



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 oxide – 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 thiencarbazon-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, *Bacillus subtilis* strain IAB/BS03, emamectin, flutolanil and imazamox in or on certain products

(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