#### MINOR USES COORDINATION FACILITY

# Survey 2022 on Minor Uses work in EU Member States, plus the United Kingdom, Norway and **Switzerland**

## Privacy notice:

We will treat all your Personal Data as confidential. The information provided by you in this questionnaire will be used in the context of the Minor Uses work in EU Member States, plus the United Kingdom, Norway and Switzerland. The data collected in this survey will be stored on Minor Uses' servers. We will keep it on a secure server and we will fully comply with all applicable privacy regulations and consumer legislation. Personal data communicated for the purpose of the survey will be handled in compliance with the EU General Data Protection Regulation nº 2016/679.

### Right of rectification:

Should you require further information concerning the processing of your personal data or to exercise your rights (e.g. access or to rectify any inaccurate or incomplete data) please contact the following email address: contact@minoruses.eu

You have the right of recourse at any time to the Data Protection Officer at: <a href="mailto:dpo@eppo.int">dpo@eppo.int</a>

# The "Survey 2022 on Minor Uses work in EU Member States, plus the United Kingdom, Norway and Switzerland" is split into 8 separate parts:

- Responsibility for minor uses in your country
- The organization of minor uses work in your country and general questions on minor uses and minor crops
- Trials
- Article 51 (Extension of authorizations for minor uses)
- Risk assessment
- Mutual recognition
- Draft registration report
- General topics

The survey is expected to take up to 1 hour.

By keeping the link received in your email you can come back to the survey in case you need to change or modify your responses at a later stage before finalising your response.

The information which you provided in the survey 2017 is accessible in the MUCF Extranet by clicking on this link.

The information which you provided in 2021 is included as prefilled answers in the new survey.

# The deadline to complete the survey is 15th of September 2022.

We thank you in advance for your participation. The information gathered should be of great help to all Minor Uses experts and stakeholders.

The results of the new survey will be presented at the beginning of 2023 in our HEG and CEG meetings.

For any questions on the survey please contact Barbara EDLER be@minoruses.eu - for any technical averable and and the functioning of the curvey, please contact Nathalic POLITION of

Please indicate hereafter your contact details if you are the person responsible for filling in the survey in your country
First name
Last name
Country
Your email

## A. Responsibility for minor uses in your country

- 1. Who is/are the national contact point(s)?
  Please correct information here below if needed in the 'Comments' box
- 2. Who is/are your representative(s) in the Commodity Expert Group(s)? Please correct information here below if needed in the 'Comments' box

# B. The organization of minor uses work in your country and general questions on minor uses and minor crops.

The information which you provided in the survey 2017 is accessible in the MUCF Extranet by clicking on this link.

- 1. Does your organisation have a national list of minor crops available? Please correct information here below if needed in the 'Comments' box
- 2. How often do you update this list of minor crops?
- 3. What are your country's criteria to define a minor crop? (e.g maximum production area (ha), production volume, dietary intake, etc.)? Please correct information here below if needed in the 'Comments' box
- 4. What is the approximate acreage [ha] of cultivated crops in your country?
- 5. Please specify the year when data was collected.
- 6. Please specify the reference from where data was collected
- 7. What is the approximate crop production value [€] of cultivated crops in your country?
- 8. Please specify the year when data was collected.
- 9. Please specify the reference from where data was collected
- 10. What is the approximate acreage [ha] of minor crops in your country?
- 11. Please specify the year when data was collected
- 12. Please specify the reference from where data was collected
- 13. What is the approximate crop production value [€] of minor crops in your country?
- 14. Please specify the year when data was collected
- 15. Please specify the reference from where data was collected
- 16. What are your country's criteria to define a minor use? (e.g.public interest)? Please comment if information hereunder is not correct
- 17. Does your organisation have a national list of minor uses available? Please comment if information hereunder is not correct
- 18. How often do you update this list of minor uses?
- 19. Please check the useful links that you provided and which are listed in the home page of EUMUDA for your country, Please correct information here below if needed in the 'Comments' box

20. Which (working) group(s) is/are involved in minor uses in you country, and what is their role?

### C. TRIALS

The information which you provided in the survey 2017 is accessible in the MUCF Extranet by clicking on this link.

