EUROPEAN COMMISSION

Health and Food Safety Directorate General

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Standing Committee on Plants, Animals, Food and Feed
Section *Phytopharmaceuticals - Legislation*
17 - 18 May 2022

CIRCABC Link: *https://circabc.europa.eu/w/browse/ff77c9fd-cbcf-4dea-9ab2-ba61051b9531*

AGENDA

Section A Information and/or discussion

A.01 Summary Report of previous meetings.

A.02 New applications, in particular basic substances:

1. Extension of use of chitosan hydrochloride
2. Extract of the wood of *Quassia amara* L.
3. Hexane

A.03 General issues on approval and renewal of approval processes, in particular:

- overview of active substances under ED stop the clock
- workshop with MS on possible grants
- renewal of active substances: allocation of Rapporteur Member States for active substances which expire between 31 January 2029 and 1 October 2035

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

- New active substances
  1. *Aspergillus flavus* MUCL54911
  2. *Trichoderma atroviride* strain AGR2
  3. *Trichoderma atroviride* strain AT10
  4. Limestone
- Renewal of approval
  5. Clofentezine
  6. Benthiavalicarb
  7. Rape seed oil

8. Oxamyl
9. Triflusulfuron-methyl
10. aluminium ammonium sulfate

- Basic substances
- Amendment of conditions of approval

A.05 Draft Review/Renewal Reports for discussion:

- New active substances
  a) Asulam-sodium
  b) Napropamid-M
- Renewal of approval
  c) Captan
  d) *Pseudomonas chlororaphis* strain MA342
  e) *Bacillus thuringiensis* (horizontal discussion)
  f) *Pythium oligandrum* strain M1
  g) Pelargonic acid
- Basic substances
  h) Hydrogen peroxide silver stabilised
  i) Extension of use of sodium chloride
  j) Lemon essential oil
  k) Yucca Schidigera extract

A.06 Confirmatory information:

1. Pyridaril (amended review report to endorse)
2. Propyzamide (amended review report to endorse)
3. Pendimethalin

A.07 Guidance Documents:

1. Guidance on the assessment of exposure of operators, workers, residents and bystanders in risk assessment of plant protection products (to endorse)
2. EFSA Guidance on aneugenicity assessment (to endorse)
3. EFSA Guidance Document on the risk assessment of plant protection products on bees (*Apis mellifera*, *Bombus* spp. and solitary bees)
4. Draft Guidance document on treatment of seeds and placing on the market of treated seeds under Regulation (EC) No 1107/2009
5. Data requirements and list of agreed test methods - Update of the Communications 2013/C 95/01 and 2013/C 95/02

6. EFSA guidance document for predicting environmental concentrations of active substances of plant protection products and transformation products of these active substances in soil
7. EFSA Scientific Committee (2017) Scientific Opinion on the guidance on the use of the weight of evidence approach in scientific assessments. EFSA Journal 2017;15(8):4971
8. EFSA (2017) Guidance on Uncertainty Analysis in Scientific Assessments. EFSA Journal 2018;16(1):512
9. EFSA Scientific Committee (2017) Guidance on the assessment of the biological relevance of data in scientific assessments. EFSA Journal 2017;15(8):4970

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

- Article 44(4)
- Article 36(3)
- Article 53

A.09 Microorganism Active Substances, in particular:

– Commission Communications in the framework of the implementation of the data requirements

A.10 Safeners and Synergists.

A.11 Updates, clarifications & questions on specific active substances:

1. Sodium hydrogen carbonate
2. Flupyradifurone
3. Acetamiprid

A.12 Article 21:

1. Ipconazole

A.13 General issues for information / discussion:

1. Scope of Regulation (EC) No 1107/2009:
   a) Scope Document rev. 69
   b) New cases: in particular Insect Protector, Cold Atmospheric Plasma
   c) Heptamaloxyloglucan (scope discussion)
   d) FAQ document Fertilising Products Regulation - products out of one single substance + plant biostimulant
2. Basic substances – general issues
3. MS updated survey on timing of regulatory procedures
4. MS-proposal PPP TARIC Code
5. PPPAMS – update

6. Incidents with phosphine products
7. Work plan for the development of test methods focusing on wild pollinators
8. Debriefing workshop on crop protection measures and pollinator protection (7 April 2022)
9. Long term toxicity effects of formulations
10. Residues on cut-flowers
11. SDHI active substances

A.14 Implementation Article 67 Regulation (EC) No 1107/2009.

