



EUROPEAN COMMISSION

Health and Food Safety Directorate General

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**Standing Committee on Plants, Animals, Food and Feed**  
**Section *Phytopharmaceuticals – Pesticide Residues***  
**21 - 22 November 2022**

**CIRCABC Link:** [https://circabc.europa.eu/ui/group/95a86e0e-0cfe-4354-8d9f-c447c6e85c1b/library/1f0b4026-83ab-48ce-8ae8-a2f5e580e131?p=1&n=10&sort=modified\\_DESC](https://circabc.europa.eu/ui/group/95a86e0e-0cfe-4354-8d9f-c447c6e85c1b/library/1f0b4026-83ab-48ce-8ae8-a2f5e580e131?p=1&n=10&sort=modified_DESC)

<b>AGENDA</b>
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**Section A      Information and/or discussion**

**A.01** Art. 12 and Art. 10 of Regulation (EC) No 396/2005 procedures:

1. Confirmatory data Art. 12 follow-up
  - a) Cases where EFSA RO has been published
  - b) Missing analytical standards follow up
2. List of non-approved substances for follow up
3. Use of footnotes under Article 12 when the MRL is set at LOQ

**A.02** Feedback from the section PPP Legislation of this Committee.

**A.03** Specific substances:

1. Glufosinate ammonium
2. Glyphosate
3. *Bacillus thuringiensis*
4. Acetamiprid
5. Thiacloprid
6. Trimethyl-sulfonium (Trimesium) cation
7. Oxamyl
8. Sodium hydrogen carbonate

**A.04** News from and files related to the European Food Safety Authority:

1. Progress under Article 10 of Regulation (EC) No 396/2005
2. Progress under Article 12 of Regulation (EC) No 396/2005
3. Update on Article 43 mandates of Regulation (EC) No 396/2005

4. Other issues

- TDMs

**A.05** Alignment of certain MRLs for pesticides and veterinary medicinal products.

**A.06** Discussion on the inclusion of certain microorganisms into Annex IV.

**A.07** Monitoring of pesticides residues:

- Outcome of the last Working Group of 17 October 2022

**A.08** Screening exercise on temporary MRLs in Regulation (EC) No 396/2005 that expire in 2022-2023:

1. General overview
2. Chlormequat and mepiquat in cultivated fungi

**A.09** International Matters:

1. OECD Guidance document on the definition for risk assessment
2. OECD Honey Guidelines
3. Codex Alimentarius/JMPR issues
  - a) Guidelines for general principles for EU coordinated positions for CCPR
  - b) Issues arising from eWGs
4. Other

**A.10** State of play on Cumulative Risk Assessment (CRA).

**A.11** **Endorsement by the Committee** of the Working Document on pesticides to be considered for inclusion in the national control programmes to ensure compliance with maximum residue levels of pesticides residues in and on food of plant and animal origin (SANCO/12745/2013, Rev. 14).

**A.12** Notifications under Article 18(4) to Regulation (EC) No 396/2005.

**A.13** Designation of Member States for maximum residue levels (MRL) applications.

**A.14** Update of the Technical Guideline on the Evaluation of Extraction Efficiency (SANTE/2017/10632 Rev. 4).

**A.15** Guidance Document on Pesticide Analytical Methods for Risk Assessment and Post-approval Control and Monitoring Purposes (SANTE/2020/12830 Rev. 1).

**A.16** Forthcoming draft Regulations (indicative only):

1. Potassium phosphonates
2. Carbendazim, thiophanate-methyl
3. Deltamethrin, metalaxyl-M, trifloxystrobin

4. Quinoxifen, lufenuron
  5. Fenoxycarb, diethofencarb, pencycuron, flutriafol, myclobutanil
  6. Measure addressing confirmatory data under Article 12
- A.17** Draft revised Communications on data requirements (Commission Regulation (EU) No 283/2013 and 284/2013).
- A.18** Other Information points:
1. Update on PRAC measures/objections
  2. Brexit
  3. Commission Directive 2002/63/EC on sampling
  4. Future organisation of PAFF meetings
  5. Planned working group with Member States on genotoxic carcinogens
  6. Inclusion of mukunuwenna (*Alternanthera sessilis*) in part B of Annex I to Regulation (EC) No 396/2005 – question from a MS

