# Supporting document for the renewal of a single national authorisation of a biocidal product (or biocidal product family) under Article 31 of Regulation (EU) No 528/2012

*Please note that if the supporting document is not included or filled in the application may not be processed.*

**Please indicate in the tick box below if you are applying for a renewal of a single authorisation of a Biocidal product (or biocidal product family) under Article 31 of the Regulation (EU) No 528/2012:**

[ ]  renewal of authorisation under Article 31 of the Regulation (EU) No 528/2012

N.B.: If the national authorisation that you want to renew is part of a group of national authorisations linked together by mutual recognition and for which there is an agreement between their authorisation holder(s) that the renewal applications are assessed together by the same reference MS, a different supporting document should be used.

**Indicate in the tick boxes below whether you are submitting new information on the active substance as part of your application:**

[ ]  yes

[ ]  no

N.B.: new information on the active substance is information that was not assessed during the approval of the active substance.

List in the table below all the new information on active substance submitted in your application.

| **Active substance[[1]](#footnote-1)** | **Location in the IUCLID dossier** | **File name in the IUCLID dossier** | **Title and author of the document** | **Area the new information concerns** | **Aim of the new information** |
| --- | --- | --- | --- | --- | --- |
| *Indicate here the name and respective CAS number of the active substance the new information is pertinent to.* | *Indicate here under which section of the active substance data set section of the IUCLID dossier is the document available.* | *Indicate here the file name of the document.* | *Indicate here the title and the author of the document concerning new information on the active substance.* | *Indicate here all the areas the new information on the active substance concerns. Indicate “APCP” if the new information concerns Analytical methods and Physico-Chemical Properties, “EFF” if it concerns efficacy, “ENV” if it concerns environment, “HH” if it concerns human health and “Animal health” if it concerns animal health.* | *Describe here more which endpoint value the new information on the active substance affects and in what way (e.g., refinement of an existing endpoint value, data for an endpoint not considered during the active substance approval etc.).* |
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**Indicate in the table below if the new information on the active substance submitted in your application has been submitted for another application and include the details of those applications:**

| **Application type** | **New information submitted in another application****(Y)** | **Name of applicant** | **Asset/case number** | **Date of submission** |
| --- | --- | --- | --- | --- |
| Technical equivalence assessment application that has been closed with the decision that technical assessment was established**[[2]](#footnote-2)** | *Indicate “Y” where the new information on the active substance submitted in your application was submitted in this application type and fill out the row accordingly, otherwise leave the entire row empty.*  | *Indicate the asset owner as stated in R4BP 3.* | *Provide the asset number assigned by R4BP 3. For ongoing applications provide the case number assigned by R4BP 3.* | *Indicate the date of submission as stated in R4BP 3.* |
| Inclusion in the Article 95 (active substance supplier) list application closed with the decision of inclusion on the Article 95 list2 | *Indicate “Y” where the new information on the active substance submitted in your application was submitted in this application type and fill out the row accordingly, otherwise leave the entire row empty.*  |  |  |  |
| Biocidal product/biocidal product family authorisation, renewal or change application2 | *Indicate “Y” where the new information on the active substance submitted in your application was submitted in this application type and fill out the row accordingly, otherwise leave the entire row empty. In case your product/product family contains more than one active substance indicate here which active substance the new information is pertinent to.* |  |  |  |

In case you indicated above that the new information on the active substance was submitted in a biocidal product or biocidal product family authorisation, change or renewal application, fill out the additional table below:

| **Asset/case number[[3]](#footnote-3)** | **Procedure type** | **Product name** | **Reference Member State/ evaluating Competent Authority** |
| --- | --- | --- | --- |
| *Provide the asset numbers assigned by R4BP 3 of all the relevant applications. For ongoing applications provide the case number assigned by R4BP 3 of all the relevant application types.* | *Indicate “NA” if the application concerns a national authorisation procedure. Indicate “SA” if the application concerns a simplified authorisation procedure. Indicate “UA” if the application concerns a Union authorisation procedure.* | *Indicate the product name as stated in R4BP 3.* | *Indicate here the reference Member State for the relevant national authorisation (NA) application, or evaluating Competent Authority for the relevant simplified (SA) or Union authorisation (UA) application.* |
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1. Add rows if necessary [↑](#footnote-ref-1)
2. Add rows if you the new information on the active substance submitted in your application has been submitted for more than one application of this type. [↑](#footnote-ref-2)
3. Add rows if necessary [↑](#footnote-ref-3)