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Health and Food Safety Directorate General

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**Standing Committee on Plants, Animals, Food and Feed**

**Section *Phytopharmaceuticals – Pesticide Residues***

**10 - 11 May 2023**

**CIRCABC Link:** <https://circabc.europa.eu/ui/group/95a86e0e-0cfe-4354-8d9f-c447c6e85c1b/library/b889e265-ea8b-4682-850d-0d85bf374c35?p=1>

<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. Statement from EFSA for substances for which no Article 12 review is necessary

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

1. General issues
2. Procedural document on setting of Toxicological Reference Values derived via an MRL application or MRL review process (outside an assessment for approval or renewal of an active substance)
3. Specific cases
  - a) Bifenthrin
  - b) Fosetyl-Al, disodium phosphonates and potassium phosphonates

**A.03** Specific substances:

1. Glufosinate ammonium
2. Glyphosate
3. Bacillus thuringiensis
  - a) Tour de table on risk management options
4. Trimethyl-sulfonium (Trimesium) cation
5. Copper
6. Folpet

- 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 Art. 43 mandates of Regulation (EC) No 396/2005
  4. Other issues
    - a) Statement on BAC/DDAC and chlorates on fish
- A.05** Alignment of certain MRLs for pesticides and veterinary medicinal products.
- A.06** Screening exercise on temporary MRLs in Regulation (EC) No 396/2005 that expire in 2023-2024:
1. General overview
  2. Chlormequat and mepiquat in cultivated fungi
- A.07** 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. WTO – Plurilateral Meeting of 17 March 2023
    - a) Wording of transitional measures in our Regulations
- A.08** Update of the Commission Working Document on drafting measures to amend pesticide MRLs following Article 12 of Regulation (EC) No 396/2005 (SANTE 11485/2012) **for endorsement by Member States.**
- A.09** Forthcoming WG on Cumulative Risk Assessment (CRA).
- A.10** Feedback from the WG on Sampling Regulation.
- A.11** State of play on genotoxic carcinogens.
- 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) **for endorsement by Member States.**
- A.15** Forthcoming draft Regulations (indicative only):
1. Dithiocarbamates

- 2. Cypermethrins
  - 3. Dithianon
  - 4. Cyproconazole and spirodiclofen
- A.16** Draft revised Communications on data requirements (Commission Regulation (EU) No 283/2013 and 284/2013).
- A.17** Issues related to Annex 1 of Regulation (EC) No 396/2005.
- A.18** Revision of the technical guidelines on data requirements for setting maximum residue levels, comparability of residue trials, and extrapolation of residue data on products from plant and animal origin (SANTE/2019/12752) **for endorsement by Member States.**
- A.19** Other Information points:
- 1) Update on PRAC measures/objections
  - 2) Brexit
  - 3) Update on F2F -measure lowering MRLs for clothianidin and thiamethoxam
  - 4) Responding to comments from third countries sent to Rapporteur Member States
  - 5) Cyflumetofen - fast track procedure for courgettes and gherkins (BE request)
  - 6) Captan - consideration for a temporary MRL for honey (BE request)

## **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 IV to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for isoxaben, metaldehyde, Metarhizium brunneum strain Ma 43, paclobutrazol and Straight Chain Lepidopteran Pheromones (SCLP) in or on certain products.

(PLAN/2023/950)

**Legal Basis:** Regulation (EC) No 396/2005 - Articles 5 and 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 Annex III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for tricyclazole in or on certain products.

(PLAN/2023/136)

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

**Procedure:** Regulatory procedure with scrutiny

- B.03** 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 pyriproxyfen in or on certain products.

(PLAN/2022/2637)

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

**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 and V to Regulation (EC) No 396/2005 of the European Parliament and of the Council 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

- B.05** Exchange of views and possible opinion of the Committee on a draft Commission Regulation (EU) .../...amending Annexes II and V to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for carbetamide, carboxin, 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

- B.06** Exchange of views and possible opinion of the Committee on a draft Commission Regulation (EU) .../...amending Annexes II and V to Regulation (EC) No 396/2005 of the European Parliament and of the Council 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

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

- C.01** Exchange of views of the Committee on a draft Commission Regulation as regards maximum residue levels for indoxacarb in or on certain products.

