**General instructions for parts A and B**

* This form is for application of new plant strengtheners (corroboranti) for the Italian Input List. Compliant products will also be included in the European Input List.
* Only registered companies may submit products. If your company is not yet registered, please complete the ‘company registration form’ and submit it simultaneously with this application.
* Fill in the form electronically only. Forms filled in hand-writing are not accepted.
* Sign and send to FiBL by e-mail (contact: see <https://italy.inputs.eu/submit-products/>).

**Part A, Basic information on composition and properties of the product**

**Instructions for part A**

* Part A of the application form concerns the composition and properties of products. If the Italian distributor of the product (see part B) does not have the necessary knowledge, part A may be submitted by another company (usually the manufacturer).
* For all information concerning product composition, FiBL will exclusively correspond with the company completing part A.
* If the composition and properties of a product has already been communicated to FiBL in the context of the **Swiss** or the **Dutch** Input List, part A does not need to be completed. Instead, the identity of the product must be confirmed with the ‘letter of access’.

**A.1 Submission of part A**

|  |  |
| --- | --- |
| Company submitting part A  *Companies must be registered with a separate form, provided on* [*https://italy.inputs.eu/submit-products.html*](https://italy.inputs.eu/submit-products.html) |  |
| Role of the company | ☐ we are the applicants for the Italian Input List (see part B) |
| ☐ we submit this information to support the application of the following entry:  company: …  product name: … |

A.2 Identification of the product

|  |  |
| --- | --- |
| Original name of the product |  |
| Other trade names of the same product  (also in other countries) |  |
| Is the product identical with another product on the European Input List? | ☐ yes ☐ no  ☐ not known for certain  If yes, please indicate the trade name here: |
| Manufacturer of the product  Name, address |  |

A.3 Description of the product’s uses

|  |  |  |
| --- | --- | --- |
| Use category | ☐ corroborante, plant strengthener, plant aid  ☐ the use category is also indicated on the product label | |
| Conditions of use | ☐ indicated on the product label *Please specify:*  ☐ not indicated on the product label | |
| Why is the product applied to crops? *Please specify* |  |  |
| Applied dose *(in kg/ha or liter/ha)* |  | |
| Number of applications per season |  | |
| In which crops can the product be used? |  | |

A.4 Ingredients of the product

* List all ingredients which are added to the product, including all additives and co-formulants (sum must add to 100%).
* Information on manufacturing of each component needs to be provided through supporting documents.
* Components given in the coloured fields may be published in the Italian Input List

|  | Name of component  *Where applicable, use standard chemical nomenclature* | CAS-Number  *as applicable* | Origin  *(natural; synthetic)* | Amount added (%) | Functionality of the component in the product |
| --- | --- | --- | --- | --- | --- |
| 1 |  |  |  |  |  |
| 2 |  |  |  |  |  |
| 3 |  |  |  |  |  |
| 4 |  |  |  |  |  |
| 5 |  |  |  |  |  |
| 6 |  |  |  |  |  |

A.5 Additional questions on the product’s composition and manufacturing

|  |  |
| --- | --- |
| **Does the product contain any of the following components?** | |
| - preservatives  *If yes, detailed information must be given in section A.4* | ☐ yes ☐ no |
| - synthetic nanoparticles | ☐ yes ☐ no |
| - other additives  *If yes, detailed information must be given in section A.4* | ☐ yes ☐ no |
| - phosphonates  *If yes (also in traces), please provide details* | ☐ yes ☐ no |
| - GMOs or products derived from GMOs *For certain components as specified in the application guidance please provide a declaration on the absence of GMO.* | ☐ yes ☐ no |
| **Further questions** | |
| Are there any synthetic processing steps involved in the manufacturing procedure?  If yes, provide a detailed description in a separate document. | ☐ yes ☐ no |
| Is the product liquid or solid? | ☐ liquid  ☐ solid |
| Location of manufacturing:  *please indicate the manufacturing site (Name, location)* | ... |
| Are there products for conventional agriculture being manufactured at the same location? | yes  no  *If yes, please indicate more details during product submission.* |
| Location where the product is packaged:  *please indicate the packaging site (Name, location)* | … |
| Are there products for conventional agriculture being packaged at the same location? | yes  no  *If yes, please indicate more details during product submission.* |

