NORWEGIAN ENVIRONMENT AGENCY

Authorisation of Biocidal Product

The authorisation is granted under the provisions of the Norwegian Biocides Regulation of 18th December 2003 no. 1848; Articles 7, 8 and 12.

The authorisation comprises:

Product name:	Wolsit KD-10
Authorisation number:	NO-2014-0041
Date of authorisation:	31.01.2014
The authorisation expires:	31.01.2024 (provided renewed inclusion of the active substance(s) on Annex I of Directive 98/8/EC. As of 1st September 2013, Annex I of Directive 98/8/EC will be replaced by a Union list of approved active substances, according to Regulations (EU) No. 528/2012, Article 9).
Product type:	PT 8
Authorisation holder in Norway:	BASF Wolman GmbH Dr. Wolman-Str. 31-33 D-76547 Sinzheim Germany
Active substances:	4.5% w/w propiconazole (4.74% technical propiconazole; CAS nr. 60207-90-1)

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Area(s) of use, restriction(s), potential risk(s) and any other specific conditions of the authorisation are stated in the Summary of Product Characteristics (SPC).

The labelling must be in accordance with the conditions stated in the SPC. In addition, the general provisions on labelling of biocidal products as stated in Article 22 of the Norwegian Biocides Regulation shall be followed. Updated labelling must be submitted to the Norwegian Environment Agency within three months of the date of authorisation.

The authorisation holder must inform the Norwegian Environment Agency immediately of any new information or data leading to changes in the labelling of the product, and revised labelling shall be submitted. Labelling changes concerning product formulation or environmental and/or health properties, including changes to the information that shall be included in the labelling according to Article 22 of the Norwegian Biocides Regulation, must be approved by the Norwegian Environment Agency. Administrative changes, such as change in contact information of manufacturer, importer or distributor, shall be communicated to the Norwegian Environment Agency in accordance with Article 16 of the Norwegian Biocides Regulation, but do not need an approval. The fees concerning changes to the product authorisation are stated in Annex 4 to the Norwegian Biocides Regulation.

We would like to inform you that an annual fee is collected for authorised biocidal products, in accordance with Article 4 of the Norwegian Biocides Regulation.

When Regulations (EU) No. 528/2012 and (EU) No. 354/2013 are implemented in Norwegian law, changes to a product authorisation will be classified and handled according to the procedures stated therein. The fees structure will be adjusted in accordance with Regulation (EU) No. 528/2012.

In accordance with Article 33 of the Norwegian Biocides Regulation, complaints against this decision may be addressed to the Ministry of the Environment. The complaint must be submitted to the Norwegian Environment Agency within three weeks of receipt of this letter, in accordance with the Public Administration Act, Article 28.

Yours sincerely,
Norwegian Environment Agency

Eli Vike
Eli Vike
Head of Section

Kjetil Haugstad
Kjetil Haugstad
Senior Adviser

Enclosure
Summary of Product Characteristic (SPC)