



EUROPEAN COMMISSION

Health and Food Safety Directorate General

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Standing Committee on Plants, Animals, Food and Feed

Section *Phytopharmaceuticals - Residues*

28 - 29 September 2026



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
3. Priority list

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

1. General issues

A.03 Specific substances

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 Monitoring Working Group

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:

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 Regulation (EC) No 396/2005

A.12 Designation of Member States for MRL applications

A.13 Forthcoming draft Regulations (indicative only):

1. Clofentezine
2. Clopyralid
3. Difluoroacetic acid and flupyradifurone

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:

1. State of play in Council and Parliament
2. Presentation of the JRC Study: “Assessing the economic impacts of stronger alignment of production standards applied to imported products on the EU’s agricultural sector”

A.17 Other Information points:

1. Corrigenda to the Maltese and Spanish language version of Commission Regulation (EU) 2026/840 of 15 April 2026 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 copper compounds in or on certain products and Commission Regulation (EU) 2018/62 of 17 January 2018 replacing Annex I to Regulation (EC) No 396/2005 of the European Parliament and of the Council
2. Reporting issues related to the pesticides monitoring data

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 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.02** Exchange of views and possible opinion of the Committee on a draft Commission Regulation (EU) amending Annex II and Annex III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for cycloxydim, ethofumesate, folpet, mepiquat and quinmerac in or on certain products
(PLAN/2026/1732)

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, 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

- 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 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

- B.05** 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 acetamiprid in or on certain products
(PLAN/2026/625)

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

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 III 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

Section C Draft(s) presented for discussion

- C.01** 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

- C.02** Exchange of views 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 dicloran, fenazaquin, fluopyram and tebufenozide in or on certain products

(PLAN/2026/1190)

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 Annex II to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for ametoctradin, isoxaben, lambda-cyhalothrin and prothioconazole in or on certain products

(PLAN/2026/1652)

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

Procedure: Regulatory procedure with scrutiny