

Riga

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Lonza Cologne GmbH

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### On an authorisation of the biocidal product Vacsol Aqua 6118

Latvian Environment, Geology and Meteorology Centre (LEGMC) has evaluated an application submitted by Lonza Cologne GmbH on 22 February 2018 concerning an authorisation of Vacsol Aqua 6118 through mutual recognition in parallel.

LEGMC has agreed with Product Assessment Report and Summary of Product Characteristics for **Vacsol Aqua 6118** developed by the reference Member States – Sweden.

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 Vacsol Aqua 6118 on the basis of mutual recognition process.

The authorisation holder for Vacsol Aqua 6118 in Latvia is:

#### Lonza Cologne GmbH

The biocidal product Vacsol Aqua 6118 contains following active substances:

- penflufen (CAS No 494793-67-8), concentration 0,38%;
- permethrin (CAS No 52645-53-1, EC No 258-067-9), concentration 0,4%.

# LEGMC assigns the authorisation number for biocidal product Vacsol Aqua 6118:

#### LV/2020/MR/015

# The authorisation is valid until 2 November 2030.

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

The authorisation of Vacsol Aqua 6118 through mutual recognition is granted on the following terms:

- Product type: 8 Wood preservatives;
- Target organisms: House longhorn beetle larvae Hylotrupes bajulus L., Brown rot fungi hyphae Basidiomycetes;
- Users: industrial;
- Product description: soluble concentrate;
- Product stability: shelf life 24 months;
- Use area: indoor areas;
- Pack sizes and packaging material: as indicated in Summary of Product Characteristics.



The authorisation through mutual recognition applies only to the product **Vacsol Aqua 6118** in the composition, form and packing for which the authorisation is granted by reference Member State to **Vacsol Aqua 6118**.

The information on the label (and if applicable an enclosed instruction of use) of the **Vacsol Aqua 6118** should be as it is indicated in the 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.

Lonza Cologne GmbH shall inform LEGMC about any changes in accordance with Commission Implementing Regulation (EU) No 354/2013 of 18<sup>th</sup> 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 **Vacsol Aqua 6118** through mutual recognition may be re-opened for review before 2 November 2030.

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 Lonza Cologne GmbH is fully responsible of the content of the biocidal product **Vacsol Aqua 6118** as well as its classification, labelling, instruction of use and safety data sheet.

LEGMC would like to ask Lonza Cologne GmbH to notify the above mentioned information down to supply chain.

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