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Health and Food Safety Directorate General

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Standing Committee on Plants, Animals, Food and Feed
Section *Phytopharmaceuticals - Legislation*
22-23 MARCH 2018

CIRCABC Link: ***https://circabc.europa.eu/w/browse/45f8503d-de53-4bc5-a291-82b6a2f865a2***

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. *Bacillus amyloliquefaciens* IT-45
   b. *Pepino mosaic virus* (*PepMV*) Chilean (CH2) strain, mild isolate Abp2 (PEPMVO)
   c. *Pepino mosaic virus* (PepMV) European (EU) strain, mild isolate Abp1 (PEPMVO)
   d. Limestone
2. Exchange of views on new European Food Safety Authority (EFSA) conclusions:
   a. Pasteuria nishizawae Pn1
   b. Metschnikowia fructicola NRRL Y-27328 (NAS 1107)
3. Commission Draft Review Report and Regulation concerning the (non-) approval of:
   a. Flutianil
   b. Fenpicoxamid (XDE-777)

A.03 Renewal of approval:

1. Annex I Renewal Projects: State of play
   a. 5th renewal programme
2. Exchange of view on EFSA conclusions:
   a. Desmedipham
   b. Phenmedipham
   c. Copper compounds

3. Draft Review/Renewal Reports and Regulations for discussion:
    a. Chlorpropham
    b. Pseudomonas chlororaphis strain MA342
    c. Quinoxyfen
    d. Diquat (short update only)
    e. Mecoprop-P
    f. Carfentrazone-ethyl
    g. Propyzamide
    h. Silthiofam
    i. Mepanipyrim
    j. Tribenuron (short update only)
    k. Flurtamone
    l. Propiconazole
    m. Etoxazole
    n. Fenamidone
    o. Trifloxystrobin
    p. Pethoxamid
    q. Methoxyfenozide
    r. Ampelomyces quisqualis AQ10

A.04 Confirmatory Data:

1. Dithianon (short update only)
2. Terbuthylazine
3. Iprovalicarb (review report to be noted)
4. Thiencarbazone-methyl (review report to be noted)
5. Mandipropamid (review report to be noted)
6. Bupimirate (review report to be noted)
7. Azimsulfuron (review report to be noted)
8. Tau-fluvalinate (review report to be noted)
9. Disodium phosphonate (review report to be noted)
10. Ipconazole (review report)
11. Urea (review report)

A.05 Article 21 Reviews.

Reports from EFSA on imidacloprid, clothianidin and thiamethoxam

A.06 Amendment of the conditions of approval:

1. New admissible dossiers to be noted:
2. Exchange of view on EFSA conclusions:
3. Draft Review/Renewal Reports and Regulations for discussion:

A.07 Basic substances:

1. Pilot projects: state of play
2. New dossiers received (only for information)
   a. Lecithin extension
   b. wheat flour
   c. clayed charcoal (extension)
3. Exchange of views on EFSA Technical Reports
4. Draft Review Reports for discussion:
   a. Landes pine tar (short update only)
   b. Lecithin extension

A.08 Exchange of views on Guidance Documents:

1. Draft revised Guidance Document on the Renewal of Authorisations according to Article 43 of Regulation (EC) No 1107/2009 (for discussion)
2. Draft revised Guidance Document on Zonal Evaluation, Mutual Recognition Withdrawal and Amendment of Authorisations under Regulation (EC) No 1107/2009 (SANCO/13169/2010 Rev. 11, feedback on comments received)

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.

1. New notifications (to be noted)
2. Differences in application of article 36(3) amongst Member States
3. On-board fumigation of grain

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

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

1. List of End-Points – update of the residue section

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)
   a. Update on the November meeting – products containing 2,4D
3. Working group on Biopesticides
4. Working group on Seed Treatments (no update)
5. Working Group on Co-formulants (no update)
6. Working Group on Low-risk criteria

A.15 OECD.

1. Pesticide Notification Information System (OECD secretariat)
2. Update on the new development about the Global Harmonised Submission Transport Standard (EFSA)
3. News on the work of the WGP and the related groups

A.16 Court cases.

