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*
19 JULY 2017 - 20 JULY 2017

CIRCABC Link: *https://circabc.europa.eu/w/browse/726e71b2-23c2-4cf3-ad04-480dbc372284*

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:

i. 24-Epibrassinolide

2. Exchange of views on new European Food Safety Authority (EFSA) conclusions:

i. Beauveria bassiana strain IMI389521

3. Commission Draft Review Report and Regulation concerning the (non-) approval of:

A.03 Renewal of approval:

1. Annex I Renewal Projects: State of play

2. Exchange of view on EFSA conclusions:

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

i. 2,4-DB
ii. Carfentrazone-ethyl
iii. Propineb
iv. Pseudomonas chlororaphis strain MA342
v. Iprodione (no discussion – only short information update for Member States)
vi. Oxasulfuron

vii. Thiram
viii. Bifenazate
ix. Bentazone

4. Update on the decision making for picoxystrobin

A.04 Confirmatory Data:

1. Bifenthrin
2. Thiamethoxam
3. Clothianidin
4. Imidacloprid
5. Tetraconazole
6. Cyflumetofen (no news, written comments before 8/9)
7. Napropamide
8. Malathion
9. Dithianon
10. Tri-allate
11. Eugenol
12. Geraniol
13. Thymol
14. Triazole Derivative Metabolites (TDM)
15. Straight Chain Lepidopteran Pheromones (SCLP) (draft revised review report referring to EFSA conclusions on confirmatory information and including new compound belonging to the SCLP)
16. Terbuthylazine
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. Fenazaquin (*no discussion – see documents on CIRCABC*)

2. Exchange of view on EFSA conclusions:

No new EFSA conclusion available

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

i. Penflufen (no news, written comments before 8/9/2017)

A.07 Basic substances:

1. Pilot projects: state of play

Quassia

2. New dossiers received (only for information)

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

4. Draft Review Reports for discussion:

i. Equisetum (extension of use) (review report to take note)
ii. Potassium sorbate
iii. Beer
iv. Mustard powder

A.08 Exchange of views on Guidance Documents:

1. Template to be used for Assessment Reports (SANCO/12592/2012 Rev. 1, to be noted)
2. Guidance Document on Data Protection (SANCO/12576/2012 Rev. 2.2, to be noted)
3. Guidance document on zonal evaluation and mutual recognition under Regulation (EC) No 1107/2009 (SANCO/13169/2010 Rev. 10, to be noted)
4. Guidance document on the presentation and evaluation of plant protection product dossiers in the format of a (draft) Registration Report (SANCO/6895/2009 Rev. 2, to be noted)
5. Terms of Reference of the Working Group on Post Approval Issues from the Standing Committee on Animals, Plants, Food and Feed: section Pesticide Legislation (SANTE/11102/2017 to be noted)

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)
3. Sustainable plant protection experts group Dutch proposal (no meeting)
4. Working group on Biopesticides
5. Working group on Seed Treatments (no meeting)
6. Working Group on Co-formulants
7. Working Group on Low-risk criteria

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
- Draft Review Report and Regulation for discussion

A.22 Exchange of information from the Pesticide Residues section of the Committee: possible impact on authorisations.

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.

A.25 Pepino Mosaic Virus – use by tomato plant propagators.

A.26 New mandate for a Working Group (WG) to set up a procedure to assess new variants of approved active substances.

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 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. 3).

(B.01_SANTE_11796_2016 Rev. 1)

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 concerning the non-renewal of approval of the active substance DPX KE 459 (flupyrsulfuron-methyl), 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/10317/2015 Rev. 3).

(B.02_SANTE 10316 2015 Rev. 0)

Legal Basis: Article 20(1) and Article 78(2) 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 concerning the non-approval of the active substance beta-cypermethrin, 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/10237/2017 Rev. 1 (formerly SANTE/12481/2015 Rev. 4)).

(B.03_SANTE_10236_2017 Rev. 1)

Legal Basis: Article 13 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 approving the basic substance sodium chloride 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/10383/2017 Rev. 1).

(B.04_SANTE_10381_2017 Rev 1)

Legal Basis: Article 23(5) 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 confirming the conditions of approval of the active substance 8-hydroxyquinoline, as set out in Implementing Regulation (EU) No 540/2011 and modifying the Commission Implementation Regulation (EU) 2015/408 of 11 March 2015 on implementing Article 80(7) of Regulation (EC) No 1107/2009 of the European Parliament and of the Council concerning the placing of plant protection products on the market and establishing a list of candidates for substitution (Draft Addendum to the Review Report SANTE/11618/2016 Rev. 0.1).

(B.05_SANTE_11620_2016 Rev. 3)

Legal Basis: Article 13 and 80(7) 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 renewing the approval of the active substance 2,4-DB in accordance with Regulation (EC) No 1107/2009 of the European Parliament and 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/10066/2017 Rev. 4).

(B.06_SANTE_10065_2017 Rev.1)

Legal Basis: Article 20(1) 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 renewing the approval of the active substance carfentrazone-ethyl in accordance with Regulation (EC) No 1107/2009 of the European Parliament and 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/10144/2017 Rev. 3).

(B.07_SANTE_10143_2017 Rev. 1)

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

B.08 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Regulation renewing the approval of the active substance imazamox 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 Renewal Report SANTE/10499/2017 Rev. 3).

(B.08_SANTE_10498_2017 Rev. 1)

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

B.09 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Regulation renewing the approval of the active substance maleic hydrazide, 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/10561/2017 Rev. 2).

(B.09_SANTE_10560_2017 Rev.1)

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

B.10 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Regulation amending Implementing Regulation (EU) No 540/2011 as regards the extension of the approval period of the active substance quizalofop-p-tefuryl.

(B.10_SANTE_10405_2017 Rev. 2)

Legal Basis: Article 17 of Regulation (EC) No 1107/2009
Procedure: Examination procedure

B.11 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Regulation renewing the approval of the active substance silthiofam in accordance with Regulation (EC) No 1107/2009 of the European Parliament and 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/11799/2016 Rev. 2).

(B.11_SANTE_11798_2016 Rev. 0)

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

B.12 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Regulation renewing the approval of the active substance acetamiprid in accordance with Regulation (EC) No 1107/2009 of the European Parliament and 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/10502/2017 Rev. 2).

(B.12_SANTE_10501_2017)

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

B.13 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Regulation amending Implementing Regulation (EU) No 540/2011 as regards the extension of the approval periods of the active substances cyflufenamid, fluopicolide, heptamaloxyloglucan and malathion.

(B.13_SANTE_10326_2017)

Legal Basis: Article 17 of Regulation (EC) No 1107/2009
Procedure: Examination procedure

B.14 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Regulation as regards the extension of the approval periods of the active substances 1-methylcyclopropene, 2,4-DB, beta-cyfluthrin, chlorothalonil, chlorotoluron, cypermethrin, daminozide, deltamethrin, dimethenamid-p, flufenacet, flurtamone, forchlorfenuron, fosthiazate, indoxacarb, iprodione, maleic hydrazide, MCPA, MCPB, silthiofam, thiophanate-methyl and tribenuron.

(B.14_SANTE_10445_2017 Rev. 1)

Legal Basis: Article 17 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.