

# EN

## ANNEX

Common Name	IUPAC Name Identification Numbers	Minimum degree of purity of the active substance <sup>1</sup>	Date of approval	Expiry date of approval	Product type	Specific conditions <sup>2</sup>
<i>Bacillus thuringiensis</i> subsp. <i>israelensis</i> serotype H14, strain SA3A	Not applicable	No relevant impurities	1 July 2016	30 June 2026	18	<p>The product assessment shall pay particular attention to the exposures, the risks and the efficacy linked to any uses covered by an application for authorisation, but not addressed in the Union level risk assessment of the active substance.</p> <p>For biocidal products, authorisations are subject to the following conditions:</p> <p>(1) For professional users, safe operational procedures and appropriate organisational measures shall be established. Where exposure cannot be reduced to an acceptable level by other means, products shall be used with appropriate personal protective equipment.</p> <p>(2) For products that may lead to residues in food or feed, the need to set new or to amend existing maximum residue levels (MRLs) in accordance</p>

<sup>1</sup> The purity indicated in this column was the minimum degree of purity of the active substance used for the evaluation made in accordance with Article 8 of Regulation (EU) No 528/2012. The active substance in the product placed on the market can be of equal or different purity if it has been proven technically equivalent with the evaluated active substance.

<sup>2</sup> For the implementation of the common principles of Annex VI to Regulation (EU) No 528/2012, the content and conclusions of assessment reports are available on the Commission website: [http://ec.europa.eu/environment/chemicals/biocides/index\\_en.htm](http://ec.europa.eu/environment/chemicals/biocides/index_en.htm).

Common Name	IUPAC Name Identification Numbers	Minimum degree of purity of the active substance <sup>1</sup>	Date of approval	Expiry date of approval	Product type	Specific conditions <sup>2</sup>
						with Regulation (EC) No 470/2009 of the European Parliament and of the Council <sup>3</sup> or Regulation (EC) No 396/2005 of the European Parliament and of the Council <sup>4</sup> shall be verified, and any appropriate risk mitigation measures shall be taken to ensure that the applicable MRLs are not exceeded.

<sup>3</sup> Regulation (EC) No 470/2009 of the European Parliament and of the Council of 6 May 2009 laying down Community procedures for the establishment of residue limits of pharmacologically active substances in foodstuffs of animal origin, repealing Council Regulation (EEC) No 2377/90 and amending Directive 2001/82/EC of the European Parliament and of the Council and Regulation (EC) No 726/2004 of the European Parliament and of the Council (OJ L 152, 16.6.2009, p. 11).

<sup>4</sup> Regulation (EC) No 396/2005 of the European Parliament and of the Council of 23 February 2005 on maximum residue levels of pesticides in or on food and feed of plant and animal origin and amending Council Directive 91/414/EEC (OJ L 70, 16.3.2005, p. 1).