



EUROPEAN COMMISSION

Health and Food Safety Directorate General

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Standing Committee on Plants, Animals, Food and Feed
Section *Phytopharmaceuticals - Residues*
18 - 19 May 2026

CIRCABC Link: https://circabc.europa.eu/ui/group/95a86e0e-0cfe-4354-8d9f-c447c6e85c1b/library/e56b3d83-1570-430a-821d-5bf90362f6e?p=1&n=-1&sort=name_ASC

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) Call for data for non-approved active substances (2nd batch)

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

1. General issues

A.03 Specific substances:

1. 1,2-dichloroethanol
2. Difluoroacetic acid
3. Chlorate
4. Guazatine
5. Choline hydrogen phosphonate
6. Trimethyl-sulfonium cation
7. Pyrimethanil

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.05 Presentation of the 2024 Annual Report on Pesticides Residues
- A.06 Alignment of certain MRLs for multiple-use substances
- A.07 Screening exercise on temporary MRLs in Regulation (EC) No 396/2005:
 - 1. General overview
- A.08 International Matters (MG):
 - 1. OECD Guidance document on the residue definition for risk assessment
 - 2. OECD Honey Guidelines
 - 3. OECD Guidance Document on Pesticide Residue Analytical Methods
 - 4. OECD Test Guideline 508 (Magnitude of the Pesticide Residues in Processed Commodities, adopted 3 October 2008) and OECD Guidance Document on Magnitude of Pesticide Residues in Processed Commodities; adopted 29 July 2008, Series of Testing and Assessment No 96
 - 5. Codex Alimentarius/JMPR issues
- A.09 Cumulative Risk Assessment (CRA)
- A.10 Other approaches for risk assessment (IESTI, PRIMO):
 - 1. Update on IESTI
 - 2. Risk management questions on PRIMO 4 and implementation plan - **for endorsement by Member States**
- A.11 Notifications under Article 18(4) to Reg. (EC) No 396/2005.
- A.12 Designation of Member States for MRL applications
- A.13 Forthcoming draft Regulations (indicative only):
 - 1. Acetamiprid
 - 2. Ametoctradin, lambda-cyhalothrin, prothioconazole
 - 3. Cinmethylin and metyltetraprole
 - 4. Dicloran, fenazaquin, fluopyram and tebufenozide
- A.14 MRLs for milk and for infant formula and follow-on formula
- A.15 Issues related to Annex I to Regulation (EC) No 396/2005
- A.16 Food and Feed Safety omnibus
- A.17 Commodities with no consumption data in PRIMo - **for endorsement by Member States**
- A.18 Sampling Regulation follow-up matters

A.19 Other Information points:

1. Feedback on the vote conducted via written procedure launched on 9 April 2026 on the draft Commission Regulation (PLAN/2024/2763, revision 6) concerning the MRLs for benomyl, carbendazim, and thiophanate-methyl
2. EU Pesticide Database Data cleaning
3. Corrigendum to the Maltese language version of Commission Regulation (EU) 2026/840 of 15 April 2026 setting MRLs for copper compounds

A.20 Poland national interim emergency measure to ban the placing on the market of products treated with certain non approved active substances (Article 54 of Regulation (EC) No 178/2002)

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 Annex II to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for acequinocyl, deltamethrin, dodine, maleic hydrazide, pinoxaden and prothioconazole in or on certain products

(PLAN/2025/3053)

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 Annex II to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for ametoctradin, cyflumetofen, diflufenican and milbemectin in or on certain products

(PLAN/2026/701)

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 Annex II to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for azoxystrobin, etofenprox, fenpropidin, flupyradifurone, hexythiazox, imazalil, spinosad and tebufenozide in or on certain products

(PLAN/2025/2744)

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

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 III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for difenoconazole in or on certain products

(PLAN/2024/2476)

Legal Basis: Regulation (EC) No 396/2005 - Articles 14(1)(a) 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 III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for alpha-cypermethrin and cypermethrin 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

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

C.01 Exchange of views 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 azocyclotin, chlorfenapyr, cyhexatin, dicofol, endosulfan, fenarimol, fenpropathrin and profenofos in or on certain products

(PLAN/2025/1425)

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

Procedure: Regulatory procedure with scrutiny

C.02 Exchange of views 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 carbofuran, imazalil, mandipropamid, propaquizafop, quizalofop-P-ethyl, and quizalofop-P-tefuryl in or on certain products

(PLAN/2025/2832)

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 (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 diazinon in or on certain products

(PLAN/2025/3159)

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 (EU) amending Annex II to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for bixlozone in or on certain products

(PLAN/2026/987)

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

Procedure: Regulatory procedure with scrutiny