### Status: Point in time view as at 13/10/2009.

Changes to legislation: There are outstanding changes not yet made to Commission Decision of 13 October 2009 establishing standard reporting formats for presenting the monitoring results of the deliberate release into the environment of genetically modified organisms, as or in products, for the purpose of placing on the market, pursuant to Directive 2001/18/EC of the European Parliament and of the Council (notified under document C(2009) 7680) (Text with EEA relevance) (2009/770/EC). Any changes that have already been made to the legislation appear in the content and are referenced with annotations. (See end of Document for details)

Commission Decision of 13 October 2009 establishing standard reporting formats for presenting the monitoring results of the deliberate release into the environment of genetically modified organisms, as or in products, for the purpose of placing on the market, pursuant to Directive 2001/18/ EC of the European Parliament and of the Council (notified under document C(2009) 7680) (Text with EEA relevance) (2009/770/EC)

Article 1 The reporting formats set out in Annexes I and II...
This Decision is addressed to the Member States.
Signature

### ANNEX I

# MONITORING FRIED CREATE TO RIGHT WAIT TO SHARE THE CHILD BY SHARE THE

- 1. General information
  - 1.1. Crop/trait(s): ...
  - 1.2. Decision authorisation number pursuant to Directive 2001/18/EC, and number and...
  - 1.3. Decision authorisation number and date of authorisation pursuant to Regulation...
  - 1.4. Unique identifier: ...
  - 1.5. Reporting period from xx/xx/xx to xx/xx/xx
  - 1.6. Other monitoring reports have been submitted in respect of:
- 2. Executive summary
- 3. Monitoring results
  - 3.1. General surveillance
    - 3.1.1. Description of general surveillance
    - 3.1.2. Details of surveillance networks used to monitor environmental effects during...
    - 3.1.3. Details of information and/or training provided to operators and users,...
    - 3.1.4. Results of general surveillance
    - 3.1.5. Additional information
    - 3.1.6. Review of peer-reviewed publications Appendix
  - 3.2. Case-specific monitoring
    - 3.2.1. Description and results of case-specific monitoring (if applicable)
    - 3.2.2. Monitoring and reporting of adverse effects resulting from accidental spillage...
  - 3.3. Concluding remarks
- 4. Summary of results and conclusions
- 5. Adaptation of the monitoring plan and associated methodology for future...

Document Generated: 2024-07-01

#### Status: Point in time view as at 13/10/2009.

Changes to legislation: There are outstanding changes not yet made to Commission Decision of 13 October 2009 establishing standard reporting formats for presenting the monitoring results of the deliberate release into the environment of genetically modified organisms, as or in products, for the purpose of placing on the market, pursuant to Directive 2001/18/EC of the European Parliament and of the Council (notified under document C(2009) 7680) (Text with EEA relevance) (2009/770/EC). Any changes that have already been made to the legislation appear in the content and are referenced with annotations. (See end of Document for details)

### Appendix 1

## REVIEW OF PEER-REVIEWED PUBLICATIONS

Some publications may contain material relevant to more than one...

## Appendix 2

### **EXPLANATORY NOTES**

- A. General comments
- B. Instructions for completing the format
- C. Confidentiality
  - C.1. Applications submitted pursuant to Directive 2001/18/EC
  - C.2. Applications submitted pursuant to Regulation (EC) No 1829/2003
    - 1. GENERAL INFORMATION
    - 2. EXECUTIVE SUMMARY
    - 3. MONITORING RESULTS
      - 3.1. General surveillance
        - 3.1.1. Description of general surveillance
        - 3.1.2. Details of surveillance networks used to monitor environmental effects during...
        - 3.1.3. Details of information and/or training provided to operators and users,...
        - 3.1.4. Results of general surveillance
        - 3.1.5. Additional information
        - 3.1.6. Review of peer-reviewed publications
      - 3.2. Case-specific monitoring
        - 3.2.1. Results of case-specific monitoring (if applicable)
        - 3.2.2. Monitoring/reporting of adverse effects resulting from accidental spillage (if applicable)...
      - 3.3. Concluding remarks
    - 4. SUMMARY OF RESULTS AND CONCLUSIONS
    - 5. ADAPTATION OF THE MONITORING PLAN AND ASSOCIATED METHODOLOGY FOR FUTURE...

### ANNEX II

## MONITORING FREDORFITH GRIEDORFITH GRIEDORF

- 1. General information
  - 1.1. Crop/trait(s): ...
  - 1.2. Decision authorisation number pursuant to Directive 2001/18/EC and number and...
  - 1.3. Decision authorisation number and date of authorisation pursuant to Regulation...
  - 1.4. Unique identifier: ...
  - 1.5. Reporting period from xx/xx/xx to xx/xx/xx
  - 1.6. Other monitoring reports have been submitted in respect of:

