

Trinol A/S
Søagervej 9,
9500 Hobro,
Denmark

Oslo, 11.11.2022

Your ref.:

Our ref.:
2018/8283

Contact person:
Marianne Stave Sekkenes

Authorisation for Trinol A/S – TRINOL VEPSESKUM NO-2022-0235

With reference to the application dated 28.06.2018 for the biocidal product TRINOL VEPSESKUM R4BP 3 case number BC-DX040720-27 as a same biocidal product to the reference product SPECIAL ONE – NO-2021-0212.

Decision

The Norwegian Environment Agency hereby grants an authorisation for the biocidal product TRINOL VEPSESKUM on the Norwegian market. The authorisation is granted from 11.11.2022 to 29.07.2031 with the authorisation number NO-2022-0235 (to be stated on the label).

Terms and conditions for the authorisation

The product is authorised as a same biocidal product in Norway under the terms and conditions as described in the final Norwegian Summary Product Characteristic (SPC) attached to the R4BP3 asset case no. NO-0029928-0000. The final SPC can also be found on the website of the European Chemicals Agency here: [Information on biocides - ECHA \(europa.eu\)](https://echa.europa.eu).

The information on the label, and, if relevant, in the Material Safety Data Sheet and Technical Data Sheet, shall be in Norwegian and in accordance with the terms and conditions provided in the final Norwegian SPC. This is the responsibility of the authorisation holder. Further requirements are described in Article 69, 70 and 72 of Regulation (EU) No 528/2012 (the Biocidal Products Regulation, BPR). An electronic copy of the label(s) for each authorised product shall be submitted to the Norwegian Environment Agency by email (biocides@miljodir.no) within three months from the authorisation date. Please mark the email with the authorisation number.

All biocidal products on the Norwegian market shall be registered in the Norwegian Product Register in accordance with the Norwegian Biocide Regulation of 18 April 2017 No 480 § 2-2, by using the biocide notification form. In addition, biocidal products which are classified as hazardous according to Regulation (EC) No 1272/2008 (the CLP Regulation) shall be fully declared if they are sold in amounts of 100 kg or more per year. The forms to be used and further

information can be found on our website <https://www.environmentagency.no/areas-of-activity/product-register/>

This authorisation is given in accordance with Article 5 of Regulation (EU) No 414/2013 (the Same Biocidal Products Regulation, amended by Regulation (EU) 2016/1802).

Background

The BPR is implemented in Norwegian law through the Norwegian Biocide Regulation of 18 April 2017 No 480.

The procedures for applications for same biocidal product are set out in Regulation (EU) No 414/2013 (amended by Regulation (EU) 2016/1802) which supplements the BPR. These applications shall be authorised under the same terms and conditions as the reference product granted by the Norwegian Environment Agency, in line with Article 7(1) of the same regulation.

The conditions for granting an authorisation of a biocidal product are laid down in Article 19 of the BPR. Additionally, the transitional measures given in Article 91 apply.

In general, a biocidal product is authorised for a period not exceeding 10 years in line with Article 17(4) of the BPR. However, for authorisations that are mutual recognised, it is an agreement among the Member States that these authorisations shall have the same expiry date as the national authorisation granted by the reference Member State (c.f. CA-Sept14-Doc.5.7 –Final).

The application concerns

Trinol A/S has applied for an authorisation for the biocidal product TRINOL VEPSESKUM on the Norwegian market as a same biocidal product as the reference product SPECIAL ONE. The biocidal product contains the active substances cyfluthrin, transfluthrin and piperonyl butoxide/PBO and is for use in product type 18 – insecticides, acaricides and products to control other arthropods. The authorisation holder in Norway is Trinol A/S.

Evaluation by the Norwegian Environment Agency

The decision is based on the evaluation of the reference Member State of the reference product, SPECIAL ONE.

Relevant information

Phase out period for existing biocidal products on the Norwegian market

In cases where the authorised biocidal product has been made available on the Norwegian market under the national transitional measurements (c.f. Article 89 of the BPR), the existing stocks must be phased out in line with Article 89(4) of the BPR. The product shall not be made available on the market with effect from 180 days after the date of this letter. Furthermore, the use of existing stocks of the biocidal product may continue for up to 365 days after the date of this letter. During this period, all advertising material related to products that do not comply with the new conditions, should also be removed from the market.

Unexpected or adverse effects

If the authorisation holder becomes aware of any unexpected or adverse effects concerning the authorised biocidal product(s) or the active substance it contains, the authorisation holder is obligated to notify without delay to the Norwegian Environment Agency (c.f., Article 47 of the BPR).

Changes to the authorisation

If it is desirable to make any changes to the product authorisation, the authorisation holder must submit an application/notification for change to the Norwegian Environment Agency, in accordance with Article 50 of the BPR independently of the reference product. This procedure is described in detail in Regulation (EU) No. 354/2013 on changes of biocidal products. The fees to be charged for applications for change are given in appendix 1A of the Norwegian Biocide Regulation.

Authorisations of a same product or of a related reference product may be changed or cancelled independently of each other.

Annual fee

For authorised biocidal products on the Norwegian market, an annual fee will be charged. Please see appendix 1B of the Norwegian Biocide Regulation for details. We kindly ask you to inform us by email (biocides@miljodir.no) if you do not intend to place the product on the Norwegian market, and therefore should not be charged with the annual fee.

Renewal of application

An application for a renewal of the national authorisation must be submitted 550 days before the authorisation period expires, at the latest, according to Article 31(1) of the BPR.

Right to appeal

This decision may be appealed to the Ministry of Climate and Environment.

An appeal shall be submitted to the Norwegian Environment Agency within three weeks after receipt of this letter.

Best regards
Norwegian Environment Agency

This document has been signed electronically

Trine-Lise Torgersen
Head of Section

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