## **Section B      Draft(s) presented for an opinion**

- B.01** Exchange of views and possible opinion of the Committee on a draft Commission Regulation (EU) .../... amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for pyridaben, pyridate, pyriproxyfen and triclopyr in or on certain products (Art. 10).  
(PLAN/2022/2334)

**Legal Basis:** Regulation (EC) No 396/2005 - Article 14(1)(a)

**Procedure:** Regulatory procedure with scrutiny

- B.02** Exchange of views and possible opinion of the Committee on a draft Commission Regulation (EU) .../...amending Annexes II, III and V to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for isoxaben, novaluron and tetraconazole in or on certain products (Art. 12).  
(SANTE/10108/2022; PLAN/2022/1062)

**Legal Basis:** Regulation (EC) No 396/2005 - Articles 14(1)(a), 18(1)(b) and 49(2)

**Procedure:** Regulatory procedure with scrutiny

- B.03** Exchange of views and possible opinion of the Committee on a draft Commission Regulation (EU) .../...amending Annex II to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for cyantraniliprole and folpet in or on certain products (Art. 10).  
(PLAN/2022/1666)

**Legal Basis:** Regulation (EC) No 396/2005 - Article 14(1)(a)

**Procedure:** Regulatory procedure with scrutiny

- B.04** Exchange of views and possible opinion of the Committee on a draft Commission Regulation (EU) .../... amending Annexes II, III and V to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for bromopropylate, chloridazon, fenpropimorph, imazaquin and tralkoxydim in or on certain products.

(SANTE/10644/2021; PLAN/2021/11187)

**Legal Basis:** Regulation (EC) No 396/2005 - Articles 14(1)(a), 18(1)(b) and 49(2)

**Procedure:** Regulatory procedure with scrutiny

- B.05** Exchange of views and possible opinion of the Committee on a draft Commission Regulation (EU) .../... correcting Commission Regulation (EU) 2022/1363 of 3 August 2022 amending Annex II to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for 2,4-D, azoxystrobin, cyhalofop-butyl, cymoxanil, fenhexamid, flazasulfuron, florasulam, fluroxypyr, iprovalicarb and silthiofam in or on certain products.

(PLAN/2022/2331)

**Legal Basis:** Regulation (EC) No 396/2005 - Article 14(1)(a)

**Procedure:** Regulatory procedure with scrutiny

## **Section C      Draft(s) presented for discussion**

- C.01** Exchange of views of the Committee on a draft Commission Implementing Regulation as regards a coordinated multiannual control programme of the Union for 2024, 2025 and 2026 to ensure compliance with maximum residue levels of pesticides and to assess the consumer exposure to pesticide residues in and on food of plant and animal origin.

(PLAN/2022/2309)

**Legal Basis:** Regulation (EC) No 396/2005 - Article 29(2)

**Procedure:** Examination procedure

- C.02** Exchange of views of the Committee on a draft Commission Regulation as regards maximum residue levels for phosmet and pyriproxyfen in or on certain products (Art. 12).

(PLAN/2022/2311)

**Legal Basis:** Regulation (EC) No 396/2005 - Articles 14(1)(a), 18(1)(b) and 49(2)

**Procedure:** Regulatory procedure with scrutiny

- C.03** Exchange of views of the Committee on a draft Commission Regulation as regards maximum residue levels for Denatonium benzoate, diuron, etoxazole, methomyl and teflubenzuron in or on certain products.

(PLAN/2022/2310)

**Legal Basis:** Regulation (EC) No 396/2005 - Articles 14(1)(a), 18(1)(b) and 49(2)

**Procedure:** Regulatory procedure with scrutiny

**C.04** Exchange of views of the Committee on a draft Commission Regulation as regards maximum residue levels for carbetamide, carboxin, fenbuconazole and triflumuron in or on certain products.

(PLAN/2022/2308)

**Legal Basis:** Regulation (EC) No 396/2005 - Articles 14(1)(a), 18(1)(b) and 49(2)

**Procedure:** Regulatory procedure with scrutiny

**C.05** Exchange of views of the Committee on a draft Commission Regulation as regards maximum residue levels for bifentazate in or on certain products.

(PLAN/2022/2307)

**Legal Basis:** Regulation (EC) No 396/2005 - Articles 14(1)(a) and 18(1)(b)

**Procedure:** Regulatory procedure with scrutiny