Regulation (EU) No 528/2012 concerning the making available on the market and use of biocidal products

**PRODUCT ASSESSMENT REPORT OF A BIOCIDAL PRODUCT FOR   
NATIONAL   
AUTHORISATION APPLICATION**

(submitted by the evaluating Competent Authority)



**CYPER DP**

Product type 18 INSECTICIDES, ACARICIDES AND

PRODUCTS TO CONTROL OTHER

ARTHROPODS

CYPERMETRHIN as included in the Union list of approved active substances

Case Number in R4BP: BC-HK088169-23

Evaluating Competent Authority: Poland

Date: -/-/2023

1. **Conclusion**

CYPER DP is an insecticide formulated as a dustable powder (DP formulation) containing cypermethrin. The product is an insecticide used indoors and outdoors by non-professional and professional users. All uses can be summarized as follow:

* Crack and crevice application against crawling insects, for non-professional and professional use.
* Direct application against wasp nests, for non-professional and professional use.
* Direct application against ant nests, for non-professional use.
* Spot application around buildings in the vicinity of windows and doorsteps against *Blatta orientalis*, for non-professional use.

CYPERD DP is identical to the biocidal product FREE LAND DUST (R4BP case number: BC-EM058913-29, authorisation number: PL/2023/0608/MR, already authorized in Poland in an NA-MRP procedure from the reference product FREE LAND DUST in Greece (authorisation number: ΤΠ18-0550, asset number in R4BP: GR-0030247-0000).

The Product Assessment Report for the authorisation of FREE LAND DUST prepared by the Greek CA (dated October 2022) is considered applicable to the authorisation of the product CYPER DP in Poland. Only the following changes in the product CYPER DP authorisation were introduced in comparison to the reference product: the name and the trade name of the product, the authorisation holder and the manufacturer of the product.

**Conditions for the authorisation of the biocidal product PX-025 in Poland:**

The conditions for granting an authorisation according to Article 19(1) of Regulation (EU) No 528/2012 are fulfilled. Description of the conditions for the authorisation of this biocidal product is presented in the summary of product characteristics (SPC).

# Administrative information

## 2.1. Identifier of the product

|  |  |
| --- | --- |
| **Identifier** | **Country (if relevant)** |
| CYPER DP | PL |

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### Authorisation holder

|  |  |  |
| --- | --- | --- |
| **Name and address of the authorisation holder** | **Name** | INDUPHARMA srl |
| **Address** | Via Sorgaglia 40, 35020 Arre (PD), Italy |
| **Authorisation number** | PL/2023/0644/MR/SBP | |
| **Date of the authorisation** | 29-11-2023 | |
| **Expiry date of the authorisation** | 18.01.2033 r. | |

### 2.3. Manufacturer of the product

|  |  |
| --- | --- |
| **Name of manufacturer** | INDUPHARMA SRL |
| **Address of manufacturer** | Via Sorgaglia 40, 35020 Arre (PD), Italy |
| **Location of manufacturing sites** | Via Sorgaglia 40, 35020 Arre (PD), Italy |