EUROPEAN COMMISSION

Health and Food Safety Directorate General

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Standing Committee on Plants, Animals, Food and Feed
Section *Phytopharmaceuticals - Legislation*
10 - 11 July 2024

CIRCABC Link: *https://circabc.europa.eu/ui/group/95a86e0e-0cfe-4354-8d9f-c447c6e85c1b/library/aaf17dac-e105-4f96-9788-a4e9d88fd935?p=1*

AGENDA

Section A Information and/or discussion

A.01 Summary Report of previous meetings.

A.02 Applications and withdrawals, in particular basic substances:

1. Withdrawal of an application for extension of use of sodium chloride

A.03 General issues on regulatory processes, in particular:

1. expected delivery dates for DAR/RAR
2. MS experiences and practices (updates and survey)
3. PIMS database: information on authorisation of PPPs

A.04 Exchange of views on EFSA conclusions/EFSA scientific reports:

- New active substances / Amendment of conditions of approval
- Renewal of approval
  1. Mecoprop-P
  2. Lenacil
- Basic substances

A.05 Draft Review/Renewal Reports for discussion:

- New active substances / Amendment of conditions of approval
  1. Pydiflumetofen
  2. Clove oil

- Renewal of approval
    3. Milbemectin
    4. Pelargonic acid
    5. Rape seed oil
    6. Flutolanil
    7. Sulfur
    8. Aluminium silicate calcinated
    9. 8-hydroxyquinoline (quinolin-8-ol)
- Basic substances

A.06 Confirmatory Information:

1. Pendimethalin

A.07 Guidance Documents, in particular:

1. EFSA Guidance Risk assessment for Birds and Mammals (for endorsement)
2. Memorandum accompanying the compendium of conditions of use to reduce exposure and risk from plant protection products
3. Guidance on emergency authorisations according to Article 53 of Regulation (EC) No 1107/2009 – draft amendment
4. Technical guidance on the assessment of negligible exposure to an active substance, safener or synergist in a plant protection product under realistic conditions of use
5. Guidance on the assessment of pesticide residues in rotational crops
6. EFSA Guidance Document on the risk assessment of plant protection products on bees (Apis mellifera, Bombus spp. and solitary bees)
7. EFSA guidance document for predicting environmental concentrations of active substances of plant protection products and transformation products of these active substances in soil
8. FOCUS surface water scenarios (ongoing mandate EFSA)
9. Statement of the Scientific Panel on Plant Protection Products and their Residues (PPR Panel) on the design and conduct of groundwater monitoring studies supporting groundwater exposure assessments of pesticides (for info)

A.08 Notifications under Regulation (EC) No 1107/2009 (for information):

1. Article 44(4)
2. Article 36(3)
3. Article 53

A.09 Microorganism and low risk Active Substances.

A.10 Updates, clarifications & questions on specific active substances:

1. Acetamiprid (amended renewal report to be endorsed)
2. Sodium hydrogen carbonate
3. Common metabolites of pyrethroids
4. Trifluoroacetic acid (TFA)
5. Thifensulfuron-methyl
6. SDHI fungicides
7. Talc
8. Labelling of mixed sodium nitro compounds

A.11 Article 21:

1. Flupyradifurone
2. Tea tree oil

A.12 General issues for information / discussion:

1. Scope of Regulation (EC) No 1107/2009:
2. Basic substances – general issues
3. PFAS
4. Cut flowers

A.13 Amendments to Regulations (EU) No 546/2011, (EU) No 283/2013 and (EU) No 284/2013.

A.14 Co-formulants and assessment of formulations, in particular:

1. Implementation of Regulation (EU) 2023/574
2. Ongoing actions

A.15 Report from Working Groups, in particular:

1. Working Group Post Approval Issues (PAI)
2. Working Group on Biopesticides
3. Working Group on comparative assessment
4. Working Group on Negligible Exposure
5. Working Group on environmental relevant topics in the context of Regulation (EC) No 1107/2009

A.16 News and updates, in particular from:

1. European Food Safety Authority (EFSA)
2. Sustainable Use Directive (Directive 2009/128/EC)
3. Health and Food Audits and Analysis (SANTE, Directorate F)

4. Minor Use Facility (MUCF)
5. OECD, FAO and EPPO activities

A.17 Court cases, requests for internal review, Ombudsman cases.

A.18 Exchange of information from the Pesticide Residues section of the Committee.

A.19 Scientific publications and information submitted by stakeholders.

A.20 Date of next meeting(s).

A.21 AoB.

