LATVIJAS VIDES, ĢEOLOĢIJAS UN METEOROLOĢIJAS CENTRS

Riga

02.02.2021 Nr. 4-6/171

Unichem d.o.o.

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On authorisation of the biocidal product Neopermin through mutual recognition in Latvia

Latvian Environment, Geology and Meteorology Centre (LEGMC) has evaluated an application submitted by **Unichem d.o.o.** on 15th April 2016 concerning an authorisation of the biocidal product **Neopermin** through mutual recognition in Latvia.

LEGMC has agreed with Product Assessment Report and Summary of Product Characteristics for *Neopermin* developed by the reference Member State – The Netherlands.

Therefore, in accordance with Article 34 of *Regulation (EU) No 528/2012 of the European Parliament and of the Council of 22 May 2012 concerning the making available on the market and use of biocidal products* (regulation 528/2012), LEGMC authorises the biocidal product *Neopermin.*

Neopermin contains **0.5%** of **permethrine** (CAS No. 52645-53-12, EC No. 258-067-9).

Additional trade names:

- TARIN;
- Pulveris skudru iznīcināšanai.

LEGMC assigns the authorisation number **LV/2021/MR/001**.

The authorisation number is valid until **5th February 2031.**

The authorisation number shall be indicated on the label of the biocidal product.

The authorisation is granted on the following terms:

- Product type: 18 – insecticides, acaricides and products to control other arthropods;
- Target organisms: crawling insects such as ants and cockroaches;
- Fields of use: indoor;
- Users: trained professionals, professionals and non-professionals;
- Product description: dustable powder;
- Pack sizes and packaging material: as indicated in Summary of Product Characteristics;
- Product stability: up to 24 months.

The authorisation applies only to the *Neopermin* in the composition, form and packing for which the first authorisation is granted by reference Member State.

The information on the label (and if applicable an enclosed instruction of use) of the *Neopermin* shall be as it is indicated in the first authorisation of above mentioned product, taking into account also the information which is stated in the Product Assessment Report and Summary of Product Characteristics issued by reference Member State.

The information on the label shall be in Latvian.

Notwithstanding content of the label specified above, requirements stated in:

- Article 69 Regulation (EU) No 528/2012 of the European Parliament and of the Council of 22 May 2012 concerning the making available on the market and use of biocidal products;
- Regulation (EC) No. 1272/2008 of the European Parliament and of the Council of 16 December 2008 on classification, labelling and packaging of the substances and mixtures, amending and repealing Directives 67/548/EEC and 1999/45/EC, and amending Regulation (EC) No 1907/2006;
- all other relevant legislation shall be applied.

Unichem d.o.o. shall inform LEGMC about any changes in accordance with *Commission Implementing Regulation (EU) No 354/2013 of 18th April 2013 on changes of biocidal products authorised in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council.*

If the first authorisation issued by reference Member State is amended or revoked, the authorisation of *Neopermin* may be re-opened for review before 5th February 2031.

Additionally, LEGMC would like to inform that Unichem d.o.o. is fully responsible of the content of the biocidal product *Neopermin,* as well as its classification, labelling, instruction of use and safety data sheet.

LEGMC would like to ask Unichem d.o.o. to notify the above mentioned information down to supply chain.

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LVGMC INFORMĀCIJAI

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