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**BROS Spółka z ograniczoną
odpowiedzialnością**

Karpia 24, 61-619, Poznan
Poland

On the authorisation of the biocidal product BROS Līdzeklis pret odiem bērniem through mutual recognition in Latvia

Latvian Environment, Geology and Meteorology Centre (LEGMC) has evaluated an application submitted by **BROS Spółka z ograniczoną odpowiedzialnością** on 18th April 2019 concerning an authorisation of **BROS Līdzeklis pret odiem bērniem** through mutual recognition in Latvia

LEGMC has agreed with Product Assessment Report and Summary of Product Characteristics developed by the reference Member States – Slovenia.

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 **BROS Līdzeklis pret odiem bērniem** on the basis of mutual recognition process.

BROS Līdzeklis pret odiem bērniem contains **17.3%** of **ethyl butylacetylaminopropionate (IR3535®)** (CAS No. 52304-36-6, EC No. 257-835-0) as active substance.

LEGMC assigns the authorisation number **LV/2022/MR/010**.

The authorisation is valid until **17th July 2032**.

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

The authorisation of is granted on the following terms:

- **Product type 19** – repellents and attractants;
- Users: general public (non-professional);
- Product description: ready-for-use repellent used on human skin;
- Target organisms: House mosquitoes, tropical mosquitoes and ticks;
- Application method: Manual spraying;
- Packaging: in accordance with the Summary of Product Characteristics;
- Product stability: 2 years.

The authorisation applies only to the product **BROS Līdzeklis pret odiem bērniem** 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 *BROS Līdzeklis pret odiem bērniem* 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 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.

BROS Spółka z ograniczoną odpowiedzialnością 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.*

Application on renewal of an authorisation shall be submitted according to *Commission Delegated Regulation (EU) No 492/2014 of 7 March 2014 supplementing Regulation (EU) No 528/2012 of the European Parliament and of the Council as regards the rules for the renewal of authorisations of biocidal products subject to mutual recognition.*

Additionally, LEGMC would like to inform that BROS Spółka z ograniczoną odpowiedzialnością is fully responsible of the content of the biocidal product as well as its classification, labelling, instruction of use and safety data sheet.

LEGMC would like to ask BROS Spółka z ograniczoną odpowiedzialnością to notify the above mentioned information down to supply chain.

Head of Information Analysis Department

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