**Supporting document for the application for authorisation of the same biocidal product under Regulation (EU) No 414/2013**

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| **Explanatory note to fill in the supporting document** |
| Under Article 1 of Regulation (EU) No 414/2013, the proposed differences between the same biocidal product or product family and the related reference product or product family shall concern only information, which can be subject to an administrative change of product(s) or product family referred to in Title 1 of the Annex to Regulation (EU) No 354/2013.According to Article 2(b) of Regulation (EU) No 414/2013, evidence shall be given that the products are identical on all other aspects. Therefore, please fill in:* the tables of the supporting document, indicating the exhaustive list of all the proposed differences between the same biocidal product or product family and the related reference product or product family. A detailed description of the change(s) is (are) required.
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*Please note that your application cannot be processed if the supporting document is not included with your application or thoroughly compiled in all its sections. Only administrative changes, as defined under Title 1 of the Annex to Regulation (EU) No 354/2013, are accepted.*

**Exhaustive list of the proposed differences between the same biocidal product or product family and the related reference product or product family**

In accordance with Article 1 of Regulation (EU) No 414/2013, the differences between the products shall concern only information that can be subject to an administrative change of product(s) or product family referred to in **Title 1** of the Annex to Regulation (EU) No 354/2013.

If the basis of your application is a product family indicate whether you are applying for the authorisation of a:

[ ]  biocidal product

[ ]  biocidal product family

By submitting this document, you declare that the product(s) are identical, except for the differences exhaustively listed below.

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 text of the related reference product or product family and the new text.

For a biocidal product family, please indicate the meta-SPC(s)/product(s) concerned by the change. In case of a biocidal product family, it is possible to apply for the authorisation of part of the family by requesting the deletion of meta-SPC(s)/product(s), but this should be clearly indicated in the description of the change.

Add rows, if necessary.

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