# Supporting document for the application for same biocidal product to a Union authorisation under Regulation (EU) No 354/2013

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| Under Article 1 of Commission Implementing Regulation (EU) No 414/2013 (“SBP Regulation”), 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 Commission Implementing Regulation (EU) No 354/2013 (“Changes Regulation”).According to Article 2(b) of the SBP Regulation, evidence shall be given that the products are identical on all other aspects.**Please note that your application cannot be processed if the supporting document is not included with your same biocidal product application or properly filled in.****By submitting this document, you declare that your same biocidal product or product family is identical to the related reference product or product family, except for the differences exhaustively listed below. In addition, you declare that the Summary of Product Characteristics (SPC) submitted with your same biocidal product application does not include any other differences than the ones reported below.** |

**Structure of your same biocidal product**

If the basis of your application is a biocidal product family and your same biocidal product application does not cover the full scope of the family, please list the meta SPCs relevant for your application. If your same biocidal product or product family has the same scope as the related reference biocidal product family, the table below does not need to be filled in.

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| **Meta SPCs of the related reference biocidal product family relevant for your application** |
| Click or tap here to enter text. |

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

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.

In case you remove tradenames or a claim, please also indicate it below.

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

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. |
| Click or tap here to enter text. |
|  | Choose an item. | The proposed change concerns Choose an item. |
| Click or tap here to enter text. |

**Letter of access**

Please provide the relevant file names for your letters of access. If you consider that a letter of access is not required to fulfil the requirements of Article 2(c) of the SBP Regulation, please indicate a justification in the “Comments” column. Any clarifications on the scope of the letter of access may be provided to the “Comments” column.

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| **Requirement** | **File name** | **Comments** |
| Valid letter of access to the active substance data[[1]](#footnote-1) | Click or tap here to enter text. | Click or tap here to enter text. |
| Valid letter of access to the related reference product dossier[[2]](#footnote-2) | Click or tap here to enter text. | Click or tap here to enter text. |
| Any other letter of access required to fulfil the requirements set in Article 2(c) of the SBP Regulation | Click or tap here to enter text. | Click or tap here to enter text. |

1. Annex II of Regulation (EU) No 528/2012 [↑](#footnote-ref-1)
2. Annex III of Regulation (EU) No 528/2012 [↑](#footnote-ref-2)