A.15 Amendment Commission Regulation (EU) No 547/2011.

A.16 Co-formulants, in particular:

1. draft procedures for listing additional unacceptable co-formulants
2. unaccepable co-formulants (mono/polymers and unacceptable concentrations)
3. data collection on co-formulants (EFSA)

A.17 Report from Working Groups, in particular:

1. Working Group on Biopesticides
2. Working Group on environmental relevant topics in the context of Regulation (EC) No 1107/2009
3. Working Group on Seed Treatments (Risk Assessment)
4. Working Group Post Approval Issues
5. Working Group on PPP Formulation Analysis

A.18 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.19 Court cases, requests for internal review, Ombudsman cases.

A.20 Exchange of information from the Pesticide Residues section of the Committee, in particular possible impact on authorizations.

A.21 Scientific publications and information submitted by stakeholders.

A.22 Date of next meeting(s).

A.23 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 substances Straight Chain Lepidopteran Pheromones (acetates) as low-risk substances, Straight Chain Lepidopteran Pheromones (aldehydes), and Straight Chain Lepidopteran Pheromones (alcohols), 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 Commission Implementing Regulation (EU) No 540/2011 (Draft Review Report SANTE/10828/2021).

(SANTE/10826/2021)

Legal Basis: Regulation (EC) No 1107/2009 - Article 20 in conjunction with Article 22(1)

Procedure: Examination procedure

B.02 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Regulation (EU) .../... concerning the non-approval of calcium propionate 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 review report SANTE/11490/2021).

(SANTE/11488/2021)

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

Procedure: Examination procedure

B.03 Exchange of views of the Committee on a draft Commission Implementing Regulation (EU) .../... concerning the non-approval of black soap E470 a 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 review report PLAN/2022/679).

(PLAN/2022/679)

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) .../... amending Commission Implementing Regulation (EU) 2015/408 to update the list of candidates for substitution.

(SANTE/10242/2022)

Legal Basis: Regulation (EC) No 1107/2009 - Articles 78(2) and 80(7)

Procedure: Examination procedure

B.05 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 period of the active substance heptamaloxyloglucan.

(PLAN/2022/1046)

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 Implementing Regulation (EU) …/… renewing the approval of the active substance fish oil as a low risk active 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 Commission Implementing Regulation (EU) No 540/2011 (Draft Review Report SANTE/10076/2022).

(SANTE/10074/2022)

Legal Basis: Regulation (EC) No 1107/2009 - Articles 20 and 22

Procedure: Examination procedure

C.02 Exchange of views of the Committee on a draft Commission Implementing Regulation (EU) …/… renewing the approval of the active substance sheep fat as a low-risk active 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 Annex to Commission Implementing Regulation (EU) No 540/2011 (Draft Review Report SANTE/10072/2022).

(SANTE/10070/2022)

Legal Basis: Regulation (EC) No 1107/2009 - Articles 20 and 22

Procedure: Examination procedure

C.03 Exchange of views of the Committee on a draft Commission Implementing Regulation (EU) …/… amending Implementing Regulation (EU) No 540/2011 as regards the conditions of approval of the active substance penflufen and repealing Implementing Regulation (EU) 2018/185 (Draft Review Report SANTE/10028/2017).

(SANTE/10574/2021)

Legal Basis: Regulation (EC) No 1107/2009 - Articles 21(3) and 78(2)

Procedure: Examination procedure

Pro memoria – TBT notification (to be) launched