LATVIJAS VIDES, ĢEOLOĢIJAS UN METEOROLOĢIJAS CENTRS

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Nr. 4-6/1390

DeLaval NV

Industriepark-Drongen 10,
9031 Gent,
Belgium

On an authorisation of biocidal product DeLaval PeraDis through mutual recognition in Latvia

Latvian Environment, Geology and Meteorology Centre (LEGMC) has evaluated an application submitted by DeLaval NV on 5th September 2017 concerning an authorisation of the biocidal product **DeLaval PeraDis** through mutual recognition in parallel.

LEGMC has agreed with Product Assessment Report and Summary of Product Characteristics for the biocidal product **DeLaval PeraDis** developed by the reference Member State – 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 **DeLaval PeraDis** on the basis of mutual recognition process.

The authorisation holder for **DeLaval PeraDis** in Latvia is:

DeLaval NV.

DeLaval PeraDis contains **4.9 %** of peracetic acid (CAS No. 79-21-0; EC No. 201-186-8) as active substance.

LEGMC assigns the authorisation number LV/2022/MR/012 for the biocidal product DeLaval PeraDis.

The authorisation is valid until 4th November 2032.

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

The authorisation of biocidal product **DeLaval PeraDis** through mutual recognition is granted on the following terms:

- Product types: 3 – Veterinary hygiene and 4 - Food and feed area;
- Target organism: bacteria, yeasts and viruses;
- Users: professional;
- Product description: water soluble concentrate;
- Product stability: 1 year;
- Pack sizes and packaging materials: as indicated in Summary of Product Characteristics.

The authorisation through mutual recognition applies only to the biocidal product **DeLaval PeraDis** in the composition, form and packing material 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 biocidal product **DeLaval PeraDis** should be as it is indicated in the first authorisation of above mentioned biocidal 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 of *regulation 528/2012*,
- *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.

DeLaval NV as the authorisation holder 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 biocidal product **DeLaval PeraDis** through mutual recognition may be re-opened for review before the 04th November 2032.

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

LEGMC would like to ask DeLaval NV to notify the above mentioned information down to supply chain.

Head of Information Analysis Department A.Jantone

Signature not validated

Digitally signed by ANTA JANTONE
Date: 2022.10.31 17:13:41 EET

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