EUROPEAN COMMISSION

Health and Food Safety Directorate General

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Standing Committee on Plants, Animals, Food and Feed
Section *Phytopharmaceuticals – Pesticide Residues*
23 - 24 September 2021

CIRCABC Link: *https://circabc.europa.eu/w/browse/1747ac72-5969-4320-9375-b105dd48f8dd*

AGENDA

Section A Information and/or discussion

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

1. Priorities under Art. 12 – updated table
2. Confirmatory data Art. 12 follow-up
3. Residue definition for risk assessment
4. List of non-approved substances for follow up

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

A.03 Specific substances:

1. Glufosinate ammonium
2. Glyphosate
3. Ethylene oxid – update on the state of play
4. Bacillus thuringiensis
5. Cyantraniliprole
6. Clethodim

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
   - Article 12 Work programme
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 Multiple source substances for which Annex IV inclusion is not recommended.

A.07 Next steps for cumulative risk assessment.

A.08 Screening exercise on temporary MRLs in Regulation (EC) No 396/2005 that will expire in 2021-2022**:**

1. General overview
2. Data analyses for decisions on t-MRLs for chlormequat, mepiquat, profenophos and nicotine

A.09 International Matters:

1. OECD Guidance document on the definition for risk assessment
2. OECD Honey Guidelines and MS experiences with the EU guidelines
3. Codex Alimentarius/JMPR issues

A.10 Information Note on Article 20 of Regulation (EC) No 396/2005 as regards processing factors and composite food and feed (SANTE/10704/2021).

A.11 **Note Taking** of the revised Technical Guidelines for MRL setting as regards the clarification of "Exceptional circumstances" under Article 16 of Regulation (EC) No 396/2005 (SANTE/2015/10595 Rev. 6).

A.12 Notifications under Article 18(4) to Regulation (EC) No 396/2005.

A.13 Designation of Member States for maximum residue levels (MRL) applications.

A.14 Questions related to the implementation of the Extraction Efficiency guidelines (SANTE/2017/10632 Rev. 3).

A.15 Classification issues related to Annex I of Regulation (EC) No 396/2005.

A.16 Update on Farm to Fork/REFIT actions.

A.17 Info on a Corrigendum to Commission Regulation (EU) 2021/1110 of 6 July 2021 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 ametoctradin, bixafen, fenazaquin, spinetoram, tefluthrin and thiencarbazone-methyl in or on certain products (SANTE/10946/2021**).**

A.18 Info on a Draft Commission Delegated Regulation (EU) …/... supplementing Regulation (EU) 2017/625 as regards additional requirements related to the use of relevant substances in food-producing animals and residues arising therefrom, for the entry into the Union of such animals, products of animal origin and composite products.

A.19 Other Information points***:***

1. Update on PRAC measures/objections
2. Brexit
3. Peeling factor/consumption of unpeeled food

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, III and IV to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for acrinathrin, fluvalinate, folpet, fosetyl, isofetamid, *'Pepino Mosaic Virus, EU strain, mild isolate Abp1', 'Pepino Mosaic Virus, CH2 strain, mild isolate Abp2'*, spinetoram and spirotetramat in or on certain products (Art. 10).

(SANTE/10884/2021)

Legal Basis: Regulation (EC) No 396/2005 - Articles 5(1) and 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 flonicamid in or on certain products (Art. 10)**.**

(SANTE/10892/2021)

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 IV to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for acequinocyl, acibenzolar-S-methyl, Bacillus subtilis strain IAB/BS03, emamectin and flutolanil in or on certain products (Art. 10).

(SANTE/11822/2019)

Legal Basis: Regulation (EC) No 396/2005 - Articles 5(1) and 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 Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for dazomet, hexythiazox, metam and methylisothiocyanate in or on certain products (Art. 12).

(SANTE/10942/2021)

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 as regards maximum residue levels for 2,4-D, azoxystrobin, cyhalofop-butyl, cymoxanil, fenhexamid, flazasulfuron, florasulam, fluroxypyr, iprovalicarb, and silthiofam following the evaluation of Article 12 confirmatory data.

(SANTE/12078/2020)

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 as regards maximum residue levels for fluoride, oxyfluorfen, pyroxsulam, quinmerac and sulfuryl fluoride in or on certain products.

(SANTE/10218/2021)

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 as regards maximum residue levels for methoxyfenozide, propoxur, spinosad and thiram in or on certain products.

(SANTE/10552/2021)

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 as regards maximum residue levels for bifenthrin, bromopropylate, chloridazone, imazaquin, fenpropimorph and tralkoxydim in or on certain products.

(SANTE/10644/2021)

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 as regards maximum residue levels for 1,4-dimethylnaphthalene, 8-hydroxyquinoline, pinoxaden and valifenalate in or on certain products (Art. 12).

(SANTE/10776/2021)

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

Procedure: Regulatory procedure with scrutiny