### Information about the product

| No | Information | |
| --- | --- | --- |
| 1 | Product name (indicate complete name of the product) | Swedish product authorisation No |
| 2 | Additional name [[1]](#footnote-1) | |
| 3 | Attach   * documentation showing that the product that will be placed on the market under an additional name is essentially similar to the authorised biocidal product [[2]](#footnote-2), i.e: * contains the same active substance in the same concentration as the authorised product * has been produced using the same method as the authorised product * has the same function and properties as the authorised product * fulfils the same safety requirements as the authorised product does. * a draft label in Swedish including risk information and safety advice as well as other prescribed label text and symbol, if any (KIFS 2022:3). | |

### Information about the applicant[[3]](#footnote-3)

|  |  |  |  |
| --- | --- | --- | --- |
| Complete company name/name | | Organisation number | |
| Street address | | Telephone number | Fax number |
| Postal code and town | | Contact person | |
| Country | | E-mail address | |
| Applicant’s registration certificate  Yes, Appendix No: | A registration certificate must be supplied by all companies that are not, or have not been, a registration holder/permanent representative or notifier of additional name for an authorised biocidal product in Sweden over the past year. Registration certificates can also be required if there have been changes since the last product authorisation or if a long time has passed since a registration certificate was submitted. | | |

### Signature

|  |  |
| --- | --- |
| Place and date | Company |
| Signature [[4]](#footnote-4) | Name (please print) |

|  |  |
| --- | --- |
| **Send the application to:**  Kemikalieinspektionen  Box 2  SE-172 13 Sundbyberg, Sweden |  |

1. The name must not lead to confusion with other biocidal products or otherwise be contrary to Chapter 4, Section 4, third paragraph of the Pesticides Ordinance (2014:425). [↑](#footnote-ref-1)
2. This may be a document for which the authorisation holder certifies that the authorised product is essentially similar to the product that will be placed on the market under the additional name. [↑](#footnote-ref-2)
3. The person who will be responsible for placing the product on the market in Sweden under the proposed additional name. [↑](#footnote-ref-3)
4. Applicant or representative (with valid attached letter of authorisation) must sign this form. Documentation proving the right to sign may be requested. [↑](#footnote-ref-4)