1. Case T-719/17- annulment of Commission Implementing Regulation (EU) No 2017/11496 concerning the non-renewal of the approval of the active substance flupyrsulfuron-methyl

A.17 Endocrine Disruptors.

1. State of play: ED criteria and draft EFSA/ECHA guidance document
2. Implementation of the new ED Criteria renewal active substances: Amending Implementing Regulation (EU) No 844/2012 in view of the scientific criteria for the determination of endocrine disrupting properties

A.18 Minor Uses.

A.19 Interpretation issues:

1. Scope of Regulation (EC) No 1107/2009:
   a. Lava meal (Belgium)
   b. Salvis freeze (Belgium)
   c. Straw pellets (Belgium)
   d. Moss control / fertilizers (Denmark)
   e. Uses against lichens on trees (Austria)

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 (EU) No 844/2012 in view of the harmonised classification of active substances
3. Report on the alignment of the classification and peer-review processes

A.21 Glyphosate.

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 Information concerning Brexit.

A.25 Draft COM Notice concerning a list of potentially low-risk substances (presentation).

A.26 Scientific publications and information submitted by stakeholders.

A.27 Confirmatory data pending and renewal ongoing – Clofentezine and Difeconazole (RMS ES).

A.28 Summary of the Workshop on Toxicological Risk Assessment of Plant Protection Products (Paris, 13-14 March 2018).

A.29 Update on acetamiprid.

A.30 Iprodione: updated toxicological reference values for a non-renewed substance (request from Belgium, for note taking).

A.31 Propargite: new toxicological reference values for setting of import tolerances (for note taking).

A.32 Revision CIRCABC Account users.

A.33 New AGM system for the organisation of meetings and reimbursement of participants.

A.34 Date of next meeting.

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 approval of active substance bentazone 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/12012/2015 rev. 7).

(SANTE/12011/2015 Rev9)

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 approving the basic substance Talc E553B 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/11639/2017 Rev1).

(SANTE/11638/2017 Rev1)

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 Decision on the establishment of a work programme for the renewal of active substances expiring in 2022, 2023 and 2024 in accordance with Article 18 of Regulation (EC) No 1107/2009.

(SANTE/11750/2017 Rev1)

Legal Basis: Regulation (EC) No 1107/2009 - Article 18

Procedure: Examination procedure

B.04 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).

(SANTE/11796/2016 Rev1)

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 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).

(SANTE/11798/2016)

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 renewing the approval of the active substance forchlorfenuron 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/11643/2017 Rev 2).

(SANTE/11642/2017 Rev2)

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

Procedure: Examination procedure

B.07 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 conditions of approval of the active substance malathion.

(SANTE/10859/2017 Rev1)

Legal Basis: Regulation (EC) No 1107/2009 - Article 7(1)

Procedure: Examination procedure

B.08 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 conditions of approval of the active substance fenazaquin (Draft addendum to the review report SANTE/11781/2017 rev 0).

(SANTE/11780/2017 Rev1)

Legal Basis: Regulation (EC) No 1107/2009 - Article 7(1)

Procedure: Examination procedure

B.09 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Regulation approving the low-risk active substance *Pasteuria nishizawae* Pn1, 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/10160/2018 rev. 0.1).

(SANTE/10159/2018 Rev1)

Legal Basis: Regulation (EC) No 1107/2009 - Article 22(1) in conjunction with Article 13(2)

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 periods of the active substances bromuconazole, buprofezin, haloxyfop-P and napropamide.

(SANTE/11624/2017 Rev1)

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

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 zoxamide 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/10052/2018).

(SANTE/10051/2018 Rev1)

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

Procedure: Examination procedure

B.12 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 thiram 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/11020/2017 Rev 2).

(SANTE/11019/2017 Rev3)

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

Procedure: Examination procedure

B.13 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 pymetrozine 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/00103/2015 Rev 3).

(SANTE/00102/2015 Rev1)

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

Procedure: Examination procedure

B.14 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 oxasulfuron 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/10886/2017 Rev 0).

(SANTE/10885/2017 Rev1)

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

Procedure: Examination procedure