European Medicines Agency
Veterinary Medicines and Inspections

December 2007

ANNEX I

MRL PROCEDURE

SUBMISSION OF DOSSIERS TO THE MEMBERS AND ALTERNATES OF THE COMMITTEE FOR MEDICINAL PRODUCTS FOR VETERINARY USE

When the application is made one full copy of the dossier (including expert reports), one extra copy of each expert report and one electronic copy of the routine analytical method are to be forwarded to the EMEA for the validation.

The Rapporteur and the Co-rapporteur should both receive one full copy of the dossier. Applicants should confirm with them their requirements with regard to the submission in paper copy versus electronic version.

Other members and alternates of the CVMP should receive the information according to the requirements specified in the consolidated list of dossier submission requirements, published on the EMEA website (http://www.emea.europa.eu/pdfs/general/contacts/46610207en.pdf).

The contact details of all CVMP members and alternates[1] are also available on the EMEA website (http://www.emea.europa.eu/htms/general/contacts/CVMP/CVMP_members.html).

[1] Please note that, depending on the Internet browser you are using, it may be possible to right-click anywhere in the area of the contact details and export the whole list into Excel, using the "Export to Microsoft Excel" option.

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