LATVIJAS VIDES, ĢEOLOĢIJAS
UN METEOROLOĢIJAS CENTRS

Riga

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Bayer Sp. z o.o.

Al. Jerozolimskie 158,
02-326 Warsaw,
Poland

On an authorisation of the biocidal product Maxforce White IC

Latvian Environment, Geology and Meteorology Centre (LEGMC) has evaluated an application submitted by **Bayer Sp. z o.o.** on 20 June 2013 concerning an authorisation of **Maxforce White IC** through mutual recognition in parallel.

LEGMC has agreed with Product Assessment Report and Summary of Product Characteristics for **Maxforce White IC** developed by the reference Member States – United Kingdom.

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 (EU) No 528/2012), LEGMC authorises the **Maxforce White IC** on the basis of mutual recognition process.

The authorisation holder for **Maxforce White IC** in Latvia is:

Bayer Sp. z o.o.

Maxforce White IC contains **2,217%** of ***imidacloprid*** (CAS No.138261-41-3, EC No.428-040-8) as an active substance.

LEGMC assigns the authorisation number for biocidal product Maxforce White IC:

LV/2019/MR/007

The authorisation is valid until 29th January 2024.

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

The authorisation of **Maxforce White IC** through mutual recognition is granted on the following terms:

- Product type: 18 – Insecticides, acaricides and products to control other arthropods;
- Target organism: german cockroach– nymphs and adults – *Blattella germanica*; oriental cockroach – nymphs and adults – *Blatta orientalis*; common cockroach - nymphs and adults – *Blattidae*: American cockroach - nymphs and adults – *Periplaneta americana*; brown cockroach - nymphs and adults – *Supella longipalpa*;
- Users: professional;
- Product description: ready to use bait;
- Product stability: up to 2 years;
- Field of use - indoor

- Pack sizes and packaging material: plastic cartridge (PP cartridge with PE tip and plug), 20-30g.

The authorisation through mutual recognition applies only to the product **Maxforce White IC** 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 **Maxforce White IC** should 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*;
- *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.

Bayer Sp. z 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 **Maxforce White IC** through mutual recognition may be re-opened for review before 29th January 2024.

Additionally LEGMC would like to inform that Bayer Sp. z o.o. is fully responsible of the content of the biocidal product **Maxforce White IC** as well as its classification, labelling, instruction of use and safety data sheet.

LEGMC would like to ask Bayer Sp. z o.o. to notify the above mentioned information down to supply chain.

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