Document Generated: 2024-07-01

#### Status: Point in time view as at 13/10/2009.

Changes to legislation: There are outstanding changes not yet made to Commission Decision of 13 October 2009 establishing standard reporting formats for presenting the monitoring results of the deliberate release into the environment of genetically modified organisms, as or in products, for the purpose of placing on the market, pursuant to Directive 2001/18/EC of the European Parliament and of the Council (notified under document C(2009) 7680) (Text with EEA relevance) (2009/770/EC). Any changes that have already been made to the legislation appear in the content and are referenced with annotations. (See end of Document for details)

- 2. Executive summary
- 3. Uses of GMOs other than cultivation
  - 3.1. Commodity imports into the Community
    - 3.1.1. Commodity crop (GM + non-GM) imports into the Community by country of...
    - 3.1.2. Commodity Crop (GM + non-GM) imports into the Community by country of...
    - 3.1.3. Analysis of data provided in tables 3.1.1 and 3.1.2
  - 3.2. General surveillance
    - 3.2.1. Description of general surveillance
    - 3.2.2. Details of industry, environmental, food and/or feed related surveillance networks...
    - 3.2.3. Details of information and/or training provided to importers, traders, handlers,...
    - 3.2.4. Results of general surveillance
    - 3.2.5. Additional information
    - 3.2.6. Review of peer-reviewed publications Appendix
  - 3.3. Case-specific monitoring
    - 3.3.1. Description and results of case-specific monitoring (if applicable)
    - 3.3.2. Processing (if applicable)
    - 3.3.3. Monitoring and reporting of adverse effects resulting from accidental spillage...
  - 3.4. Concluding remarks
- 4. Summary of results and conclusions
- 5. Adaptation of the monitoring plan and associated methodology for future...

#### Appendix 1

## REVIEW OF PEER-REVIEWED PUBLICATIONS

Some publications may contain material relevant to more than one...

## Appendix 2

## **EXPLANATORY NOTES**

- A. General comments
- B. Instructions for completing the format
- C. Confidentiality
  - C.1. Applications submitted pursuant to Directive 2001/18/EC
  - C.2. Applications submitted pursuant to Regulation (EC) No 1829/2003
    - 1. GENERAL INFORMATION
    - 2. EXECUTIVE SUMMARY
    - 3. GMO USES OTHER THAN CULTIVATION
      - 3.1. Commodity imports into the Community
        - 3.1.1. Commodity Crop (GM + non-GM) imports into the Community by country of...

Document Generated: 2024-07-01

Status: Point in time view as at 13/10/2009.

Changes to legislation: There are outstanding changes not yet made to Commission Decision of 13 October 2009 establishing standard reporting formats for presenting the monitoring results of the deliberate release into the environment of genetically modified organisms, as or in products, for the purpose of placing on the market, pursuant to Directive 2001/18/EC of the European Parliament and of the Council (notified under document C(2009) 7680) (Text with EEA relevance) (2009/770/EC). Any changes that have already been made to the legislation appear in the content and are referenced with annotations. (See end of Document for details)

- 3.1.2. Commodity Crop (GM + non-GM) imports into the Community by country of...
- 3.1.3. Analysis of data provided in tables 3.1.1 and 3.1.2
- 3.2. General surveillance
  - 3.2.1. Description of general surveillance
  - 3.2.2. Details of industry, environmental, food and/or feed related surveillance networks...
  - 3.2.3. Details of information and/or training provided to importers, traders, handlers,...
  - 3.2.4. Results of general surveillance
  - 3.2.5. Additional information
  - 3.2.6. Review of peer-reviewed publications Appendix
- 3.3. Case-specific monitoring
  - 3.3.1. Results of case-specific monitoring (if applicable)
  - 3.3.2. Processing (if applicable)
  - 3.3.3. Monitoring and reporting of adverse effects resulting from Accidental Spillage...
- 3.4. Concluding remarks
- 4. SUMMARY OF RESULTS AND CONCLUSIONS
- 5. ADAPTATION OF THE MONITORING PLAN AND ASSOCIATED METHODOLOGY FOR FUTURE...

Status: Point in time view as at 13/10/2009.

Changes to legislation: There are outstanding changes not yet made to Commission Decision of 13 October 2009 establishing standard reporting formats for presenting the monitoring results of the deliberate release into the environment of genetically modified organisms, as or in products, for the purpose of placing on the market, pursuant to Directive 2001/18/EC of the European Parliament and of the Council (notified under document C(2009) 7680) (Text with EEA relevance) (2009/770/EC). Any changes that have already been made to the legislation appear in the content and are referenced with annotations. (See end of Document for details)

- (1) OJ L 106, 17.4.2001, p. 1.
- (2) OJ L 268, 18.10.2003, p. 1.
- (**3**) OJ L 280, 18.10.2002, p. 27.

### **Status:**

Point in time view as at 13/10/2009.

## **Changes to legislation:**

There are outstanding changes not yet made to Commission Decision of 13 October 2009 establishing standard reporting formats for presenting the monitoring results of the deliberate release into the environment of genetically modified organisms, as or in products, for the purpose of placing on the market, pursuant to Directive 2001/18/EC of the European Parliament and of the Council (notified under document C(2009) 7680) (Text with EEA relevance) (2009/770/ EC). Any changes that have already been made to the legislation appear in the content and are referenced with annotations.