Substances considered as not falling within the scope of Regulation (EC) No. 470/2009\(^1\), with regard to residues of veterinary medicinal products in foodstuffs of animal origin

1. **Background information**


Article 1(a) of Regulation (EC) No. 470/2009 defines its scope as follows:

“For the purposes of ensuring food safety, this Regulation lays down rules and procedures in order to establish:

a) the maximum concentration of a residue of a pharmacologically active substance which may be permitted in food of animal origin (maximum residue limit);

(...)

Article 1(3) of the Regulation states that:

“This Regulation shall not apply to ‘active principles of biological origin intended to produce active or passive immunity or to diagnose a state of immunity, used in immunological veterinary medicinal products’.”

In Article 2(a) it is specified that “residues of pharmacologically active substances” means all pharmacologically active substances, expressed in mg/kg or μg/kg on a fresh weight basis, whether active substances, excipients or degradation products, and their metabolites which remain in food obtained from animals.

Previously, in the context of the evaluation of MRL applications under Council Regulation (EC) No. 2377/90, the CVMP discussed the concept of "pharmacologically active substances", in particular considering the need for MRL evaluations for excipients and manufacturing materials. As detailed in the CVMP publication entitled *Position Paper on the definition of substances capable of pharmacological action in the context of Directive 2001/82/EC, as amended, with a particular reference to excipients*.

and manufacturing materials (EMEA/CVMP/072/97-Rev.1²), it was concluded that "substances capable of pharmacological action are substances pharmacodynamically active at the dose at which they are administered to the target animal by means of the veterinary medicinal product in which they are included". It follows that if an excipient has not pharmacodynamically activity at the relevant dose, then no MRL evaluation would be needed.

Subsequently, the Committee adopted a paper entitled Reflection Paper on consideration of adjuvants and preservatives under Council Regulation (EEC) No. 2377/90 laying down a community procedure for the establishment of maximum residue limits of veterinary medicinal products in foodstuffs of animal origin (EMEA/CVMP/339116/2007-CONSULTATION), in which it is concluded that the approach used to determine whether an MRL evaluation is required for preservatives and adjuvants should be the same as is used in relation to excipients (i.e. it should be based on the presence or absence of pharmacodynamic activity at the intended dose).

Since the implementation of Council Regulation (EEC) No. 2377/90, the CVMP has deliberated on many substances (including excipients, adjuvants and preservatives) to be used in veterinary medicinal products intended for food producing species, and regularly receives requests (either scientific advice or ad hoc requests) to consider whether such substances fall within the scope of the MRL regulation. The substances for which the CVMP has concluded that no MRL evaluation is required are listed in the CVMP publication "Substances considered as not falling within the scope of Regulation (EEC) No. 2377/90" (EMEA/CVMP/046-00), also often referred to as the 'out of scope list'.

The list also includes a small number of compounds that do not fall within the categories of excipients, adjuvants or preservatives but are natural substances essential for life or are biologically active constituents. Due to the nature of these specific compounds the CVMP considered that an evaluation for the establishment of maximum residue limits would not be appropriate.

Following the implementation of Regulation (EC) No. 470/2009 there was a need to update the background information and legal references included in the document containing the 'out of scope list'. This document performs that function and supersedes the CVMP publication Substances considered as not falling within the scope of Council Regulation (EEC) No. 2377/90. The list presented on the following pages includes all the substances included in the superseded document.

It should be noted that this list of substances is in no way exhaustive and includes only substances for which requests in this respect were made to CVMP by a company or a national authority.

Any enquiries may be posted to mrl@ema.europa.eu

² Initially adopted in April 1997 and further revised in July 2004
## II. Substances considered as not falling within the scope of Regulation (EC) No 470/2009

