

Riga

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Detia Freyberg GmbH

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On an authorisation of AMEISEN MITTEL through mutual recognition in Latvia

Latvian Environment, Geology and Meteorology Centre (LEGMC) has evaluated an application submitted by **Detia Freyberg GmbH** on 20th April 2016 concerning an authorisation of **AMEISEN MITTEL** through mutual recognition in parallel.

LEGMC has agreed with Product Assessment Report and Summary of Product Characteristics for **AMEISEN MITTEL** developed by the reference Member State – Germany.

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 AMEISEN MITTEL on the basis of mutual recognition process.

The authorisation holder for AMEISEN MITTEL in Latvia is:

Detia Freyberg GmbH.

Additional trade names for AMEISEN MITTEL in Latvia are:

- Skudru-Ex:
- Skudru Granulas.

AMEISEN MITTEL contains **0.54** % (w/w) of permethrin (CAS No. 52645-53-1; EC No. 258-067-9) as active substance.

LEGMC assigns the authorisation number for biocidal product AMEISEN MITTEL through mutual recognition:

LV/2019/MR/005

The authorisation of AMEISEN MITTEL through mutual recognition is valid until $25^{\rm th}$ March 2029 based on reference Member State – Germany decision.

The authorisation number of **AMEISEN MITTEL** through mutual recognition shall be indicated on the label of the biocidal product.

The authorisation of AMEISEN MITTEL through mutual recognition is granted on the following terms:



- Product type: 18 Insecticides, acaricides and products to control other arthropods.
- Target organism black garden ant (Lasius niger);
- Users: professional and general public;
- Product description: granule and soluble granule;
- Product stability: shelf life 12 months;
- Use area: Around houses on paved ways, balconies and terraces.

The authorisation through mutual recognition applies only to the biocidal product AMEISEN MITTEL 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 AMEISEN MITTEL 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.

Detia Freyberg GmbH 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 AMEISEN MITTEL through mutual recognition may be re-opened for review before the 25th March 2029.

Additionally LEGMC would like to inform that Detia Freyberg GmbH is fully responsible of the content of the biocidal product AMEISEN MITTEL as well as its classification, labelling, instruction of use and safety data sheet.

LEGMC would like to ask Detia Freyberg GmbH to notify the above mentioned information down to supply chain.

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