EUROPEAN COMMISSION

Health and Food Safety Directorate General

**sante.ddg2.g.5(2018)110237**

**Standing Committee on Plants, Animals, Food and Feed**

**Section *Phytopharmaceuticals - Legislation***

**25 JANUARY 2018 - 26 JANUARY 2018**

***CIRCABC Link:*** *https://circabc.europa.eu/w/browse/a67ec92e-9989-498e-a48f-94361ee0a66e*

# 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:
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. Flutianil

**A.03** Renewal of approval:

1. Annex I Renewal Projects: State of play
   a. Draft Commission Decision concerning the fifth renewal project (AIR 5)
   b. AIR 5 work programme
2. Exchange of view on EFSA conclusions:
   a. Methoxyfenozide
   b. Trifloxystrobin
3. Draft Review/Renewal Reports and Regulations for discussion:
   a. Chlorpropham
   b. Pseudomonas chlororaphis strain MA342
   c. Oxasulfuron
   d. Thiram
   e. Diquat (short update only)
   f. Mecoprop-P
   g. Carfentrazone-ethyl
   h. Propyzamide
   i. Silthiofam
   j. Clonostachys rosea J1446

k. Forchlorfenuron
l. Mepanipyrim
m. Tribenuron
n. Flurtamone (short update only)
o. Propiconazole
p. Etoxazole
q. Fenamidone (short update only)
r. Zoxamide

**A.04** Confirmatory Data:

1. Malathion
2. Dithianon
3. Tri-allate
4. Terbuthylazine
5. Iprovalicarb (to be noted)
6. Metazachlor
7. Picloram (to be noted)
8. Chlorsulfuron
9. Pseudomonas sp. Strain DSMZ 13134 (review report to be noted)
10. Pyroxsulam (review report to be noted)
11. Chlorantraniliprole (review report to be noted)
12. Halauxifen-methyl (review report to be noted)

**A.05** Article 21 Reviews (no news).

**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. Fenazaquin

**A.07** Basic substances:

1. Pilot projects: state of play
2. New dossiers received (only for information)
   a. Lecithin extension
3. Exchange of views on EFSA Technical Reports
4. Draft Review Reports for discussion:
   a. *Saponaria officinalis* root extract
   b. Sodium hydrogen carbonate (updated review report to be noted)

**A.08** Exchange of views on Guidance Documents:

1. 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)
2. Proposed Mandate to revise the Guidance Document on significant and non-significant changes of the chemical composition of authorised plant protection products under Regulation (EC) No 1107/2009 (SANCO/12638/2011, rev. 2, 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.

1. New notifications (to be noted)
2. Differences in application of article 36(3) amongst Member States
3. On-board fumigation of grain: rejection of mutual recognition for a phosphine product

**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).

**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. 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 follow-up discussion)
   b. Update on the November meeting
3. Sustainable plant protection experts group Dutch proposal (no meeting)
4. Working group on Biopesticides
5. Working group on Seed Treatments (short update)
6. Working Group on Co-formulants
7. Working Group on Low-risk criteria

**A.15** OECD.

1. Pesticide Notification Information System

**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. Plant strenghteners (request by Lithuania)
   b. Fertinema (request by Belgium)
   c. A Polyvinyl alcohol-based product to reduce pod shattering on rapeseed crops (request by Belgium)

**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** Mandate for a Working Group (WG) to set up a procedure to assess new variants of approved active substances (to be noted).

**A.25** Information concerning Brexit.

**A.26** Draft COM Notice concerning a list of potentially low-risk substances (presentation).

**A.27** Scientific publications and information submitted by stakeholders.

**A.28** Confirmatory data pending and renewal ongoing – Clofentezine and Difeconazole (RMS ES).

**A.29** 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 Rev7).

(SANTE/12011/2015 Rev7)

**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 amending Implementing Regulation (EU) No 540/2011 as regards the conditions of approval of the active substance bifenthrin (Draft Review Report SANCO/12946/2011 final Rev4).

(SANTE/10317/2017)

**Legal Basis:** Regulation (EC) No 1107/2009 - Article 21(3) in conjuction with Article 78(2)

**Procedure:** Examination procedure

**B.03** 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)

**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 concerning the non-renewal of approval of active substance propineb 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/11034/2017 Rev0).

(SANTE/11035/2017)

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

**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 the active substance Reynoutria sachalinensis extract, 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.

(SANTE/10102/2016 Rev3)

**Legal Basis:** Regulation (EC) No 1107/2009 - Article 13(2)(b)

**Procedure:** Examination procedure

**B.06** 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 *Bacillus subtilis* (Cohn 1872) Strain QST 713, identical with strain AQ 713, clodinafop, clopyralid, cyprodinil, dichlorprop-P, fosetyl, mepanipyrim, metconazole, metrafenone, pirimicarb, *Pseudomonas chlororaphis* strain: MA 342, pyrimethanil, quinoxyfen, rimsulfuron, spinosad, thiacloprid, thiamethoxam, thiram, tolclofos-methyl, triclopyr, trinexapac, triticonazole and ziram, amending the Annex to Implementing Regulation (EU) No 540/2011.

(SANTE/11935/2017 Rev1)

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

**Procedure:** Examination procedure