A.6 Checklist for the documentation for Part A

Use this section to check whether your application contains all necessary documents.

|  |  |  |
| --- | --- | --- |
| **For all** products | *Full composition and manufacturing process* | ☐ enclosed  ☐ provided by: … |
| **For all** products | *Label* | ☐ enclosed  ☐ provided by: … |
| **For all** products | *Technical specification of the product* | ☐ enclosed  ☐ provided by: … |
| **For all** products | *Product datasheet* | ☐ enclosed  ☐ provided by: … |
| **For all** products | Material safety data sheets | ☐ enclosed  ☐ provided by: … |
| *Only for components with a* ***GMO risk*** | *Confirmation of the absence of GMOs (please use the form provided on the project website)* | ☐ enclosed  ☐ not relevant |

A.8 Confirmation and signature for part A

The undersigned confirms that the information given in this form is correct and complete, that the placing on the market of the product complies with the relevant EU law and national legislation for the respective country and that no plant protection effects will be claimed for this product. The undersigning party moreover ensures that any changes in the composition and manufacturing will be communicated to FiBL immediately. The company has read and fully agrees with the General Business Contract. The undersigned takes note that FiBL checks the correctness of applications as part of the analytical quality monitoring programme.

|  |  |
| --- | --- |
| Place and date |  |
| Name |  |
| Position in the company |  |
| Signature |  |

**Part B, Supplementary information relevant for the marketing of the product in Italy**

**Instructions for part B**

* Part B of the application form concerns the marketing of the product in Italy. The company submitting part B will be **shown in the Italian Input List**.
* For all administrative questions, FiBL will correspond with the company completing part B.
* The company completing part B is responsible for paying all fees.

B.1 Company shown in the Italian input List

|  |  |
| --- | --- |
| Company name  *Companies must be registered with a separate form.* |  |
| Role  *(multiple answers possible)* | ☐ manufacturer of the product  ☐ distributor of the product in Italy  ☐ other: … |
| Information on full composition | ☐ information on the full composition is supplied in part A together with this application.  ☐ the full composition will be communicated to FiBL by the following company: … |

B.2 Entry in the Italian input List

|  |  |
| --- | --- |
| Trade name of the product in Italy |  |
| Other names of the product  *(must be identical with part A.2)* |  |
| Suggested categorization in the Italian Input List |  |
| Suggested wording concerning composition / uses |  |

B.4 Placing on the market and conformity to legal requirements

|  |  |
| --- | --- |
| Letter sent to MIPAAFT, communicating that the product has been placed on the market | ☐ Copy of the letter attached  ☐ was/ will be submitted by ……. |
| The product is compliant to the presidential decree DPR 55 of 2012 as well as to Annex 2 of the decree DM no. 6793 from 18 July 2018 | ☐ yes ☐ no |
| The product is compliant to the principles of organic farming as specified in the EU Reg. 834/2007 and 889/2008 | ☐ yes ☐ no |

B.5 Product label

*The product label used for commercialization purposes in Italy must be provided to FiBL (PDF)*

B.6 Checklist of necessary documentation for part B

Use this section to check whether your application contains all necessary documents.

|  |  |  |
| --- | --- | --- |
| For **all** products | *Product label* | ☐ enclosed |
| For **all** products | *Product datasheet* | ☐ enclosed |
| For **all** products | *Copy of the letter sent to MIPAAFT, communicating that the product has been placed on the market* | ☐ enclosed |

B.7 Confirmation and signature for part B

The undersigned confirms that the information given in this form is correct and complete, that the placing on the market of the product complies with the relevant EU law and national legislation for the respective country and that no plant protection effects will be claimed for this product. The undersigning party moreover ensures that any changes in the composition and manufacturing will be communicated to FiBL immediately. The company has read and fully agrees with the General Business Contract. The undersigned takes note that FiBL checks the correctness of applications as part of the analytical quality monitoring programme.

|  |  |
| --- | --- |
| Place and date |  |
| First and last name |  |
| Position in the company |  |
| Signature |  |