

Jotun A/S
Postboks 2021
3248 Sandefjord
Iceland

27. April 2018
UST201801-199/H.I.I.
07.06.04

Minor Change on request for the product Visir Oljegrinning Pigmentert

The Environment Agency of Iceland (Umhverfisstofnun) received your application for minor change on request of Visir Oljegrinning Pigmentert on 27th February 2014. The case was accepted by the agency on 16th December 2015. In addition to the material in the application documents the agency has based the authorisation on the authorisations of the Norwegian Environment 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 Visir Oljegrinning Pigmentert, with the authorisation number IS-2012-0003 under Icelandic Biocidal Regulation No 1101/2004.

We hereby confirm that we accept the notification for a minor change on request for replacement of a non-active substance intentionally incorporated in the product.

The notification concerns

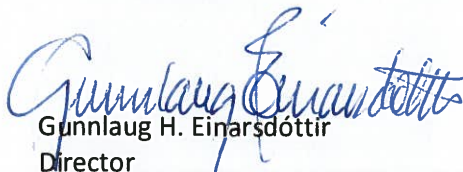
- Change in the formulation, where the non-active raw material DRIER CO/ZR is substituted with FE DRIER and water.

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 authorisation number IS-2012-0003 in the following terms:

1. A Summary of the Product Characteristics, compliant with Article 22 (2) of Regulation (EU) No 528/2012 is listed in Appendices 1 – the relevant criteria for this biocidal product authorisation applies as described therein.
2. As a result of the change in formulation, the classification has been replaced by "Not Classified".

Sincerely



Gunnlaug H. Einarsdóttir
Director



Hafdis Inga Ingvarsdóttir
Advisor

Appendix 1.1: Summary of Product Characteristics for a Biocidal Product

Appendix 1.2 Conditions of Authorisation

Appendix 1.2 Conditions of Authorisation

A failure to comply with any conditions contained in this Appendix may result in cancellation of the authorisation under Article 48 of Regulation (EU) No 528/2012.

1. Without prejudice to the duties imposed on the Authorisation holder of the biocidal product by Article 69 of Regulation (EU) No 528/2012, the Authorisation holder must include on the product labels the information contained in relevant meta summary of the product characteristics for the biocidal product, other than,
 - The name and address of the manufacturer(s) of the product (including site details);
 - The name and address of the manufacturer(s) of the active substance(s) (including site details); and
 - The list of all authorised pack sizes and types (however the relevant pack size must be on the product label).