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Standing Committee on Plants, Animals, Food and Feed
Section *Phytopharmaceuticals - Legislation*
05 OCTOBER 2017 - 06 OCTOBER 2017

CIRCABC Link: *https://circabc.europa.eu/w/browse/2a806ef7-3337-4dcb-b0fb-fd58c3f28ca2*

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:
   a. Beauveria bassiana 203
2. Exchange of views on new European Food Safety Authority (EFSA) conclusions:
3. Commission Draft Review Report and Regulation concerning the (non-) approval of:
   a. Beauveria bassiana strain IMI389521

A.03 Renewal of approval:

1. Annex I Renewal Projects: State of play
2. Exchange of view on EFSA conclusions:
   a. Pethoxamid
   b. Mepanipyrim
   c. Bromoxynil
   d. Chlorpropham
   e. Propiconazole
3. Draft Review/Renewal Reports and Regulations for discussion:
   a. Propineb
   b. Pseudomonas chlororaphis strain MA342
   c. Oxasulfuron
   d. Thiram
   e. Bifenazate (short update only)
   f. Bentazone (short update only)
   g. Mecoprop-P

h. Carfentrazone-ethyl
i. Laminarin
j. Propyzamide
k. Silthiofam
l. Pymetrozine
m. Isoxaflutole
n. Clonostachys rosea J1446

A.04 Confirmatory Data:

1. Bifenthrin
2. Thiamethoxam (short update only)
3. Clothianidin (short update only)
4. Imidacloprid (short update only)
5. Cyflumetofen (discussion only)
6. Malathion
7. Dithianon
8. Tri-allate
9. Eugenol
10. Geraniol
11. Thymol
12. Terbuthylazine
13. Iprovalicarb
14. Metazachlor
15. Pyretrins (drr to take note)
16. Acetic acid (drr to take note)
17. Picloram
18. Chlorsulfuron
19. Triazine amine (common metabolite)
20. AOB

A.05 Article 21 Reviews (no news).

A.06 Amendment of the conditions of approval:

1. New admissible dossiers to be noted:
   a. Fenazaquin (no discussion – see documents on CIRCABC)
2. Exchange of view on EFSA conclusions:
   a. No new EFSA conclusion available
3. Draft Review/Renewal Reports and Regulations for discussion:
   a. Penflufen (no news)

A.07 Basic substances:

1. Pilot projects: state of play
2. New dossiers received (only for information)
3. Exchange of views on EFSA Technical Reports.
   a. Talc
   b. Vinegar
4. Draft Review Reports for discussion:
   a. Saponaria officinalis root extract

A.08 Exchange of views on Guidance Documents:

1. Template to be used for Assessment Reports (SANCO/12592/2012 Rev. 1, for discussion and possible noting)
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 for discussion)
6. Report from the Danish EPA workshop on data requirements for acute inhalation toxicity testing – follow-up
7. Guidance document on the Evaluation Efficiency of Residue Analytical Methods (SANTE/10632/2017, for discussion)
8. Guidance document on the establishment of the residue definition for dietary risk assessment (SANTE/11644/2017, 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.

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

1. Notifications (to be noted)
2. Update of the Working document on emergency authorisations according to Article 53 (for information)
3. Mandate to EFSA under Article 53(2) in relation with emergency authorisations granted in 2017 for products containing imidacloprid, clothianidin, and thiametoxam

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.

A.14 Reports 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 (no meeting)
5. Working group on Seed Treatments (no meeting)
6. Working Group on Co-formulants
7. Working Group on Low-risk criteria

A.15 Organisation for Economic Co-operation and Development (OECD).

A.16 Court cases.

New court case: T-476/17 Arysta LifeScience v. Commission – Application for the annulment of Commission Implementing Regulation (EU) 2017/855 on diflubenzuron.

A.17 Endocrine Disruptors.

1. State of play: ED-criteria and development of EFSA/ECHA guidance document
2. Implementation of the new ED Criteria renewal active substances: Amending Implementing Regulation 844/2012 in view of the scientific criteria for the determination of endocrine disrupting properties
3. Implementation of the new ED Criteria new active substances

A.18 Minor Uses.

A.19 Interpretation Uses.

1. Scope of Regulation (EC) No 1107/2009:
   a. 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 – Amending Implementation Regulation 844/2012 in view of the harmonised classification of active substances
3. Follow-up of the merging of CLH and xAR templates (discussion only)

A.21 Glyphosate.

a. State of the dossier
b. 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 Adoption of the mandate for a Working Group (WG) to set up a procedure to assess new variants of approved active substances.

A.27 2,4-D – Revision of AOEL, ADI and ARfD (revised review report to be noted).

A.28 Protection goals for environmental risk assessment – update on next steps.

A.29 Pest management changes after neonicotinoid and fipronil restrictions: results from a survey (Article publicly available).

A.30 Initial information concerning Brexit.

A.31 Scientific publications and information submitted by stakeholders.

A.32 Date of next meeting.

Section B <u>Draft(s) presented for an opinion</u>

B.01 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 Rev2).

(SANTE/11620/2016)

Legal Basis: Regulation (EC) No 1107/2009 - Article 13 and 80(7)

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 iprodione 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 Renewal Report SANTE/10627/2017 Rev1).

(SANTE/10626/2017)

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

Procedure: Examination procedure

B.03 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 flonicamid (IKI-220), metalaxyl, penoxsulam and proquinazid.

(SANTE/10327/2017)

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

Procedure: Examination procedure

B.04 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Regulation concerning the non-approval of paprika extract (capsanthin, capsorubin E 160c) 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 Doc. SANTE/10068/2017)

(SANTE/10067/2017)

Legal Basis: Regulation (EC) No 1107/2009 - Articles 13(2) and 23(5)

Procedure: Examination procedure

B.05 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Regulation concerning the non-approval of *Achillea millefolium* L. 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 Doc. SANTE/10142/2017).

(SANTE/10100/2017)
Legal Basis: Regulation (EC) No 1107/2009 - Articles 13(2) and 23(5)
Procedure: Examination procedure

B.06 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Regulation concerning the approval of beer 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 Doc. SANTE/11038/2017 Rev1).

(SANTE/11037/2017)

Legal Basis: Regulation (EC) No 1107/2009 - Articles 13(2) and 23(5)

Procedure: Examination procedure

B.07 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Regulation concerning the non-approval of potassium sorbate 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 Doc. SANTE/11031/2017).

(SANTE/11029/2017)

Legal Basis: Regulation (EC) No 1107/2009 - Articles 13(2) and 23(5)

Procedure: Examination procedure

B.08 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Regulation concerning the approval of mustard seeds 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 Review Report Doc. SANTE/11309/2017).

(SANTE/11307/2017)

Legal Basis: Regulation (EC) No 1107/2009 - Articles 13(2) and 23(5)

Procedure: Examination procedure