# Supporting document for the pre-submission phase for Union authorisation under Regulation (EU) No 528/2012

**Company identifier (asset owner/ prospective authorisation holder):**

|  |  |
| --- | --- |
| **Company name** | **Company UUID** |
|  |  |

**Name of the biocidal product / biocidal product family:**

**Name of the active substance(s):**

**Product Type(s) (PT):**

**Evaluating Competent Authority (eCA):**

**Existing authorisations or notifications:**

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
| **Member State** | **Name** | **Identifier** | **Current legal status[[1]](#footnote-1)**  | **Comments** |
|  |  |  |  |  |

**Comments**

**Comments:**

The provided information will represent the basis for the pre-submission consultation, in order to:

* obtain the confirmation required in accordance with Article 43(1) of BPR that the product would have similar conditions of use across the Union;
* seek confirmation that the product falls within the scope of the BPR; and
* identify the appropriate Product Type (PT).

In case ECHA needs to contact you in reference to the current application please indicate your contact details, **if different from the ones specified in your R4BP 3/Reach-IT account**. Regulation (EC) 45/2001 or Directive 95/46/EC on the processing of personal data apply. The Data Subject shall at any time have the right to access and rectify its Personal Data.

|  |  |
| --- | --- |
| **Name** |  |
| **Telephone number** |  |
| **E-mail address** |  |

1. Please indicate whether the product is considered as a biocidal product or other. If other, please specify (e.g. medicinal product). [↑](#footnote-ref-1)