

Plant protection products in storage and on the field

Request for approval in accordance with Regulation (EU) 2018/848 or NOP

If you plan to use plant protection products, please submit this application form to bio.inspecta and do not use the input before approval. **Using unauthorised inputs can lead to decertification of your crops/fields.**

Please send the completed form with annexes to: international@bio-inspecta.ch

Input approvals can be complex and demanding, thus bio.inspecta recommends to submit the application at least four weeks before the planned date of use.

Information on requirements of Regulation (EU) 2018/848:

According to Reg. (EU) 2018/848, plant protection products as listed in Reg. (EU) 2021/1165, Annex I may be used if plants cannot be adequately protected by promoting natural enemies, choice of adapted varieties, crop rotation, cultivation techniques. Besides particular restrictions for use in organic agriculture indicated in Reg. (EU) 2021/1165, general EU restrictions on use as defined in Reg. (EU) 540/2011 apply. Safeners, synergists, co-formulants and adjuvants according to Regulation (EC) No 1107/2009 are allowed.

Please be aware that only plant protection products in accordance with the national legislation in your country may be used. Instructions on the label of the product have to be followed.

1. Applicant/operator

Name of operator		b.i Number	
------------------	--	------------	--

2. Information about the plant protection product

Product is purchased <input type="checkbox"/>	Product is self manufactured <input type="checkbox"/>
Trade name of the product	
Is the plant protection product registered in your country? Y/N If yes, give the Registration No.	
From which company/distributor do you plan to buy the product?	
Manufacturer name (if known)	
Manufacturer address and website (if known)	

Composition of the product (sum 100% of volume or weight)			
Active substance(s)	CAS no.	%	
Inert substance(s)	CAS no.	%	Function (e.g. synergist, safener, co-formulant (such as e.g. surfactant, emulsifier, anti-dusting agent, anti-caking agent, etc.), adjuvant)

Possible genetic modification of ingredients or substances/microorganisms used for production: provide evidence for absence		
Product/process	Possible use/contamination with GMO	Evidence for absence (name document here and submit it with application form)
Microbial products	Used mirco-organisms may have been subject to recombinant DNA techniques or other forms of genetical engineering.	
Production of one or more ingredient in a fermenter system using micro-organisms	Used micro-organisms may have been subject to genetical engineering.	
Ingredients of agricultueal origin (e.g. vegetable oil)	Crop may have been genetically modified.	
Mineral oil		
Mineral oil named in the list of ingredients above (incl. CAS-Number) is composed predominately of paraffinic and naphthenic fractions with 50 percent boiling point (10 mm Hg) between 415 and 440 °F (212 and 226°C)?		<input type="checkbox"/> yes <input type="checkbox"/> no
Additional information considered relevant		

3. Application

The plant protection product shall be evaluated in accordance with	<input type="checkbox"/> Reg. (EU) 2018/848 <input type="checkbox"/> NOP <input type="checkbox"/> Other
Crops concerned	
Quantity/ha on which the product is planned to be applied	
Mode of application (spraying, dipping etc.)	
Reason of use (specify pest/disease threatening the crop)	

4. Please attach the following

- ☐ Technical data sheet / product specifications (formulation)
- ☐ Safety data sheet, other relevant data sheets (if applicable)
- ☐ Products approved for use according to an EU-equivalent standard or NOP by an accredited certification body: add copy of the certificate or approval documents / Link to OMRI-list for NOP
- ☐ Labels and recommendations for use
- ☐ Registration in your country
- ☐ **Important:** For products containing microorganisms or risky ingredients (corn, canola, soy etc): a declaration of compliance with the prohibition of genetically modified organisms pursuant to Reg. (EU) No 2018/848 or USDA National Organic Program NOP, Please refer to the website of bio.inspecta. www.bio-inspecta.ch, *Vendor Declaration non-GMO, 24_2636EN*)

☐ Other:

5. Signature

- a) **This form may be filled and signed by the operator** if one or several of the following points apply:
- (EU) A detailed technical data sheet by the manufacturer is attached.
 - (NOP) A detailed technical data sheet by the manufacturer is attached, providing information about all active and inactive ingredients and how they are obtained.
 - (EU, NOP) A confirmation that the product qualifies for use in organic agriculture according to the relevant standard by a certification body accredited to the relevant standard is attached.
 - (NOP) The product is listed by OMRI.

Operator name		
..... Place/Date Name Signature

- b) **This form must be filled and signed by the manufacturer of the product** if the points under a) are not applicable.

Manufacturer name		
..... Place/Date Name Signature and Stamp

6. To be completed by bio.inspecta

Final assessment		
<input type="checkbox"/> Request is approved	<input type="checkbox"/> Reg. (EU) 2018/848	
	<input type="checkbox"/> NOP	<input type="checkbox"/> Other
<input type="checkbox"/> Request is rejected	<input type="checkbox"/> Reg. (EU) 2018/848	
	<input type="checkbox"/> NOP	<input type="checkbox"/> Other
Explanation & justification		
.....		
..... Place/Date Signature & Name	
<p>This approval is valid for three years from the date of singnatur of bio.inspecta. It is in the responsibility of the operation to use only inputs, which have been approved for the respective standard.</p>		