decentralised Procedures (Veterinary)
CMS	Concerned Member State
Co.	Company
Co-Rapp	Co-rapporteur

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CTA	Clinical Trial Application
CV	Curriculum Vitae
CVMP	Committee for Medicinal Products for Veterinary Use
CXMP	CVMP or CHMP
DCP	Decentralised Procedure
DDPS	Detailed Description of Pharmacovigilance System
DPRA	Drug and Pharmacies Regulation Act (Canada)
EEA	European Economic Area
EC	European Commission
ECDC	European Centre for Disease Prevention and Control
eCTD	Electronic Common Technical Document
EDQM	European Department for the Quality of Medicines
EFSA	European Food Safety Authority
EGGVP	European Group for Generic Veterinary Products
EMA	European Medicines Agency
EMEA	Old acronym for European Medicines Agency (replaced by EMA)
END	National expert
EPAA	The European Partnership for Alternative Approaches to Animal Testing
EPAR	European Public Assessment Report
EPMAR	European Public MRL Assessment Report
ERA	Environmental Risk Assessment
ERAWP	Environmental Risk Assessment Working Party
ESVAC	The European Surveillance of Veterinary Antimicrobial Consumption
EU	European Union
EU-RL	European Union Reference Laboratory
EURL ECVAM	European Union Reference Laboratory for Alternatives to Animal Testing
EVVet	EudraVigilance Veterinary
EWG	Expert Working Group
EWP	Efficacy Working Party

FBS	Fetal Bovine Serum
FDA	Food and Drug Administration (USA)
FVE	Federation of Veterinarians of Europe
GCP	Good Clinical Practice
GL	Guideline
GMO	Genetically Modified Organism
GMP	Good Manufacturing Practice
HMA	Heads of Medicines Agencies
ICH	International Conference on Harmonisation of Technical Requirements for Registration of Pharmaceuticals for Human Use
IFAH	International Federation for Animal Health
INN	International Non-proprietary Name
IVMP	Immunological Veterinary Medicinal Product
IWP	Immunologicals Working Party
JECFA	Joint FAO/WHO Expert Committee on Food Additives
JEG-3Rs	Joint CVMP/CHMP Ad-hoc Expert Group on the Application of the 3Rs in Regulatory Testing of Medicinal Products
LoOI	List of Outstanding Issues
LoQ	List of Questions
MA	Marketing Authorisation
MAH	Marketing Authorisation Holder
MB	Management Board
MMD	Managing Meeting Documents system
MRL	Maximum Residue Limit
MR	Mutual Recognition
MRP	Mutual Recognition Procedure
MS	Member State
MUMS	Minor Use and Minor Species
N/a	Not applicable
NA	National Authority
NCA	National Competent Authority
OE	Oral Explanation

OIE	Office International des Epizooties
OMCL	Official Medicines Control Laboratories
PARERE	EURL ECVAM's Network for Preliminary Assessment of Regulatory Relevance
PBPK modeling	Physiologically Based Pharmacokinetic modeling
PBT	Persistent Bioaccumulative Toxic (chemical)
PET	Polyethylene Terephthalate
Ph. Eur.	European Pharmacopoeia
PhV	Pharmacovigilance
PhVWP	Pharmacovigilance Working Party
PI	Product Information
PIQ	Product Information Quality
PL	Package Leaflet
PM	Project Manager
PSUR	Periodic Safety Update Report
Q&A	Questions and Answers
QRD	Quality Review of Documents
QWP	Joint CHMP/CVMP Quality Working Party
Rapp	Rapporteur
RAR	Rapporteur Assessment Report
RBA	Risk Based Approach
Reg.	Regulation
Rev.	Revision
RMS	Reference Member State
SA	Scientific Advice
SAG	Scientific Advisory Group
SAGAM	Scientific Advisory Group on Antimicrobials
SAR	Serious Adverse drug Reaction
SAWP	Scientific Advice Working Party
SME	Small and Medium-sized Enterprises
SPC	Summary of Product Characteristics
SPG	Strategic Planning Group

SWP	Safety Working Party
Tbc	To be confirmed
Tbd	To be discussed/decided
TIGes	Telematic Implementation Group for electronic submissions
TOPRA	The Organisation for Professionals in Regulatory Affairs
V	Veterinary
VeDDRA	Veterinary Dictionary for Drug Related Affairs
VICH	International Cooperation on Harmonisation of Technical Requirements for Registration of Veterinary Medicinal Products
VMP	Veterinary Medicinal Product
WP	Working Party

Country codes of EU/EEA countries

ISO country code	Country (short name in English)
AT	Austria
BE	Belgium
BG	Bulgaria
CY	Cyprus
CZ	Czech Republic
DE	Germany
DK	Denmark
EE	Estonia
ES	Spain
FI	Finland
FR	France
GR	Greece
HR	Croatia
HU	Hungary
IE	Ireland
IS	Iceland
IT	Italy
LI	Liechtenstein

ISO country code	Country (short name in English)
LT	Lithuania
LU	Luxembourg
LV	Latvia
MT	Malta
NL	Netherlands
NO	Norway
PL	Poland
PT	Portugal
RO	Romania
SE	Sweden
SI	Slovenia
SK	Slovakia
UK	United Kingdom