# Supporting document for the application for a minor change to a Union authorisation under Regulation (EU) No 354/2013

*Please note that your application cannot be processed if the supporting document is not included with your change application or properly filled in.*

**Authorisation affected by the proposed change(s)**

|  |  |
| --- | --- |
| **Product name\*** | **Asset number\*** |
| Click or tap here to enter text. | Click or tap here to enter text. |

\*as indicated in R4BP 3.

**Description of the proposed change(s) to the product**

##### Minor change(s)

Please provide a detailed description of the proposed change(s) clearly indicating whether the change is an addition or modification. In addition, please justify that the proposed change(s) do not adversely affect the conclusions previously reached concerning the compliance with the conditions set out in Article 19 of Regulation (EU) No 528/2012 (“BPR”). You should reflect on all following items:

* Do the physico-chemical properties, physical hazards and chemical identity fulfil the conditions of Art. 19(1)(c),(d) of the BPR following the change?
* Is efficacy of the product(s) still sufficiently demonstrated following the change?
* Does the human health risk assessment remain unaffected by the change?
* Does the environmental risk assessment remain unaffected by the change?

For a biocidal product family, please indicate the meta SPC(s)/product(s) concerned by the change.

| **#** | **Detailed description of the change** | **Detailed justifications** |
| --- | --- | --- |
|  | Click or tap here to enter text. | **Physico-chemical properties, physical hazards and chemical identity:**Click or tap here to enter text. |
| **Efficacy:**Click or tap here to enter text. |
| **Human Health:**Click or tap here to enter text. |
| **Environment:**Click or tap here to enter text. |

##### Administrative change(s) (optional)[[1]](#footnote-1)

Please select the nature of the change from the dropdown menu and provide a detailed description of the change. It should be clearly indicated whether the change concerns an addition, modification or deletion. In case of modification, please provide both the old and new text.

For a biocidal product family, please indicate the meta SPC(s)/product(s) concerned by the change.

Please note that a transfer of the authorisation to a new holder (Change number 3 of Section 1 of Title 1 of the Annex to the Regulation (EU) No 354/2013) is only feasible through the submission of a UA-TRS application.

Add rows, if necessary.

| **#** | **Nature of the change** | **Detailed description of the change** |
| --- | --- | --- |
|  | Choose an item. | Click or tap here to enter text. |
|  | Choose an item. | Click or tap here to enter text. |

**Annex I. List of all approved or pending changes for the biocidal product/family, following the first authorisation or the last renewal of the product/family**

Please list all changes that you applied for since the first authorisation or the last renewal of the biocidal product/family, whichever is the latest. For each change, please indicate the case number, the type: administrative (ADC), minor (MIC) or major change (MAC), the status: approved / pending and provide a brief description of the change.

| **#** | **R4BP 3 case No** | **Type of change** | **Status** | **Brief description of the change** |
| --- | --- | --- | --- | --- |
|  | Click or tap here to enter text. | Choose an item. | Choose an item. | Click or tap here to enter text. |
|  | Click or tap here to enter text. | Choose an item. | Choose an item. | Click or tap here to enter text. |

1. Used only if the minor change application is combined with administrative changes. [↑](#footnote-ref-1)