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Subject: Renewal of authorisation subject to mutual recognition for placing the biocidal product, *ContraC Blox*, on the market in Iceland

The Environment Agency of Iceland (Umhverfisstofnun) received your application for renewal of national authorisation subject to mutual recognition of *ContraC Blox* on 24th October 2017. The case was accepted by the Agency on 9th July 2018 and validated on 10th July 2018.

The Agency based the decision on the evaluation of the reference Member State, United Kingdom, Health and Safety Executive.

The Environment Agency of Iceland hereby grants a renewal of authorisation for placing the biocidal product **ContraC Blox** on the market in Iceland, by mutual recognition of product authorisation UK-2015-0941 issued by the United Kingdom, Health and Safety Executive, in accordance with Article 5 of Icelandic Regulation No 878/2014 on biocidal products, which implements Regulation (EU) No 528/2012 into Icelandic legislation. The Product Assessment Report is accessible under the authorisation in the R4BP 3 database.

This authorisation is granted in exercise of the powers conferred by Articles 17(3) and 19(1) of Regulation (EU) No 528/2012 with the authorisation number IS-2016-0001.

The Environment Agency of Iceland has determined that the conditions in Articles 19 and 32(2) of Regulation (EU) No 528/2012 are still met and renews this authorisation with the following terms:

1. This authorisation revokes the authorisation document reference number UST201510-049 (as supplemented or amended from time to time).
2. The composition and formulation established for the biocidal product is detailed in the Summary of the Product Characteristics in Appendices 1 and 2 – the relevant criteria for this biocidal product authorisation apply as described therein.
3. Subject to compliance with the conditions as listed in Appendix 3, the authorisation holder is authorised to place on the market the biocidal product detailed in the Summary of the Product Characteristics (Appendix 1) for the use(s) set out in that document.
4. This authorisation and associated documents outlined in the Summary of the Product Characteristics may be amended in accordance with Article 48 and 50 of Regulation (EU) No 528/2012.

5. This authorisation and associated documents outlined in the Summary of the Product Characteristics may be cancelled in the circumstances set out in Article 48 and 49 of Regulation (EU) No 528/2012.
6. When placing the above mentioned biocidal product on the market in Iceland, the products shall be labelled according to Article 69 of Regulation (EU) No 528/2012 and if the biocidal product is classified as hazardous according to Regulation (EU) No 1272/2008 (CLP), such labelling shall be in Icelandic, cf. Article 4 of Regulation No 878/2014 on biocidal products
7. Subject to paragraphs 4 and 5, this authorisation remains in force until midnight of 10th June 2023, on the condition that the active substance is registered in EU list of approved active substances

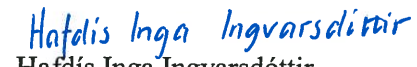
This administrative decision may be appealed before the Minister for the Environment and Natural Resources, in accordance with Article 26 of the Icelandic Administrative Act No 37/1993

Appeals should be directed, within three months as of 11th July 2018, to the Ministry for the Environment and Natural Resources, Skuggasundi 1, 101 Reykjavík, Iceland.

Sincerely



Gunnlaug H. Einarsson
Director



Hafdis Inga Ingvarsdottir
Advisor

Appendix 1: Summary of Product Characteristics for a Biocidal Product

Appendix 2: Confidential Biocidal Product Characteristics

Appendix 3: Conditions of Authorisation