

Riga

Czech Republic

On the authorisation of the biocidal product Bochemit Antiflash Family through mutual recognition in Latvia

Latvian Environment, Geology and Meteorology Centre (LEGMC) has evaluated an application submitted by **Bochemie a.s.** on 3th March 2022 concerning an authorisation of **Bochemit Antiflash Family** through mutual recognition in Latvia

LEGMC has agreed with Product Assessment Report and Summary of Product Characteristics for *Bochemit Antiflash Family* developed by the reference Member States – Czech Republic.

Therefore, in accordance with Article 33 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 Bochemit Antiflash Family on the basis of mutual recognition process.

Trade names of the products:

- *Bochemit Antiflash brown;*
- Bochemit Antiflash green;
- Bochemit Antiflash.

The Bochemit Antiflash Family contains **20.0%** of **Boric acid** (CAS No.10043-35-3, EC No.233-139-2) as active substance.

LEGMC assigns the authorisation number LV/2015/MR/003.

The authorisation is valid until 31 December 2022.

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

The authorisation of is granted on the following terms:

- Product type: 8 Wood preservative;
- Target organisms: wood rotting fungi, wood destroying insects,
- Users: industrial and professional;
- Product description: aqueous liquid concentrate (the ready to use solution is prepared by diluting the concentrate tenfold with water);
- Product stability: up to 3 years.



 Pack sizes and packaging material: as indicated in Summary of Product Characteristics.

The authorisation applies only to the product *Bochemit Antiflash Family* 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 *Bochemit Antiflash Family* 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.

Bochemie a. s. 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 Bochemie a. s. is fully responsible of the content of the biocidal product *Bochemit Antiflash Family* as well as its classification, labelling, instruction of use and safety data sheet.

LEGMC would like to ask Bochemie a. s. to notify the above mentioned information down to supply chain.

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