

23 rol March, 2018 No. 4-6/435 Riga

Reckitt Benckiser Healthcare (UK) Ltd.

> Dansom Lane Hull HU8 7DS United Kingdom

On major changes of the national authorisation of the HCl Family B

Latvian Environment, Geology and Meteorology Centre (LEGMC) received an application submitted by **Reckitt Benckiser Healthcare (UK) Ltd.** On the 9 December 2016 for major change via Register for Biocidal product (Case No. BC-KR028319-19) in accordance with Article 7 of Commission Implementing Regulation (EU) No 354/2013 of 18 April 2013 on changes of biocidal products authorised in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council for the biocidal product family **Hydrochloric Acid Family B** with authorisation number **LV/16/NA/02**.

LEGMC as the Competent Authority have developed Product Assessment Report and Summary of Product Characteristics for biocidal product family **Hydrochloric Acid Family B** based on application for major changes.

With this application for evaluation in accordance with Article 50 of *Regulation (EU) 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 all the conditions listed in authorisation letter No. 4-6/1087 issued by LEGMC on the 21st of June 2016 are fulfilled.

After evaluation of submitted changes it has been concluded that EthomeenT/12 and Arquad T50 does not pose a risk to environment and reclassification of the biocidal product family is not appropriate.

Updated Product Assessment Report and Summary of Product Characteristics are available in Register for Biocidal Products.

This letter is integral part of authorisation letter No. 4-6/1087 issued by LEGMC on the 21st of June 2016.

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