service public fédéral
SANTE PUBLIQUE,
SECURITE DE LA CHAINE ALIMENTAIRE
ET ENVIRONNEMENT

SPF Santé publique, Sécurité de la Chaîne alimentaire et Environnement

Direction générale Environnement

Maîtrise des risques

Eurostation II

NOS RÉF. Dossier n° xxxxx

DATE 17/01/2017

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OBJET: Prolongation of the existing national authorisation during the evaluation of the Union authorisation application

Dear Madam, Sir,

You have been granted a Belgian national authorization to place on the Belgian market the following biocidal products :

Ioklar Multi **notification number NOTIF461. (date first authorization : 28/07/2011)**

Veloucid Spray D **notification number NOTIF471. (date first authorization : 25/11/2011)**

In view of the pending Union Authorization application submitted to the European Chemical Agency on 23/07/2015 and according to the transitional measures as laid out in article 89, §2 of the Biocidal Product Regulation (EU) No 528/2012 we can provide you with a prolongation of making your products Ioklar Multi and Veloucid Spray D available on the Belgian market until 3 years after the date of approval of the last active substances to be approved in this given biocidal product. **Your product may remain on the Belgian market**, under the same conditions as those mentioned in the Belgian authorization NOTIF461 and NOTIF471, and at the latest until **1/09/2018**.

This letter is the **official notification of the new date of validity** of the mentioned Belgian authorizations NOTIF461 and NOTIF471. The biocidal products concerned will continue to be included in the list of the authorized biocidal products, which can be found at www.biocide.be. The already granted authorizations will not be amended. You can use this letter as proof of the validity of the existing authorizations.

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This letter remains **valid until the Union Authorization** is granted and will terminate on 1/09/2018 at the latest. The granted Union Authorization will provide a period of grace to continue to market the biocidal products concerned with the current authorization numbers NOTIF461 and NOTIF471 (180 days for the making available on the Belgian market and an additional maximum period of 180 days). If the Commission shall adopt an implementing decision stating that the Union Authorization has not been granted, the biocidal products concerned will no longer be made available on the market 180 days after the implementing regulation has been adopted. The use of existing stocks remains authorized until a maximum of 365 days after the date of adoption of the Commission decision.

Best regards,

A. Rihoux