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DIRECTORATE-GENERAL FOR HEALTH AND FOOD SAFETY

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STANDING COMMITTEE ON PLANTS, ANIMALS, FOOD AND FEED

Section *Phytopharmaceuticals - Plant Protection Products - Legislation*

26 JANUARY 2015 - 27 JANUARY 2015

CIRCABC Link: *https://circabc.europa.eu/w/browse/2e04b70e-4392-49f3-91a9-0d99b25f0422*

AGENDA

Section A **Information and/or discussion**

A.01 Summary Report of previous meetings.

A.02 Stage 4 of the review programme under Directive 91/414 – "Green Track".

A.03 New active substances:

1. New admissible dossiers - *no new dossiers*
2. EFSA conclusions
    i. *Flumetralin*
    ii. *Halauxifen-methyl*
    iii. *3-decene-2-one*
    iv. *Pepino mosaic virus CH2*
3. First discussion of a Commission Draft Review Report and Regulation concerning the approval of:
    i. Cyantraniliprole
4. *Chromobacterium subtsugae* PRAA4-1 (MBI-203)

A.04 Renewal of approval:

1. Draft Working Document Renewal Programme (Doc. SANCO/11284/2012 Rev. 15) (For information)
2. Applications for renewal of approval of active substances submitted under Article 14 of Regulation (EU) No 1107/2009 and in accordance with Regulation (EU) No 844/2012 (SANCO/10148/2014 Rev. 4) (For information)

3. EFSA conclusions:
    i. Esfenvalerate
    ii. Cyhalofop-butyl
    iii. Florasulam
4. Draft Review Reports for discussion:
    i. *Flupyrsulfuron-methyl*
    ii. *Thiabendazole*
    iii. *Lambda-cyhalothrin*
    iv. *Acybenzolar-S-methyl*
    v. *Amitrole*
    vi. *Pyridate*
    vii. *Flumioxazin*
    viii. *Sulfosulfuron*
    ix. *Fenhexamid*
    x. *Prosulfuron*
    xi. *Pymetrozine*

A.05 Confirmatory data:

1. Flurochloridone (updated review report to be noted)
2. Tall oil pitch
3. 8-Hydroxiquinoline
4. Etridiazole
5. Dazomet
6. Diflubenzuron
7. Dithianon
8. Dodine
9. AOB

A.06 Amendment of the conditions of approval.

A.07 Basic substances:

1. Pilot projects: state of play
2. New dossiers received
3. EFSA Technical Reports
    i. *Sodium hydrogen carbonate*
4. Draft Review Reports for discussion
    i. *Salix alba*
    ii. *Rheum officinale*
    iii. *Vinegar*
    iv. *Lecithins*
    v. *Artemisia vulgaris*

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

1. Draft Guidance Document on the assessment of certain applications for which reference is made to Article 34 of Regulation (EC) No 1107/2009 (SANCO/11371/2014 Rev. 2) (for discussion)
2. Draft Guidance Document on renewal, withdrawal and amendment of authorisation under Regulation (EC) No 1107/2009 (SANCO/13170/2010 Rev. 10) (for discussion)
3. Draft Guidance Document on the assessment of negligible exposure of an active substance in a plant protection product under realistic conditions of use (points 3.6.3 to 3.6.5, and 3.8.2 of Annex II of Regulation (EC) No 1107/2009) (SANCO/12096/2014) (for information)
4. EFSA Guidance Document on clustering and ranking of emissions of plant protection products and transformation products of these active substances from protected crops (greenhouses and crops grown under cover) to relevant environmental compartments (implementing document SANCO/12184/2014) (to be noted)
5. EFSA Guidance Document on the assessment of exposure of operators, workers, residents and bystanders in risk assessment for plant protection products. EFSA Journal 2014; 12(10):3874 (for discussion)
6. Draft Annexes of the Guidance Document on the presentation and evaluation of dossiers according to Annex III of Directive 91/414/EEC in the format of a (draft) Registration Report - (Doc. SANCO/6895/2009) (for information)

A.09 Notifications under Article 53 of Regulation (EC) No 1107/2009 (to be noted)

A.10 Notifications under Article 44(4) of Regulation (EC) No 1107/2009 (to be noted)

A.11 Notifications under Article 36(3) of Regulation (EC) No 1107/2009 (to be noted)

A.12 Notifications under Article 56 of Regulation (EC) No 1107/2009 (to be noted)

A.13 Sustainable Use Directive (Directive 2009/128/EC):

- State of play

A.14 News from European Food Safety Authority (EFSA)

A.15 Report from working groups:

i. Authorisation database
ii. Low risk

A.16 Bees

1. Review of Neonicotinoids – state of play and next steps
2. EFSA Guidance Document on the risk assessment of plant protection products on bees –and implementation plan (SANCO/10606/2014) state of play
3. Uniform principles – Amendment to the Regulation 546/2011 as regards the trigger value for honeybees to align to the EFSA Guidance Document.
4. AOB

A.17 Court cases

- C-405/12 (and C-404/12) – Judgement in the Appeal to judgment in the case T-338/0 -COM v. Stichting natuur en Milieu & PAN - Internal review under Aarhus Regulation.
- C-108/13 – Judgement in the preliminary ruling in Mac GmbH – Parallel trade of parallel traded plant protection products.
- C-442/14 – Request for a preliminary ruling – Request for access to studies – Questions related to the definition of "emissions into the environment" under Aarhus Regulation.
- T-521/14 - Action for failure to act - Sweden v. COM – adoption of the scientific criteria for the determination of endocrine-disrupting properties pursuant to Regulation 528/2012 on biocidal products.
- T-729/14 – Action for annulment – PAN v. COM - Request to set MRLs for the active substance imidacloprid to protect bee health.

A.18 Endocrine disruptors:

1. State of play
2. Interpretation of the 2nd interim criterion (to be published in the Questions & Answers document)

A.19 19 Minor Uses:

- State of play

A.20 Interpretation issues:

1. Scope of Regulation (EC) No 1107/2009
2. Questions and answers

A.21 Status of harmonised classifications under Regulation (EC) No 1272/2008.

A.22 Glyphosate:

1. State of the dossier
2. Court case T 545-2011

A.23 Chlorpyrifos - state of the dossier.

A.24 Chlorpyrifos-methyl – state of the dossier.

A.25 EFSA Scientific Opinion on the developmental neurotoxicity potential of acetamiprid and imidacloprid.

A.26 Data requirements and acceptance of waivers/implementation of doc. SANCO/10181/2013.

A.27 Follow-up workshop "Harmonisation in Toxicology".

A.28 Imidacloprid revised review report for discussion.

A.29 Note taking procedures.

A.30 Study on the trade of illegal and counterfeit pesticides in the EU – presentation of the draft final report by the Food Chain Evaluation Consortium (FCEC).

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 approving the active substance COS-OGA, 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 Doc. SANTE/00036/2015 Rev. 0)

(B.01_SANTE_00035_2015 Rev. 0)

Legal Basis: Article 13(2) and Article 78(2) 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 approving the active substance cerevisane, 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 doc. SANTE/00033/2015 Rev. 0)

(B.02_SANTE_00032_2015 Rev. 0)

Legal Basis: Article 13(2) 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 Regulation on Implementing Regulation 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 (Doc. SANTE/12589/2013 Rev. 3) (Legal Base: Article 78(2) of Regulation (EC) No 1107/2009) (Opinion of the Committee via the examination procedure)

(B.03_SANCO_12589_2013)

Legal Basis: Article 78(2) of Regulation (EC) No 1107/2009
Procedure: Examination procedure