

Bayer A/S
Arne Jacobsen Alle 13,
2300 Copenhagen S,
Denmark

Oslo, 16.11.2021

Your ref.:

Our ref.:
2021/7736

Contact person:
Marianne Stave Sekkenes

Authorisation – Racumin Expert – NO-2021-0216

We refer to your application for mutual recognition in sequence of the biocidal product Racumin Expert, R4BP 3 case number BC-LG068388-27, containing the active substance coumatetralyl. The Norwegian Environment Agency hereby grants authorisation.

Background

Regulation (EU) No. 528/2012 concerning the making available on the market and use of biocidal products (the Biocidal Products Regulation, BPR), is implemented in Norwegian law through the Norwegian Biocide Regulation of 18 April 2017 No. 480. 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.

According to Article 17(4) of the BPR, an authorisation can be granted for a maximum of 10 years. To facilitate the renewal procedure, it is agreed (CA-Sept14-Doc.5.7 -Final) that authorisations granted by the concerned Member States should have the same expiry date as the authorisation which is granted by the reference Member State.

Evaluation

Coumatetralyl is considered a candidate of exclusion since it meets the criteria for being classified as Repr. 1B. Coumatetralyl, however, satisfies the conditions laid down in Article 5(2)(c), meaning that not authorising products containing this active substance would have disproportionate negative consequences for society in comparison with the risk associated with the use. This implies that coumatetralyl satisfies the criteria given in Article 10(1)(a) and must be viewed as a substance eligible for substitution. Under Article 23(1) of the BPR, Member States evaluating biocidal products containing an active substance that is a candidate for substitution in accordance with Article 10(1), are required to perform a comparative assessment. A comparative assessment in accordance with Article 23 of the BPR has been carried out, and as a result, the criteria of Article 23(3) of BPR are found not to be met. The product can therefore be authorised for a period not exceeding 5 years.

Derogations from mutual recognition

According to national restrictions in Norway, there is only one professional user category for rodenticides: trained professionals (and others with special authorisation). In addition, only tamper resistant bait stations are allowed for indoor and outdoor use around buildings. A derogation from mutual recognition is made for Racumin Expert in accordance with Article 37(1) (b) of the BPR, adjusting the terms and conditions of the Norwegian authorisation by restricting the user categories and area of use in accordance with the national policy.

Another derogation from the mutual recognition is made in accordance with Article 37(1) (e) of the BPR, adjusting the terms and conditions of the Norwegian authorisation by replacing common vole (*Microtus arvalis*) with field vole (*Microtus agrestis*) as a target organism. Common vole is not present in Norway and, hence, is replaced with field vole which is a similar organism present in Norway. The derogation has been communicated to the applicant and agreed upon in the evaluation process.

Decision

Subject to Articles 19 and 91 of the BPR, cf. § 1 of the Norwegian Biocide Regulation, the Norwegian Environment Agency grants an authorisation of Racumin Expert until 26.02.2024

The product is mutual recognised in Norway under the terms and conditions as described in the Summary Product Characteristic (SPC). The decision is based on the evaluation of the reference Member State Germany, some adjustments according to the national restrictions concerning the user categories and area of use, cf. Article 37 of the BPR.

The authorisation concerns

Product name:	Racumin Expert
Trade name(s):	Racumin Expert
Active substance(s):	Coumatetralyl (CAS no. 5836-29-3)
Product type:	PT 14 – Rodenticides
Authorisation holder in Norway:	Bayer A/S
Authorisation number:	NO-2021-0216
Authorisation date:	16.11.2021
Expiry date:	26.02.2024

Additionally, the conditions provided in the Norwegian Summary of Product Characteristics (SPC) apply. The SPC is uploaded to R4BP 3.

The Norwegian Environment Agency may, in accordance with Article 47 of the BPR, cancel or amend the authorisation should new information on the product or the active substance come to our attention that may affect the authorisation. Should the authorisation holder be aware of such information, the Norwegian Environment Agency should be notified without delay.

According to Article 31(1) of the Biocidal Products Regulation, an application for a renewal of the authorisation must be submitted 550 days before the authorisation period expires, at the latest.

Label

The information on the label, and, if relevant, in the Material Safety Data Sheet and Technical Data Sheet, shall be in accordance with the conditions provided in the attached SPC. Furthermore, Article 69(1), (2) and Article 70 of the BPR also apply.

The authorisation holder is responsible for ensuring that the information given in the above-mentioned documents is accurate, and is translated to Norwegian, cf. Article 69(3) of the BPR.

An electronic copy of the label with the Norwegian authorisation number NO-2021-0216 shall be submitted to the Norwegian Environment Agency within three months from the authorisation date, using the email address biocides@miljodir.no. In case of a biocidal product family, please submit an electronic copy of the label for one individual product in each meta SPC.

Phase-out period of existing stocks, when relevant

In line with Article 89(4), existing products that do not comply with the conditions of this authorisation, 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.

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. 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.

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 using the e-mail address 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.

Registration in the Norwegian Product Register

All biocidal products on the Norwegian market must be registered in the Product Register by using the biocide notification form. In addition, biocidal products which are classified as hazardous must be fully declared, using the declaration form, if they are sold in amounts of 100

kg or more per year. Forms and further information can be found on our website <https://www.environmentagency.no/areas-of-activity/product-register/>.

Appeal

This decision can be appealed to the Ministry of Climate and Environment, in accordance with Article 28 of the Public Administration Act. The complaint must be submitted to the Norwegian Environment Agency within 3 weeks after receipt of this letter, in accordance with Article 29 of the Public Administration Act.

Best regards
Norwegian Environment Agency

This document has been signed electronically

Trine-Lise Torgersen
Head of Section

Marianne Stave Sekkenes
Senior Adviser