EUROPEAN COMMISSION

HEALTH AND FOOD SAFETY DIRECTORATE-GENERAL

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Standing Committee on Plants, Animals, Food and Feed
Section *Phytopharmaceuticals - Plant Protection Products - Legislation*
17 MAY 2017 - 18 MAY 2017

CIRCABC Link: *https://circabc.europa.eu/w/browse/3b2f1718-61bc-4d2d-b713-a3dedb8bed9c*

AGENDA

Section A Information and/or discussion

A.01 Summary Report of previous meetings.

A.02 New active substances:

1. New admissible dossiers to be noted: (no new dossiers)

2. Exchange of view on new European Food Safety Authority (EFSA) conclusions (*no specific conclusions identified*)

3. Commission Draft Review Report and Regulation concerning the (non-) approval of (*no new documents available*)

A.03 Renewal of approval:

1. Annex I Renewal Projects: State of play (no news)

2. Exchange of view on EFSA conclusions:

   i. Thiram
   ii. Propineb
   iii. Oxasulfuron

3. Draft Review/Renewal Reports and Regulations for discussion:

   i. Maleic hydrazide
   ii. 2,4-DB
   iii. Carfentrazone-ethyl
   iv. Acetamiprid (*no discussion – only short information to Member States*)

v. Silthiofam
vi. Isoxaflutole
vii. Imazamox
viii. *Pseudomonas chlororaphis* strain MA342
ix. Iprodione

4. AOB

i. Corrigendum as regards information on purity in the Annex to Regulation renewing the approval of Coniothyrium minitans strain CON/M/91-08.

A.04 Confirmatory Data:

1. Bifenthrin
2. Thiamethoxam
3. Clothianidin
4. Imidacloprid
5. Tetraconazole
6. Cyflumetofen
7. Napropamide
8. Malathion
10. Quinmerac (*revised Review Report to be noted*)
11. Dithianon
12. Tri-allate
13. Eugenol
14. Geraniol
15. Thymol
16. Triazole Derivative Metabolites (TDM)
17. AOB

A.05 Article 21 Reviews (no news)

A.06 Amendment of the conditions of approval:

1. New admissible dossiers to be noted:

i. Penflufen
ii. Metam

A.07 Basic substances:

1. Pilot projects: state of play

2. New dossiers received (only for information):

i. Flavan-gallo tannins

ii. Grape cane tannins

3. Exchange of view on EFSA Technical Reports - (no specific report identified)

4. Draft Review Reports for discussion:

i. Equisetum (extension of use)

A.08 Exchange of views on Guidance Documents.

There are no documents available.

A.09 Notifications under Article 44(4) of Regulation (EC) No 1107/2009 (to be noted).

A.10 Notifications under Article 36(3) of Regulation (EC) No 1107/2009 (to be noted).

A.11 Notifications under Article 53 of Regulation (EC) No 1107/2009 (to be noted).

A.12 News from European Food Safety Authority (EFSA).

A.13 News from the Directorate General for Health and Food Safety (SANTE) Directorate F, Health and Food Audits and Analysis (former FVO).

A.14 Report from working groups:

1. Plant Protection Products Application Management System (PPPAMS)
2. Post Approvals Issues group (PAI) (reminder on zonal applications)
3. Sustainable plant protection experts group Dutch proposal
4. Working group on Biopesticides (no meeting)
5. Working group on Seed Treatments (no meeting)
6. Working Group on implementation of Ruling in C-442/14

A.15 OECD.

A.16 Court cases.

A.17 Endocrine disruptors.

A.18 Minor Uses.

A.19 Interpretation issues:

1. Scope of Regulation (EC) No 1107/2009:

    i. Plant strenghteners (request by Lithuania)

2. Questions and answers

A.20 Classifications under Regulation (EC) No 1272/2008 / REACH:

1. Status of harmonised classifications
2. Preparation of Harmonised Classification and Labelling dossiers (CLH dossiers) by Member States
3. Report from the Working Group (WG) on Assessment Reports (AR template) (merging CLH and xAR templates)

A.21 Glyphosate:

- State of the dossier

A.22 Exchange of information from the Pesticide Residues section of the Committee: possible impact on authorisations (no new meeting has taken place since March 2017).

A.23 Evaluation of the EU legislation on plant protection products and pesticides residues (Regulation (EC) No 1107/2009 and Regulation (EC) No 396/2005).

A.24 Exposure of florists to plant protection products from cutflowers (follow-up from March meeting).

A.25 New rules on availability of draft documents for discussion.

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 renewing the approval of the active substance propoxycarbazone 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 Renewal Report SANTE/11954/2016 Rev. 1).

(B.01_SANTE_11953_2016 Rev. 2))

Legal Basis: Article 20(1) of Regulation (EC) No 1107/2009
Procedure: Examination procedure

B.02 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Regulation renewing the approval of the active substance benzoic acid 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 Renewal Report SANTE/10147/2017 Rev 1).

(B.02_SANTE_10146_2017 Rev. 1))

Legal Basis: Article 20(1) of Regulation (EC) No 1107/2009
Procedure: Examination procedure

B.03 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Regulation renewing the approval of the active substance pendimethalin, as a candidate for substitution, 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/11656/2016).

(B.03_SANTE_11655_2016 Rev. 0)

Legal Basis: Article 20(1) of Regulation (EC) No 1107/2009
Procedure: Examination procedure

B.04 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Regulation concerning the non-renewal of approval of the active substance picoxystrobin, 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 Renewal Report SANTE/11601/2016 Rev. 2).

(B.04_SANTE_11600_2016 Rev. 3)

Legal Basis: Article 20(1) and Article 78(2) of Regulation (EC) No 1107/2009
Procedure: Examination procedure

B.05 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Regulation renewing the approval of the active substance propyzamide,

as a candidate for substitution, 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 11797/2016 Rev. 1).

(B.05_SANTE_11796_2016 Rev. 0)

Legal Basis: Article 20(1) of Regulation (EC) No 1107/2009
Procedure: Examination procedure

B.06 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Regulation withdrawing the approval of the active substance repellents by smell of animal or plant origin/tall oil pitch, 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 SANCO/2632/08 Rev. 5).

(B.06_SANTE_11587_2016 Rev. 0)

Legal Basis: Article 21(3) and Article 78(2) of Regulation (EC) No 1107/2009
Procedure: Examination procedure

B.07 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Regulation withdrawing the approval of the active substance repellents by smell of animal or plant origin/tall oil crude, 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 SANCO/2631/08 Rev. 5).

(B.07_SANTE_11586_2016 Rev. 0)

Legal Basis: Article 21(3) and Article 78(2) of Regulation (EC) No 1107/2009
Procedure: Examination procedure

Miscellaneous

M.01 Scientific publications and information submitted by stakeholders.

M.02 AOB

M.03 Date of next meeting.