



Strasbourg, **XXX**
SANTE/10268/2022 ANNEX
(POOL/E4/2022/10268/10268-EN
ANNEX.docx)
[...] (2022) **XXX** draft

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to the

Commission Implementing Regulation (EU) .../...

approving *Chrysanthemum cinerariaefolium* extract from open and mature flowers of *Tanacetum cinerariifolium* obtained with supercritical carbon dioxide for product type 19 in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council

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Common Name	IUPAC Name Identification Numbers	Minimum degree of purity of the active substance ¹	Date of approval	Expiry date of approval	Product type	Specific conditions
<i>Chrysanthemum cinerariaefolium</i> extract from supercritical carbon dioxide	<i>Chrysanthemum cinerariaefolium</i> extract from open and mature flowers of <i>Tanacetum cinerariifolium</i> obtained with supercritical carbon dioxide EC No: 289-699-3 CAS No: 89997-63-7	100% w/w of <i>Chrysanthemum cinerariaefolium</i> extract from open and mature flowers of <i>Tanacetum cinerariifolium</i> obtained with supercritical carbon dioxide	1 February 2024	31 January 2034	19	The authorisation of biocidal products is subject to the following conditions: (1) 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; (2) The product assessment shall pay particular attention to the exposure of and potential risks for non-professional users and the general public; (3) 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 with Regulation (EC) No 470/2009 of the European Parliament and of the Council ² or Regulation (EC) No 396/2005 of the European Parliament and of the Council ³ shall be verified, and any appropriate risk mitigation measures shall be taken to ensure that such MRLs are not exceeded.

¹ The purity indicated in this column was the minimum degree of purity of the active substance evaluated. The active substance in the product placed on the market can be of equal or different purity if it has been proven to be technically equivalent to the evaluated active substance.

² 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).

³ 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).