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HEALTH AND FOOD SAFETY DIRECTORATE-GENERAL

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Standing Committee on Plants, Animals, Food and Feed
Section *Phytopharmaceuticals - Plant Protection Products - Legislation*
22 MARCH 2017 - 23 MARCH 2017

CIRCABC Link: *https://circabc.europa.eu/w/browse/a9b05b29-e99a-437b-b980-9f13c1799e72*

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. Lavandulyl senecioate
   ii. Sweet Lupin (seeds), *Lupinus albus* L., germ., ext.

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:

   i. *Beauveria bassiana* strain 147
   ii. *Beauveria bassiana* NPP111B005

A.03 Renewal of approval:

1. Annex I Renewal Projects: State of play

2. Exchange of view on EFSA conclusions:

   i. Thiram

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

i. Coniothyrium minitans strain CON/M/91-08
ii. Maleic hydrazide (No detailed discussion)
iii. Mesotrione
iv. Pendimethalin
v. 2,4-DB
vi. Carfentrazone-ethyl
vii. Acetamiprid
viii. Propyzamide
ix. Bentazone
x. Silthiofam
xi. Isoxaflutole

A.04 Confirmatory Data:

1. Bifenthrin
2. Thiamethoxam
3. Clothianidin
4. Imidacloprid
5. Tetraconazole
6. Cyflumetofen
7. Napropamide
8. Malathion
9. Fluroxypyr (review report to be noted in conjunction with item B.10)
10. Quinmerac
11. Dithianon
12. AOB

A.05 Article 21 Reviews:

i. Thiametoxam, clothianidin, imidacloprid (other uses than seed treatments and granules) (*amended schedule of review*)

A.06 Amendment of the conditions of approval:

1. New admissible dossiers to be noted.

A.07 Basic substances:

1. Pilot projects: state of play

2. New dossiers received (only for information):

i. Equisetum (extension of use)

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

4. Draft Review Reports for discussion:

i. Honey from rhododendron (*No detailed discussion; Member States are requested to send in comments after the meeting.*)
ii. Sodium chloride (*No detailed discussion; Member States are requested to send in comments after the meeting.*)

A.08 Exchange of views and possible taking note of the following Guidance Documents:

1. Guidance Document on the renewal of approval of active substances to be assessed in compliance with Regulation (EU) No 844/2012 (for discussion - changes of specification).

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 since December)
4. Drift Risk Assessment Workshop (DRAW). Improving Representation, Management and Mitigation of Spray Drift for Plant Protection Products in Arable Crops (8-9 February 2017).

A.15 OECD.

A.16 Bees:

1. AOB

A.17 Court cases.

A.18 Endocrine disruptors.

A.19 Minor Uses.

A.20 Interpretation issues:

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

   i. Colour spray
   ii. Plant strenghteners (request by LT)

2. Questions and answers

A.21 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.22 Glyphosate:

- State of the dossier

A.23 Exchange of information from the Pesticide Residues section of the Committee: possible impact on authorisations.

A.24 Evaluation of the EU legislation on plant protection products and pesticides residues (Regulation (EC) No 1107/2009 and Regulation (EC) No 396/2005).

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 Regulation amending Regulation (EC) No 1107/2009 of the European Parliament and the Council concerning the placing of plant protection products on the market as regards the criteria for the approval of low risk active substances.

(B.01_SANTE_12376_2015)

Legal Basis: Article 22(3) and Article 78(1)(a) of Regulation (EC) No 1107/2009
Procedure: Regulatory procedure with scrutiny

B.02 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Regulation renewing the approval of the active substance flazasulfuron 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/10153/2017 Rev. 1)

(B.02_SANTE_10152_2017)

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 approving the low-risk active substance Bacillus amyloliquefaciens strain FZB24, 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. (Draft Review Report SANTE/12037/2016 Rev. 1)

(B.03_SANTE_12036_2016)

Legal Basis: Article 13(2)(a) and 22(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 approving the active substance Beauveria bassiana strain NPP111B005, 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/10398/2016 Rev. 1)

(B.04_SANTE_10397_2016 Rev. 2)

Legal Basis: Article 13(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 approving the active substance *Beauveria bassiana* strain 147, 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/10424/2016 Rev. 1)

(B.05_SANTE_10423_2016 Rev. 2)

Legal Basis: Article 13(2) 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 mesosulfuron 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/11827/2016 Rev. 2)

(B.06_SANTE_11825_2016 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 mesotrione 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/11654/2016 Rev. 1)

(B.07_SANTE_11653_2016)

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 cyhalofop-butyl 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/10879/2015 Rev. 2)

(B.08_SANTE_10878_2015 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 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. 1)

(B.09_SANTE_11600_2016 Rev. 1)

Legal Basis: Article 20(1) and Article 78(2) 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 conditions of approval of the active substance fluroxypyr. (Draft Review Report SANCO/11019/2011 Rev. 5)

(B.10_SANTE_10978_2016 Rev. 0)

Legal Basis: Article 21(3) and Article 78(2) 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 withdrawing the approval of the active substance methyl nonyl ketone, 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 Implementing Regulation (EU) No 540/2011.

(B.11_SANCO_11754_2016 Rev. 2)

Legal Basis: Article 21(3) and Article 78(2) 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 (EU) No 540/2011 as regards the extension of the approval periods of the active substances alpha-cypermethrin, *Ampelomyces quisqualis* strain: aq 10, benalaxyl, bentazone, bifenazate, bromoxynil, carfentrazone ethyl, chlorpropham, cyazofamid, cyhalofop butyl, desmedipham, diquat, DPX KE 459 (flupyrsulfuron-methyl), etoxazole, famoxadone, fenamidone, flumioxazine, foramsulfuron, *Gliocladium catenulatum* strain: j1446, imazamox, isoxaflutole, laminarin, mesotrione, metalaxyl-m, methoxyfenozide, milbemectin, oxasulfuron, pendimethalin, phenmedipham, pymetrozine, s-metolachlor, and trifloxystrobin

(B.12_SANTE_10211_2017 Rev. 1)

Legal Basis: Article 17 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 conditions of approval of the active substance diflubenzuron. (Draft Review Report SANCO/831/08 Rev. 6)

(B.13_SANTE_11216_2016 Rev. 2)

Legal Basis: Article 21(3) and Article 78(2) 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 concerning the non-approval of the active substance orthosulfamuron, 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 SANTE/11756/2016 Rev. 0)

(B.14_SANTE_11753_2016 Rev.1)

Legal Basis: Article 13(2) and Article 80(1) of Regulation (EC) No 1107/2009
Procedure: Examination procedure

Miscellaneous

M.01 New Scientific publications and information submitted by stakeholders.

M.02 AOB

1. Antibiotics – Yearly reporting by Member States

M.03 Date of next meeting.