EUROPEAN COMMISSION

Health and Food Safety Directorate General

sante.g.3(2023)9016619

Standing Committee on Plants, Animals, Food and Feed
Section *Phytopharmaceuticals – Pesticide Residues*
18 - 19 September 2023

CIRCABC Link: *https://circabc.europa.eu/ui/group/95a86e0e-0cfe-4354-8d9f-c447c6e85c1b/library/78eb369a-150b-46bc-ab1f-2b7eb013a882?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
   b) Missing analytical standards follow up
   c) Commission Staff Working Document on the Evaluation of data submitted to confirm MRLs following their review in accordance with Article 12 of Regulation (EC) No 396/2005 and risk management decisions in the absence of such data (SANTE/10235/2016, Rev. 5.0)
3. Non-approved substances for follow up
   a) Next mandate to EFSA
   b) Procedural issues

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

1. General issues

A.03 Specific substances:

1. Glyphosate
2. Bacillus thuringiensis
3. Copper
4. Folpet
5. Acetamiprid

6. Captan
7. Ethephon
8. Bifenazate
9. Flupyradifurone/DFA

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
   - Zoxamide
   - Clopyralid
3. Update on Art. 43 mandates of Regulation (EC) No 396/2005
4. Other issues

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. Codex Alimentarius/JMPR issues
   a) Guidelines for general principles for EU coordinated positions for CCPR **for endorsement by Member States**
   b) Follow up for CXLs for which reservations were made in earlier years and proposed amendments of the Terms of reference for the next EFSA scientific report
   c) Feedback from CCPR meeting and preparations for CAC

A.08 Cumulative Risk Assessment (CRA):

1. Follow up on CRA EFSA/SANTE Action Plan
2. Feedback from RIVM training/ WG on risk management issues
3. Working document on risk management aspects related to the assessment of cumulative exposure (SANTE/10216/2015, Rev. 8) for endorsement by Member States
4. Training needs of MS and views on implementation

A.09 Sampling Regulation – Feedback from WG and next steps.

A.10 State of play on genotoxic carcinogens and Guidelines for harmonised risk management approaches and enforcement action in cases of incidents involving food products containing genotoxic carcinogens **(for endorsement by the Member States**).

A.11 Notifications under Article 18(4) to Regulation (EC) No 396/2005.

A.12 Designation of Member States for maximum residue levels (MRL) applications.

A.13 Monitoring of pesticide residues:

1. Next Working Group
2. Measurement uncertainty for laboratories that are not designated for official controls in the sense of Regulation (EU) 2017/625
3. Enforcement of residue definitions that include conjugates of substances

A.14 Forthcoming draft Regulations (indicative only):

1. Dithiocarbamates
2. Fenarimol and fenpropathrin
3. Fluopyram, napropamide, pyridaben and tebufenpyrad

A.15 Issues related to Annex 1 of Regulation (EC) No 396/2005.

A.16 Regular review of designations of EU Reference Laboratories.

A.17 Data extraction from TRACES for the purpose of EFSA's reporting.

A.18 Wording of transitional measures in our Regulations.

A.19 Other Information points:

1. Update on PRAC measures/objections and update on tricyclazole
2. Brexit
3. Draft revised Communications on data requirements (Commission Regulation (EU) No 283/2013 and 284/2013)
4. Update on F2F -measure lowering MRLs for clothianidin and thiamethoxam
5. Furathiocarb (Question from BE)
6. Future organisation of PAFF meetings
7. MRLs for chili peppers

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

(PLAN/2023/947)

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

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, III and V to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for desmedipham, etridiazole, flurtamone, profoxydim, difenacoum, and potassium permanganate in or on certain products.

(PLAN/2023/946)

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

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 indoxacarb in or on certain products.

(PLAN/2023/242)

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

(PLAN/2023/1703)

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

Procedure: Regulatory procedure with scrutiny

B.05 Exchange of views and possible opinion of the Committee on a draft Commission Regulation (EU) …/…amending and correcting Annex II to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for mandipropamid in or on certain products.

(PLAN/2023/750)

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

(PLAN/2023/962)

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

Procedure: Regulatory procedure with scrutiny

B.07 Exchange of views and possible opinion 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 diethofencarb, fenoxycarb, flutriafol and pencycuron in or on certain products.

(PLAN/2023/194)

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

Procedure: Regulatory procedure with scrutiny

B.08 Exchange of views and possible opinion 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 (Z)-13-hexadecen-11-yn-1-yl acetate, (Z,Z,Z,Z)-7,13,16,19-docosatetraen-1-yl isobutyrate, acrinathrin, azimsulfuron, famoxadone, prochloraz and sodium hypochlorite in or on certain products.

(PLAN/2023/145)

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

Procedure: Regulatory procedure with scrutiny

B.09 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 haloxyfop in or on certain products.

(PLAN/2023/897)

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

Procedure: Regulatory procedure with scrutiny

B.10 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 thiacloprid in or on certain products.

(PLAN/ 2023/961)

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

Procedure: Regulatory procedure with scrutiny

B.11 Exchange of views and possible opinion of the Committee on a draft Commission Regulation (EU) …/… amending Annex IV to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards *Pythium oligandrum* strain M1, *Trichoderma atroviride* strain AGR2 and *Trichoderma atroviride* strain AT10.

(PLAN/2023/1697)

Legal Basis: Regulation (EC) No 396/2005 - Articles 5(1) and 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 as regards maximum residue levels for glufosinate in or on certain products.

(PLAN/2023/1772)

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 as regards maximum residue levels for cypermethrins in or on certain products.

(PLAN/2023/1863)

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

Procedure: Regulatory procedure with scrutiny

C.03 Exchange of views of the Committee on a draft Commission Regulation as regards maximum residue levels for dithianon in or on certain products.

(PLAN/2023/1782)

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

Procedure: Regulatory procedure with scrutiny

C.04 Exchange of views of the Committee on a draft Commission Regulation as regards maximum residue levels for 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

C.05 Exchange of views of the Committee on a draft Commission Regulation 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)

Procedure: Regulatory procedure with scrutiny

C.06 Exchange of views of the Committee on a draft Commission Regulation as regards maximum residue levels for deltamethrin, metalaxyl-M, thiabendazole and trifloxystrobin.

(PLAN/ 2023/326)

Legal Basis: Regulation (EC) No 396/2005 - Articles 14(1)(a) and 49(2), Regulation (EC) No 178/2002 - Articles 5(3) and 13(e)

Procedure: Regulatory procedure with scrutiny

C.07 Exchange of views of the Committee on a draft Commission Regulation as regards maximum residue levels for bifenthrin.

(PLAN/ 2023/951)

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

Procedure: Regulatory procedure with scrutiny

C.08 Exchange of views of the Committee on a draft Commission Regulation as regards maximum residue levels for 2,4-DB, iodosulfuron-methyl, mesotrione and pyraflufen-ethyl in or on certain products.

(PLAN/2022/2563)

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

Procedure: Regulatory procedure with scrutiny

C.09 Exchange of views of the Committee on a draft Commission Regulation as regards maximum residue levels for cyproconazole, isopyrazam 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

C.10 Exchange of views of the Committee on a draft Commission Regulation as regards maximum residue levels for bispyribac, lemon essential oil, metosulam, oryzalin, oxasulfuron and triazoxide in or on certain products.

(PLAN/2023/948)

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

Procedure: Regulatory procedure with scrutiny

C.11 Exchange of views of the Committee on a draft Commission Regulation as regards maximum residue levels for nicotine in or on certain products.

(PLAN/2023/1999)

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

Procedure: Regulatory procedure with scrutiny