

Rīga

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**GLOBUZ**

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### **On authorisation of the biocidal product PROCULAC under the simplified authorisation procedure**

Latvian Environment, Geology and Meteorology Centre (LEGMC) on 21<sup>st</sup> September 2023 received application for a simplified authorisation of **PROCULAC** in accordance with Chapter V of the *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) 528/2012) submitted by **GLOBUZ** through Register for Biocidal Products (Case No. BC-BX088824-03).

LEGMC as Latvian competent authority has evaluated the above mentioned application and considers that biocidal product meets the conditions of the Article 25 of the Regulation (EU) No 528/2012:

- the active substance Lactic acid (CAS No. 50-21-5, EC No. 200-018-0) in the biocidal product appears in Annex I of the Regulation (EU) No 528/2012 and satisfy the restriction specified in that Annex;
- the biocidal product does not contain any substances of concern;
- the biocidal product does not contain nanomaterials;
- the biocidal product is effective;
- the handling of the biocidal product does not require personal protective equipment.

Therefore, LEGMC grants authorisation for *PROCULAC* with **Lactic acid** (CAS No. 50-21-5, EC No. 200-018-0) as active substance at the concentration **0.9% w/w** for **product type 2** and **4** as ready-to-use disinfectant for hard clean non-porous surfaces with bactericidal and yeasticidal efficacy in domestic, institutional and industrial area.

**LEGMC assigns an authorisation number EU-0031834-0000.** The authorisation number is valid until **15<sup>th</sup> January 2034.**

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

The authorisation applies only to the biocidal product *PROCULAC* in the composition claimed by the authorisation holder.

The information on the label (and if applicable an enclosed instruction of use) of the biocidal product *PROCULAC* should be in line with Summary of Product Characteristics.

Notwithstanding content of the label specified above, requirements stated in Article 69 of regulation 528/2012 and all other relevant legislation shall be applied.

GLOBUZ is fully responsible of the composition of the biocidal product, label, instruction of use and safety data sheet.

GLOBUZ 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 GLOBUZ wants to continue to make available on the market the biocidal product beyond the 15<sup>th</sup> January 2034, a new application for authorisation under the simplified authorisation procedure must be submitted.

The biocidal product authorised in accordance with Article 26 of the Regulation (EU) 528/2012 may be made available on the market in all Member States according to conditions laid down in Article 27 of the Regulation (EU) 528/2012.

Head of Information Analysis Department

signature\*

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