EUROPEAN COMMISSION

Health and Food Safety Directorate General

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Standing Committee on Plants, Animals, Food and Feed
Section *Phytopharmaceuticals - Residues*
5 - 6 February 2026

CIRCABC Link: *https://circabc.europa.eu/ui/group/95a86e0e-0cfe-4354-8d9f-c447c6e85c1b/library/fa620bb0-c1d3-4f95-b9c0-bab460d7f76e?p=1*

AGENDA

Section A Information and/or discussion

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

1. Priority list
2. Confirmatory data Art. 12 follow-up
   a) Cases where EFSA RO has been published
3. Non-approved substances for follow-up

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

1. General issues

A.03 Specific substances:

1. Chlorate
2. Maleic hydrazide
3. Diquat
4. Copper

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 Alignment of certain MRLs for multiple-use substances.

A.06 Screening exercise on temporary MRLs in Regulation (EC) No 396/2005:

1. General overview

A.07 International Matters:

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)
5. OECD Guidance Document on Magnitude of Pesticide Residues in Processed Commodities; adopted 29 July 2008, Series of Testing and Assessment No 96)
6. Codex Alimentarius/JMPR issues

A.08 Cumulative Risk Assessment (CRA).

A.09 Other approaches for risk assessment (IESTI, PRIMO).

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

A.11 Designation of Member States for MRL applications.

A.12 Forthcoming draft Regulations (indicative only):

1. Bixlozone and pydiflumetofen
2. Fluopyram, fenazaquin, dicloran and tebufenozide
3. Candidate substances for Annex IV inclusion
4. Correcting act EUMACP Regulation

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

A.14 Food and Feed Safety omnibus.

A.15 Other Information points:

1. Feedback on RASFF working instruction
2. Question from a Member States on commodities with no consumption data in PRIMo
3. Conclusion of the European Court of Justice on the Case T-247/23 on MRLs for clothianidin and thiamethoxam
4. Objection of European Parliament on the MRL draft Regulation for acetamiprid

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 product

(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 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.03 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Regulation as regards a coordinated multiannual control programme of the Union for 2027, 2028 and 2029 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/2025/1929)

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

Procedure: Examination procedure

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 1,4-dimethylnaphthalene, chlormequat, metribuzin, metribuzin-desamino-diketo (metribuzin-DADK), terbuthylazine and triclopyr in or on certain products

(PLAN/2025/1086)

Legal Basis: Regulation (EC) No 396/2005 - Articles 14(1)(a), 18(1) 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 Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for fenpyrazamine in or on certain products

(PLAN/2025/2793)

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

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 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

C.02 Exchange of views 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 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.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 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.04 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.05 Exchange of views of the Committee on a draft Commission Regulation (EU) as regards maximum residue levels for benomyl, carbendazim and thiophanate-methyl in or on certain products

(PLAN/2024/2763)

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

Procedure: Regulatory procedure with scrutiny

C.06 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