### Excipients

<table>
<thead>
<tr>
<th>Substance</th>
<th>CAS No.</th>
<th>Use Limitations</th>
</tr>
</thead>
<tbody>
<tr>
<td>Acetone</td>
<td>67-64-1</td>
<td>for cutaneous use only</td>
</tr>
<tr>
<td>Aqua purificata</td>
<td>7732-18-5</td>
<td></td>
</tr>
<tr>
<td>2-(2-n-Butoxyethoxy)ethanol</td>
<td>112-34-5</td>
<td></td>
</tr>
<tr>
<td>Carbomer</td>
<td>9003-01-4</td>
<td></td>
</tr>
<tr>
<td>Carbomer copolymer A</td>
<td>1456857-02-1</td>
<td>for topical administration at doses of up to 1.6 mg/kg bw</td>
</tr>
<tr>
<td>Casein hydrolysate</td>
<td>65072-00-6</td>
<td></td>
</tr>
<tr>
<td>Cetearyl ethylhexanoate</td>
<td>59130-70-7</td>
<td>for cutaneous use only at doses of up to 35.0 mg/kg bw</td>
</tr>
<tr>
<td>Chlorobutanol</td>
<td>57-15-8</td>
<td>at concentrations up to 1%</td>
</tr>
<tr>
<td>Coconut oil</td>
<td>8001-31-8</td>
<td></td>
</tr>
<tr>
<td>Collagen hydrolysate</td>
<td>9007-34-5</td>
<td></td>
</tr>
<tr>
<td>Copolymer of polyvinylpyrrolidone and vinyl acetate</td>
<td>25086-89-9</td>
<td>for cutaneous use only</td>
</tr>
<tr>
<td>Corn oil</td>
<td>8001-30-7</td>
<td></td>
</tr>
<tr>
<td>Cotton seed oil</td>
<td>8001-29-4</td>
<td></td>
</tr>
<tr>
<td>Denatonium benzoate</td>
<td>3734-33-6</td>
<td>for topical administration at doses of up to 0.25 mg/kg bw</td>
</tr>
<tr>
<td>Diethylaminoethyl (DEAE)-Dextran</td>
<td>9013-34-7</td>
<td>at concentrations up to 150 mg/ml</td>
</tr>
<tr>
<td>Diethanolamine</td>
<td>111-42-2</td>
<td>at doses up to 0.3 mg/kg bw/day</td>
</tr>
<tr>
<td>Dimethylidocadecylammonium bromide</td>
<td>3700-67-2</td>
<td>for use as an adjuvant at a total dose of not greater than 21 mg/animal.</td>
</tr>
<tr>
<td>Dimethyl ether</td>
<td>115-10-6</td>
<td>for use as a propellant for topical administration</td>
</tr>
<tr>
<td>Ethoxyquin</td>
<td>91-53-2</td>
<td>at concentrations up to 0.1 mg/g</td>
</tr>
<tr>
<td>Fatty acid methyl esters</td>
<td>67762-38-3</td>
<td>for topical administration</td>
</tr>
<tr>
<td>Fibrous materials of plant origin</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Gamma hexalactone</td>
<td>685-06-7</td>
<td>for topical administration at doses of up to 14 mg/kg bw</td>
</tr>
<tr>
<td>Gelatin</td>
<td>9000-70-8</td>
<td></td>
</tr>
<tr>
<td>Glycerol dimethylketal</td>
<td>100-79-8</td>
<td>at concentrations up to 150 mg/ml</td>
</tr>
<tr>
<td>4-(2-Hydroxyethyl)-1-piperazineethanesulfonic acid (HEPES)</td>
<td>7365-45-9</td>
<td>for use as a buffering agent in vaccines and vaccine diluents</td>
</tr>
<tr>
<td>Isopropyl myristate</td>
<td>110-27-0</td>
<td>at doses up to 5 mg/kg bw</td>
</tr>
<tr>
<td>Macrogol cetostearyl ether</td>
<td>68439-49-6</td>
<td>for administration by the intramammary route at doses of up to 0.95 mg/kg bw</td>
</tr>
<tr>
<td>Maleic acid</td>
<td>110-16-7</td>
<td>for use in buffering systems at doses up to 0.39 mg/kg bw</td>
</tr>
</tbody>
</table>
Meglumine (CAS No: 6284-40-8): at doses up to 1.5 mg/kg bw
Metacresol (CAS No: 108-39-4): at concentrations up to 0.2%
2-Octyl-dodecanol (CAS No: 5333-42-6): when administered topically at doses up to 20 mg/kg bw
Oleic acid (CAS No: 112-80-1)
Olive oil (CAS No: 8001-25-0)
Peanut oil (CAS No: 8002-03-7)
Polybutene (CAS No: 9003-29-6)
Polyethylene glycol-75 lanolin (CAS no 8039-09-6 and 61790-81-8): for topical use only
Polysaccharides naturally occurring such as celluloses and hydroxycelluloses, dextrans and glucans
Polymyxin B (CAS No: 1404-26-8): for use as an endotoxin neutralising agent in vaccines at doses of not more than 500 μg/dose (approximately 5000 IU per dose) or not more than 8 μg/kg bw (approximately 80 IU/kg bw), whichever is the lower
Polyoxyethylene oleate (CAS No. 9004-96-0): at doses up to 1.15 mg/kg bw
Polyoxyethylene oleic alcohol (CAS No 9004-98-2): at doses up to 0.95 mg/kg bw
Polyoxyethylene (40) sorbitol septaoleate (CAS No: 63089-85-0): for cutaneous use only
Polyoxypropylene (PPG-2) myristyl ether propionate (CAS No: 74775-06-7): for cutaneous use only
Propolis
Sesame oil (CAS No: 8008-74-0)
Silicones (CAS No: 9006-65-9)
Simethicone (CAS No: 8050-81-5)
Sodium starch glycolate (CAS: 9063-38-1): for cutaneous use only
Soybean (milled and hulled)
Soybean oil, including epoxidized soybean oil (CAS No: 8001-22-7 and 8013-07-8)
Squalane (CAS No: 111-01-3): as a component of the adjuvant system
Starches normally found in food and food grade starches
Sulfolipo-cyclodextrin
Triethanolamine (CAS No: 102-71-6)
Trometamol (CAS No: 77-86-1): for use in buffering systems at doses up to 0.65 mg/kg bw
Vermiculite (CAS No: 1318-00-9): including expanded vermiculite
Zymosan A (CAS No: 58856-93-2): for use as an adjuvant at doses of up to 0.12 mg/kg bw

**Normal foodstuffs**

Avena (oats)

Carbohydrates naturally occurring
Cereals
Chocolate flavour
*Coffea arabica*
Honey
Lipids as constituents of the human diet
Royal jelly
*Petroselinum crispum* (parsley)
Peptides and proteins as constituents of the human diet
Pulses

**Chemically complex substances of natural origin**

Organ autolysates
Immunoglobulines
Dried dialysate derived from blood
Lyophilised ruminal fluid

**Natural substances essential for animal and human life**

Oxygen (CAS No: 7782-44-7)

**Biologically active constituents**

Probiotic components including bacteria and yeasts
Stem cells