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Health and Food Safety Directorate General

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Standing Committee on Plants, Animals, Food and Feed

Section *Phytopharmaceuticals - Residues*

23 - 24 November 2026

[redacted]

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.02 Feedback from the section PPP Legislation of this Committee

1. General issues

A.03 Specific substances

1. Benomyl
2. Pyraclostrobin
3. Trimethyl-sulfonium cation

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

1. Feedback from the Working Group
2. Guidance document on Analytical Quality Control and Method Validation Procedures for pesticide residues analysis in food and feed (SANTE/11312/2021, v2027) **for endorsement by the Member States**
3. Working Document on pesticides to be considered for inclusion in the national control programmes to ensure compliance with maximum residue levels of pesticide residues in and on food of plant and animal origin (SANCO/12745/2013, Rev.18) **for endorsement by the Member States**

A.06 Alignment of certain MRLs for multiple-use substances

A.07 Screening exercise on temporary MRLs in Regulation (EC) No 396/2005

A.08 International Matters:

1. OECD Guidance document on the residue definition for risk assessment
2. OECD Honey Guidelines
3. 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)
4. Codex Alimentarius/JMPR issues

A.09 Cumulative Risk Assessment (CRA)

A.10 Other approaches for risk assessment

1. Update on IESTI
2. PRIMO 4

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)

- Possible corrigendum to Commission Implementing Regulation (EU) 2026/765
- Flufenacet and mepanipyrim

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 Simplification Package

A.17 Other Information points

1. CIRCABC memberships (PAFF Legislation and Residues) and update of Member States contact lists
2. Updated Standard Rules of Procedure

Section B **<u>Draft(s) presented for an opinion</u>**

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 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

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 acequinocyl, aclonifen, dodine, potassium phosphonates and propyzamide in or on certain products

(PLAN/2026/2405)

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, hexythiazox and spinosad

(PLAN/2026/2454)

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

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 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.02 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

C.03 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 clopyralid in or on certain products

(PLAN/2026/2245)

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 Implementing Regulation as regards a coordinated multiannual control programme of the Union for 2028, 2029 and 2030 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/2026/1730)

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

Procedure: Examination procedure

C.05 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 difluoroacetic acid and flupyradifurone in or on certain products

(PLAN/2026/2039)

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

Procedure: Regulatory procedure with scrutiny