4 December 2014
EMA/CVMP/666932/2014
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/003225/MODF/0002
EMEA/V/MRL/002867/MODF/0002
EMEA/V/MRL/002880/MODF/0003

Name of the substance: Potassium selenate (INN)
Sodium selenate (INN)
Sodium selenite (INN)

Basis for the opinion

Pursuant to Article 11 of Regulation (EC) No 470/2009 of 6 May 2009, European Commission submitted to the European Medicines Agency, on 12 May 2014, a request to review the maximum residue limits established for potassium selenate, sodium selenate and sodium selenite in all food producing species.

Recommendation

The Committee, having considered the request from the European Commission, recommends by consensus the maintenance of the existing entries for the above substances in Table 1 of the Annex to Commission Regulation (EU) No 37/2010.

The Icelandic and Norwegian CVMP members agree with the above-mentioned recommendation of the Committee.

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

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

London, 4 December 2014

Signature on file

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

Annex I

European public MRL assessment report (EPMAR)