



EUROPEAN COMMISSION

Health and Food Safety Directorate General

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Standing Committee on Plants, Animals, Food and Feed
Section *Phytopharmaceuticals – Pesticide Residues*
22 - 23 April 2024

CIRCABC Link: <https://circabc.europa.eu/ui/group/95a86e0e-0cfe-4354-8d9f-c447c6e85c1b/library/09141bfb-f622-4fb9-9fb9-57d378c69401?p=1>

AGENDA

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
2. Non-approved substances for follow up
 - a) Update
 - b) Next mandate to EFSA

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

1. General issues
2. Endorsed TRV for cypermethrin

A.03 Specific substances:

1. Difenoconazole
2. Sodium silver thiosulfate
3. Straight Chain Lepidopteran Pheromones
4. Imazalil
5. *Bacillus thuringiensis* (Bt)
6. Bifenazate
7. Etoxazole
8. Glufosinate
9. Matrine

10. Ethiprole
11. Nicotine
12. Dimethomorph

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: Presentation of the 2022 Annual Report on Pesticides Residues

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

A.07 International Matters:

1. OECD Guidance document on the definition for risk assessment
2. OECD Honey Guidelines
3. OECD Guidance on Stability of Pesticide Residues in Stored Commodities
4. OECD Guidance Document on Pesticide Residue Analytical Methods
5. Codex Alimentarius/JMPR issues
 - a) Update on EFSA mandates
 - b) Substances for CCPR 55
 - c) Updates from working groups

A.08 Cumulative Risk Assessment (CRA).

A.09 Sampling Regulation – Feedback from WG.

A.10 Notifications under Article 18(4) to Regulation (EC) No 396/2005.

A.11 Designation of Member States for maximum residue levels (MRL) applications.

A.12 Forthcoming draft Regulations (indicative only):

1. Gamma-cyhalothrin Article 12

A.13 Issues related to Annex IV to Regulation (EC) No 396/2005.

A.14 Issues related to Annex I to Regulation (EC) No 396/2005.

A.15 New proposals for Table 3 of the extrapolation guidelines (SANTE/2019/12752 Rev01).

A.16 EFSA Guidelines on rotational crops – **for endorsement by Member States.**

A.17 Other Information points:

- 1) Article 19 of Regulation (EC) No 395/2005 translations
- 2) Chlorate - question from a Member State
- 3) Piperonylbutoxide - question from a Member State
- 4) Presence of chlormequat in sunflower
- 5) Member States' submission of the multiannual national control programmes for pesticides residues

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, III and IV to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for chitosan, clopyralid, difenoconazole, fat distillation residues, flonicamid, hydrolysed proteins, and lavandulyl senecioate in or on certain products**

(PLAN/2024/791)

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 Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council 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), Regulation (EC) No 178/2002 - Articles 5(3) and 13 (e)

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, III and V to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for benomyl, 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

B.04 Exchange of views and possible opinion of the Committee on a draft Commission Regulation (EU) .../... amending Annex I to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards as regards radish leaves

(PLAN/2023/2900)

Legal Basis: Regulation (EC) No 396/2005 - Article 4

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 III to Regulation (EC) No 396/2005 as regards maximum residue levels for cyproconazole and spirodiclofen in or on certain products

(PLAN/2023/1960)

Legal Basis: Regulation (EC) No 396/2005 - Articles 14(1)(a) 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 Annex II to Regulation (EC) No 396/2005 as regards maximum residue levels napropamide, pyridaben and tebufenpyrad in or on certain products

(PLAN/2023/2190)

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

Procedure: Regulatory procedure with scrutiny

B.07 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 azoxystrobin, famoxadone, flutriafol, mandipropamid and mefentrifluconazole in or on certain products

(PLAN/2024/817)

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

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 cypermethrins in or on certain products

(PLAN/2023/1863)

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

Procedure: Regulatory procedure with scrutiny

- C.02 Exchange of views of the Committee on a draft Commission Regulation as regards maximum residue levels for dithiocarbamates in or on certain products**
(PLAN/2023/2019)
Legal Basis: Regulation (EC) No 396/2005 - Article 14(1)(a) 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 fenbuconazole and penconazole in or on certain products**
(PLAN/2024/23)
Legal Basis: Regulation (EC) No 396/2005 - Articles 14(1)(a) 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 isopyrazam in or on certain products**
(PLAN/2022/2927)
Legal Basis: Regulation (EC) No 396/2005 - Articles 14(1)(a) 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 thiacloprid in or on certain products**
(PLAN/2023/961)
Legal Basis: Regulation (EC) No 396/2005 - Articles 14(1)(a), 18(1) and 49(2)
Procedure: Regulatory procedure with scrutiny
- C.06 Exchange of views of the Committee on a draft Commission Implementing Regulation as regards methods of sampling and analysis for the control of pesticide residues in and on products of plant origin and repealing Directive 2002/63/EC**
(PLAN/2023/636)
Legal Basis: Regulation (EU) 2017/625 - Article 34(6)
Procedure: Examination procedure
- C.07 Exchange of views of the Committee on a draft Commission Regulation as regards maximum residue levels for zoxamide in or on certain products**
(PLAN/2024/307)
Legal Basis: Regulation (EC) No 396/2005 - Articles 14(1)(a) and 49(2)
Procedure: Regulatory procedure with scrutiny