Section B Draft(s) presented for an opinion

B.01 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Regulation (EU) renewing the approval of the active substance captan in accordance with Regulation (EC) No 1107/2009 of the European Parliament and of the Council concerning the placing of plant protection products on the market, and amending Commission Implementing Regulation (EU) No 540/2011 (Draft Review Report SANTE/12270/2020)

(SANTE/12268/2020)

Legal Basis: Regulation (EC) No 1107/2009 - Article 20(1)

Procedure: Examination procedure

B.02 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Regulation (EU) approving the basic substance *Vitis vinifera* L. seed extract (grape seed extract) in accordance with Regulation (EC) No 1107/2009 of the European Parliament and of the Council, and amending Commission Implementing Regulation (EU) No 540/2011 (Draft Renewal Report PLAN/2024/800 RR Rev1)

(PLAN/2024/800)

Legal Basis: Regulation (EC) No 1107/2009 - Article 23(5) in conjunction with Article 13(2)

Procedure: Examination procedure

B.03 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Regulation (EU) concerning the non-approval of eggshell powder as a basic substance in accordance with Regulation (EC) No 1107/2009 of the European Parliament and of the Council concerning the placing of plant protection products on the market (Draft Renewal Report PLAN/2024/799 RR Rev2)

(PLAN/2024/799)

Legal Basis: Regulation (EC) No 1107/2009 - Article 23(5) in conjunction with Article 13(2)

Procedure: Examination procedure

B.04 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Regulation (EU) renewing the approval of the active substance metrafenone in accordance with Regulation (EC) No 1107/2009 of the European Parliament and of the Council, and amending Commission Implementing Regulation (EU) No 540/2011 (Draft Renewal Report PLAN/2023/2534 RR)

(PLAN/2023/2534)

Legal Basis: Regulation (EC) No 1107/2009 - Article 20(1)

Procedure: Examination procedure

B.05 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Regulation (EU) renewing the approval of the active substance folpet in accordance with Regulation (EC) No 1107/2009 of the European Parliament and of the Council, and amending Commission Implementing Regulation (EU) No 540/2011 (Draft Renewal Report PLAN/2023/646 RR)

(PLAN/2024/646)

Legal Basis: Regulation (EC) No 1107/2009 - Article 20(1)

Procedure: Examination procedure

B.06 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Regulation (EU) amending Implementing Regulation (EU) No 540/2011 as regards the extension of the approval periods of the active substances acequinocyl, aluminium silicate, captan, emamectin, fatty acids C7 to C20, metrafenone, pendimethalin, plant oils / rape seed oil and triclopyr.

(PLAN/2024/1404)

Legal Basis: Regulation (EC) No 1107/2009 - Article 17

Procedure: Examination procedure

Section C **Draft(s) presented for discussion**

C.01 Exchange of views of the Committee on a draft Commission Regulation (EU) amending Commission Regulation (EU) No 547/2011 of 8 June 2011 implementing Regulation (EC) No 1107/2009 of the European Parliament and of the Council as regards labelling requirements for plant protection products

(PLAN/2022/1649)

Legal Basis: Regulation (EC) No 1107/2009 - Article 65(1) and (3), Article 78(1)(m)

Procedure: Regulatory procedure with scrutiny

C.02 Exchange of views of the Committee on a draft Commission Implementing Regulation (EU) concerning the non-renewal of the approval of the active substance metribuzin, in accordance with Regulation (EC) No 1107/2009 of the European Parliament and of the Council, and amending Commission Implementing Regulation (EU) No 540/2011 and Commission Implementing Regulation (EU) 2015/408 (Draft Renewal Report PLAN/2024/1249 RR)

(PLAN/2024/1249)

Legal Basis: Regulation (EC) No 1107/2009 - Articles 20(1) and 78(2)

Procedure: Examination procedure

C.03 Exchange of views of the Committee on a draft Commission Implementing Regulation (EU) concerning the non-renewal of approval of the active substance tritosulfuron in accordance with Regulation (EC) No 1107/2009 of the European Parliament and of the Council and amending Commission Implementing Regulation (EU) No 540/2011 (Draft Renewal Report PLAN/2024/1025 RR)

(PLAN/2024/1025)

Legal Basis: Regulation (EC) No 1107/2009 - Articles 20(3) and 78(2)

Procedure: Examination procedure

C.04 Exchange of views of the Committee on a draft Commission Implementing Regulation (EU) concerning the non-approval of 1,3,7-trimethyl xanthine (caffeine) as a basic substance in accordance with Regulation (EC) No 1107/2009 of the European Parliament and of the Council concerning the placing of plant protection products on the market, and amending the Annex to Implementing Regulation (EU) No 540/2011 (Draft Review Report SANTE/10846/2021)

(SANTE/10844/2021)

Legal Basis: Regulation (EC) No 1107/2009 - Article 23

Procedure: Examination procedure

C.05 Exchange of views of the Committee on a draft Commission Implementing Regulation (EU) concerning the approval of *Allium fistulosum*, processed, as a basic substance in accordance with Regulation (EC) No 1107/2009 of the European Parliament and of the Council concerning the placing of plant protection products on the market, and amending the Annex to Implementing Regulation (EU) No 540/2011 (Draft Review Report PLAN/2024/798)

(PLAN/2024/798)

Legal Basis: Regulation (EC) No 1107/2009 - Article 23

Procedure: Examination procedure