COMMISSION IMPLEMENTING REGULATION (EU) No 436/2012 of 23 May 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 azamethiphos

(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 (hereinafter ‘MRL’) 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) Azamethiphos is currently included in Table 1 of the Annex to Regulation (EU) No 37/2010 as an allowed substance, for salmonidae species.

(4) An application for the extension of the existing entry for azamethiphos applicable to fin fish species has been submitted to the European Medicines Agency.

(5) The Committee for Medicinal Products for Veterinary Use recommended the extension of that entry and the absence of the need to establish an MRL for azamethiphos in fin fish species.

(6) The entry for azamethiphos in Table 1 of the Annex to Regulation (EU) No 37/2010 should therefore be amended accordingly.

(7) 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 that of its publication in the Official Journal of the European Union.

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

Done at Brussels, 23 May 2012.

For the Commission
The President
José Manuel BARROSO

## ANNEX

The entry corresponding to azamethiphos 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>‘Azamethiphos’</td>
<td>NOT APPLICABLE</td>
<td>Fin fish</td>
<td>No MRL required</td>
<td>NOT APPLICABLE</td>
<td>NO ENTRY</td>
<td>Antiparasitic agents/Agents against ectoparasites</td>
</tr>
</tbody>
</table>