ENVIRONMENT AGENCY

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Icelandic authorisation of the product, Bioquell HPV-AQ, 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) 2022/1226 of 14 July 2022, granting a Union authorisation for the biocidal product, Bioquell HPV-AQ. 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, Bioquell HPV-AQ, by publishing a summary of the decision on the Agency's website (ust.is).

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 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 Summary of Product Characteristics (SPC)).

Sincerely

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