II
(Non-legislative acts)

REGULATIONS

COMMISSION IMPLEMENTING REGULATION (EU) No 84/2012
of 1 February 2012
amending the Annex to Regulation (EU) No 37/2010 on pharmacologically active substances and their classification regarding maximum residue limits in foodstuffs of animal origin, as regards the substance phenoxymethylpenicillin
(Text with EEA relevance)

THE EUROPEAN COMMISSION,

Having regard to the Treaty on the Functioning of the European Union,


Having regard to the opinion of the European Medicines Agency formulated by the Committee for Medicinal Products for Veterinary Use,

Whereas:

(1) The maximum residue limit for pharmacologically active substances intended for use in the Union in veterinary medicinal products for food-producing animals or in biocidal products used in animal husbandry should be established in accordance with Regulation (EC) No 470/2009.


(3) Phenoxymethylpenicillin is currently included in Table 1 of the Annex to Regulation (EU) No 37/2010 as an allowed substance, for porcine species, applicable to muscle, liver and kidney and for poultry species, applicable to muscle, skin and fat, liver and kidney, excluding animals from which eggs are produced for human consumption.

(4) An application for the extension of the existing entry for phenoxymethylpenicillin to include eggs for poultry species has been submitted to the European Medicines Agency.

(5) The Committee for Medicinal Products for Veterinary Use has recommended establishing a maximum residue limit (hereinafter ‘MRL’) for phenoxymethylpenicillin for porcine species, applicable to muscle, liver and kidney and for poultry species, applicable to muscle, skin and fat, liver, kidney and eggs.

(6) The entry for phenoxymethylpenicillin in Table 1 of the Annex to Regulation (EU) No 37/2010 should therefore be amended to include the MRL for eggs for poultry species.

(7) It is appropriate to provide for a reasonable period of time for the stakeholders concerned to take measures that may be required to comply with the newly set MRL.

(8) The measures provided for in this Regulation are in accordance with the opinion of the Standing Committee on Veterinary Medicinal Products,

HAS ADOPTED THIS REGULATION:

Article 1
The Annex to Regulation (EU) No 37/2010 is amended as set out in the Annex to this Regulation.

Article 2

This Regulation shall enter into force on the third day following its publication in the Official Journal of the European Union.

It shall apply from 2 April 2012.

This Regulation shall be binding in its entirety and directly applicable in all Member States.

Done at Brussels, 1 February 2012.

For the Commission
The President
José Manuel BARROSO
The entry for phenoxymethylpenicillin in Table 1 of the Annex to Regulation (EU) No 37/2010 is replaced by the following:

<table>
<thead>
<tr>
<th>Pharmacologically active substance</th>
<th>Marker residue</th>
<th>Animal species</th>
<th>MRL</th>
<th>Target tissues</th>
<th>Other provisions (according to Article 14(7) of Regulation (EC) No 470/2009)</th>
<th>Therapeutic classification</th>
</tr>
</thead>
<tbody>
<tr>
<td>Phenoxymethylpenicillin</td>
<td>Phenoxymethylpenicillin</td>
<td>Porcine</td>
<td>25 μg/kg</td>
<td>Muscle, Liver, Kidney</td>
<td>NO ENTRY</td>
<td>Anti-infectious agents/ Antibiotics'</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Poultry</td>
<td>25 μg/kg</td>
<td>Muscle, Skin and fat, Liver, Kidney, Eggs</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

EN 2.2.2012 Official Journal of the European Union L 30/3