ENVIRONMENT AGENCY

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Minor Change on request of National Authorisation for the biocidal product family HCl Family A

The Environment Agency of Iceland (Umhverfisstofnun) received your application for National Authorisation - minor change on request of HCl Family A 27th December 2017. The case was accepted by the agency 3rd April 2018. In addition to the material in the application documents the agency has based the authorisation on the authorisations of the Danish Environmental Protection Agency.

This authorisation is granted in exercise of the powers conferred by Articles 50 (2) and (3)(b) of Regulation (EU) No 528/2012 and Article 7 (7) of Regulation (EU) No 354/2013 (on changes) in accordance with Article 1 (6) of Icelandic Regulation No 878/2014 on biocidal products, which implemented Regulation (EU) No 354/2013 into Icelandic legislation.

The Environment Agency of Iceland granted authorisation to HCl Family A, with the asset number IS-0016070-0000 under Icelandic Biocidal Regulation 1101/2004.

We hereby confirm that we accept the notification for a minor change on request for addition of non-active substances intentionally incorporated in the product.
The notification concerns

- Addition of new dye, Duasyn Ink Blue SLK.
- Addition of new fragrance, Fresh 291674.

as referred to in Section 1 of Title 2 to the Annex to Regulation (EU) No 354/2013.

The Environment Agency of Iceland amends the asset number IS-0016070-0000 in the following terms:

1. The biocidal product and restrictions outlined in document number UST201610-115 are updated by the relevant conditions and restrictions as listed in Appendix 1 to this certificate.
2. A Summary of the Product Characteristics, compliant with Article 22 (2) of Regulation (EU) No 528/2012 is listed in Appendices 2.1 – the relevant criteria for this biocidal family authorisation applies as described therein.

Sincerely

Gunnlaug H. Einarsdóttir
Director

Hafdís Inga Ingvarsdóttir
Advisor

Appendix 1: Modified Biocidal Product Characteristics

Appendix 2.1: Summary of Product Characteristics for a Biocidal Product

Appendix 2.2 Conditions of Authorisation

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