(PLAN/2023/242)

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

**Procedure:** Regulatory procedure with scrutiny

- C.02** Exchange of views of the Committee on a draft Commission Regulation as regards maximum residue levels for carbendazim and thiophanate-methyl in or on certain products.

(PLAN/2022/2853)

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

**Procedure:** Regulatory procedure with scrutiny

- C.03** Exchange of views of the Committee on a draft Commission Regulation as regards maximum residue levels for fosetyl-Al, potassium phosphonates and disodium phosphonates in or on certain products.

(PLAN/2023/138)

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

**Procedure:** Regulatory procedure with scrutiny

- C.04** Exchange of views of the Committee on a draft Commission Regulation as regards maximum residue levels for diethofencarb, fenoxycarb, flutriafol, myclobutanil and pencycuron in or on certain products.

(PLAN/2023/194)

**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 (Z)-13-hexadecen-11yn-1-ylacetate and (Z,Z,Z,Z)-7,13,16,19-docosatetraen-1-ylisobutyrate, acrinathrin, azimsulfuron, famoxadone, methyl-nonylketone, prochloraz, sodium hypochlorite in or on certain products.

(PLAN/2023/145)

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

**Procedure:** Regulatory procedure with scrutiny

- C.06** Exchange of views of the Committee on a draft Commission Regulation as regards maximum residue levels for 2,4 DB, methoxyfenozide, iodosulfuron-methyl, mesotrione and pyraflufen-ethyl in or on certain products.

(PLAN/2022/2563)

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

**Procedure:** Regulatory procedure with scrutiny

- C.07** Exchange of views of the Committee on a draft Commission Regulation as regards maximum residue levels for deltamethrin, metalaxyl-M, thiabendazole and trifloxystrobin.

(PLAN/ 2023/326)

**Legal Basis:** Regulation (EC) No 396/2005 - Articles 14(1)(a) and 49(2), Regulation (EC) No 178/2002 - Articles 5(3) and 13(e)

**Procedure:** Regulatory procedure with scrutiny

**C.08** Exchange of views of the Committee on a draft Commission Regulation as regards maximum residue levels for bifenthrin.

(PLAN/ 2023/951)

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

**Procedure:** Regulatory procedure with scrutiny

**C.09** Exchange of views of the Committee on a draft Commission Regulation as regards maximum residue levels for haloxypop.

(PLAN/2023/897)

**Legal Basis:** Regulation (EC) No 396/2005 - Article 14(1)(a) and Article 49(2)

**Procedure:** Regulatory procedure with scrutiny

**C.10** Exchange of views of the Committee on a draft Commission Regulation as regards maximum residue levels for oxamyl in or on certain products.

(PLAN/2023/947)

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

**Procedure:** Regulatory procedure with scrutiny

**C.11** Exchange of views of the Committee on a draft Commission Regulation as regards maximum residue levels for desmedipham, etridiazole, flurtamone, profoxydim, difenacoum, and potassium permanganate in or on certain products.

(PLAN/2023/946)

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

**Procedure:** Regulatory procedure with scrutiny

**C.12** Exchange of views of the Committee on a draft Commission Regulation as regards maximum residue levels for thiacloprid.

(PLAN/ 2023/961)

**Legal Basis:** Regulation (EC) No 396/2005 - Article 14(1)(a) and possibly Article 49(2)

**Procedure:** Regulatory procedure with scrutiny

**C.13** Exchange of views of the Committee on a draft Commission Regulation as regards maximum residue levels for mandipropamid in or on certain products.

(PLAN/2023/750)

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

**Procedure:** Regulatory procedure with scrutiny

**C.14** Exchange of views of the Committee on a draft Commission Regulation as regards maximum residue levels for fipronil in or on certain products.

(PLAN/2023/962)

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

**Procedure:** Regulatory procedure with scrutiny

- C.15** Exchange of views of the Committee on a draft Commission Regulation as regards maximum residue levels for bispyribac, lemon essential oil, metosulam, oryzalin, oxasulfuron and triazoxide in or on certain products.

(PLAN/2023/948)

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

**Procedure:** Regulatory procedure with scrutiny