- 1. Do you have research facilities to carry out trials (GEP, GLP, monitoring trials, etc.)?
- 2. If no, which organization is carrying out trials (efficacy and residues) in minor uses in your country?
- 3. Is your organization able to finance trials to generate data for minor uses extensions (efficacy or residue)?
- 4. If you have data, would you be willing to share them with other countries?
- 5. If yes, what would you require in order to share the data (e.g., data sharing fee, etc)?
- 6. If you do not have trial data yet, does your organization intend to produce trial data (efficacy or residue) in the near future?

# D. Article 51 (Extension of authorizations for minor uses)

- 1. Who is eligible to apply for Art.51 extensions in your country? Please select below (multiple answers possible)
- 2. If other, please specify
- 3. Who primarily applies for Art.51 extensions in your country? Please select below (multiple answers possible)
- 4. If other, please specify
- 5. Which organization in your country evaluates the applications?
- 6. How much do you charge for an Art. 51 extension (please specify if the fee mentioned is per crop, per application, are there different fees for zonal applications )?
- 7. Does your country require efficacy data for an Art.51 extension?
- 8. If yes, please specify or provide details
- 9. Have you dealt with the data protection claims for minor uses (acc. to Art.51) in your country?
- 10. If yes, please specify or provide details
- 11. Have you experienced any challenges in your country's minor uses authorization (acc. to Art.51)?
- 12. If yes, please specify or provide details
- 13. How is the zonal application (acc. to Art. 51) for minor uses carried out in your country? Please provide some details
- 14. Do you use the 'public interest' as criterion to grant a registration according to Art. 51?
- 15. How is the 'public interest' assessed in your country? Please specify or provide some details

# E. Risk assessment

- 1. How is the risk assessment for minor uses carried out in your country? Please provide some details
- 2. Do you have special national legislation or provisions for the risk assessment? Please provide some details
- 3. If the risk envelope concept does not cover uses, do you have similar data requirements for Art.51 applications as for label extension after Art. 33 ?
- 4. If yes, what are the data requirements? Please provide some details

# F. Mutual recognition

- 1. Please describe the procedure for mutual recognition in your country
- 2. What obstacles do you see within the procedure of mutual recognition acc. to Art. 51? Please provide some details
- 3. Have you received or approved any minor use applications (acc. Art.51) for mutual recognition from other countries?
- 4. If yes, who is the applicant (e.g. the authorization holder, grower's association)? Please specify
- 5. Have you received an application where the applicant was a third party, and the authorization holder refused its consent (the country's competent authority may still accept the application on public interest grounds)?
- 6. If yes, How did you handle that authorization procedure, for example, to access the registration dossier? Please provide some details

# **G.** Draft registration report

An EU format for the draft Registration Report (dRR) is available to ensure a uniform reporting method for the plant protection product assessment.

Applicants must use the draft Registration Report (dRR) when applying for a plant protection product authorization.

- 1. Is this complied by applicants in your country?
- 2. What obstacles do you see using the dRR for the minor use (acc. to ARt. 51)? Please provide some details
- 3. If it is impossible to prepare a registration report, part A (acc. to ARt. 51 authorizations), is it possible to provide some basic information about the authorization acc. to Art.51 in your country (e.g., a complete GAP table)?

### H. General topics

- 1. What information will be produced after a minor use authorization (Art. 33, 40(1), 40(2), 51(1)-(6), 51(7)? For example the registration report, Part A, and the GAP table. Please provide some details
- 2. How do you ensure that the information from the authorized GAP is available for other countries? Please provide some details
- 3. Do you have a specific procedure in place for minor use applications for low-risk PPPs (e.g., reduced application fees, a shortened evaluation timeline, etc.)?
- 4. If yes, please provide some details