10 April 2014
EMA/CVMP/176966/2014
Committee for Medicinal Products for Veterinary Use

Opinion of the Committee for Medicinal Products for Veterinary Use on the establishment of maximum residue limits

Procedure no: EMEA/V/MRL/003071/MODF/0002

Name of the substance: Barium selenate (INN)

Basis for the opinion

Pursuant to Article 11 of Regulation (EC) No 470/2009 of 6 May 2009, Germany submitted to the European Medicines Agency, on 2 December 2013, a request to review the maximum residue limits established for barium selenate in bovine and ovine species.

Recommendation

The Committee, having considered the request from Germany, recommends, by consensus, the amendment of the entry for barium selenate in Table 1 of the Annex to Commission Regulation (EU) No 37/2010, in accordance with the following table:

Pharmaco-logically active substance	Marker residue	Animal species	MRLs	Target tissues	Other provisions	Therapeutic classification
Barium selenate	NOT APPLICABLE	All food producing species	No MRL required	NOT APPLICABLE	Not for administration by injection	Alimentary tract and metabolism/ mineral supplements

The Icelandic CVMP member agrees with the above-mentioned recommendation of the Committee.

The scientific conclusions of the Committee are presented in the European public MRL assessment report (EPMAR), provided in Annex I of this opinion.

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The present opinion is forwarded to the European Commission and to the applicant together with its appendices.

London, 10 April 2014

Signature on file

Dr. A. Holm
Chair, on behalf of the CVMP

Annex I

European public MRL assessment report (EPMAR)