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Icelandic authorisation of the product family, CVAS Biocidal Product Family based on L(+) Lactic Acid, authorised by a Union authorisation

Regulation (EU) No 528/2012 concerning the making available on the market and use of biocidal products, is implemented into Icelandic legislation through the Icelandic Regulation No 878/2014 on biocidal products.

The Environment Agency of Iceland (Umhverfisstofnun) refers to Commission Implementing Regulation (EU) 2023/1311 of 27 June 2023, granting a Union authorisation for the biocidal product family, CVAS Biocidal Product Family based on L (+) Lactic Acid. When the Commission grants a Union authorisation or decides that a Union authorisation has not been granted, the EFTA states will, according to the EEA agreement Annex II Chapter XV point 12n (e), simultaneously and within 30 days of the Commission Act take corresponding decisions.

The Environment Agency of Iceland (Umhverfisstofnun) hereby accepts the European Commission's decision on granting a Union Authorisation for the biocidal product family, CVAS Biocidal Product Family based on L (+) Lactic Acid, by publishing a summary of the decision on the [Agency's website](#). The product is authorised under the terms and conditions as described in the Icelandic Summary of Product Characteristics (SPC) attached to the R4BP3 asset.

When placing the above-mentioned biocidal product family on the market in Iceland, the products shall be labelled in accordance with the terms and conditions in the Icelandic SPC and Article 69 of Regulation (EU) No 528/2012. If the products are classified as hazardous according to Regulation (EU) No 1272/2008 (CLP), such labelling shall be in Icelandic cf. Article 32 of the Chemicals Act No 61/2013 (see section 6 of the Icelandic SPC).

Sincerely

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