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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*

20 MARCH 2015

CIRCABC Link: *https://circabc.europa.eu/w/browse/0305ed0c-ad9e-4559-80a4-28e6a0dba8ab*

AGENDA

Section A **Information and/or discussion**

A.01 Summary Reports of previous meetings.

A.02 New active substances:

1. New admissible dossiers (to be noted):

    i. *Oxathiapiprolin*
    ii. *XDE-777 (suggested ISO name: Lyserphenvalpyr)*

2. EFSA conclusions:

    i. *Flumetralin*
    ii. *Rescalure*
    iii. *3-decene-2-one*
    iv. *Flupyradifurone*

3. Commission draft Review Report and Regulation concerning the approval of:

    i. *Cyantraniliprole*
    ii. *Terpenoid blend QRD-460*
    iii. *Pepino mosaic virus CH2*
    iv. *Halauxifen-methyl*

4. *Chromobacterium subtsugae PRAA4-1 (MBI-203)*

A.03 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. State of play Annex I Renewal Project (AIR):

- Next stage of renewal programme

4. EFSA conclusions:

1. Metalaxyl-M
2. Pyraflufen-ethyl

5. Draft Review Reports for discussion:

1. *Flupyrsulfuron-methyl*
2. *Thiabendazole*
3. *Lambda-cyhalothrin*
4. *Acybenzolar-S-methyl*
5. *Amitrole*
6. *Pyridate*
7. *Flumioxazin*
8. *Sulfosulfuron*
9. *Fenhexamid*
10. *Prosulfuron*
11. *Pymetrozine*
12. *Metsulfuron-methyl*
13. *Esfenvalerate*
14. *Florasulam*
15. *Ferric phosphate*

A.04 Confirmatory data:

1. *Tall oil pitch*
2. *Etridiazole*
3. *Dazomet (updated review report to be noted)*
4. *dithianon*
5. *dodine*
6. *Haloxyfop-P*
7. *Chlormequat*
8. *Metamitron (updated review report to be noted)*
9. *Buprofezin*
10. *Pyridaben*
11. *Azoxystrobin (updated review report to be noted)*

12. AOB

A.05 Article 21 Reviews:

1. *Diflubenzuron*
2. *Chlorpyrifos – state of the dossier*

A.06 Amendment of the conditions of approval:

- *Bacillus subtilis QST 713*

A.07 Basic Substances:

1. Pilot projects: state of play
2. New dossiers received
3. EFSA Technical Reports
4. Draft Review Reports for discussion

    i. *Salix alba*
    ii. *Vinegar*
    iii. *Lecithins*
    iv. *Artemisia vulgaris*
    v. *Artemisia absinthium*
    vi. *Tanacetum vulgaris*
    vii. *Fructose*

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 (doc. SANCO/11371/2014 Rev. 4) (to be noted)

2. Draft Guidance Document on the Renewal of Authorisations according to Article 43 of Regulation (EC) No 1107/2009 (doc. SANCO/13170/2010 Rev. 12) (to be noted)

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) (doc. SANCO/12096/2014) (for information)

4. 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)

5. 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) (to be noted)

6. Draft Guidance Document on the Interpretation of the Transitional Measures for the Data Requirements for Chemical Active Substances and Plant Protection Products according to Regulation (EU) No 283/2013 and Regulation (EU) No 284/2013 (SANCO/11509/2013 Rev. 4) (for discussion)

7. Draft Template to be used for the List of Endpoints (SANCO/12483/2014 Rev. 3) (to be noted)

8. Guidance Document for applicants on preparing dossiers for the approval or renewal of approval of microorganisms including viruses according to Regulation (EU) No 283/2013 and Regulation (EU) No 284/2013 (SANCO/ 12545/2014 Rev. 2) (to be noted)

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 the European Food Safety Authority (EFSA).

A.15 Report from working groups:

1. Authorisation database
2. Low risk
3. Post Approval Issues
4. Zonal Workshop
5. OECD-Risk Indicators

A.16 OECD

A.17 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 values for bees to take into account the new scientific development.
4. AOB

A.18 Court cases:

- *T-51/15 – PAN v. Commission - Request to annul a decision of the Commission pertaining to access to some documents*

A.19 Endocrine disruptors:

- State of play

A.20 Minor uses:

- State of play

A.21 Interpretation issues:

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

A.22 Status of harmonised classifications under Regulation (EC) No 1272/2008.

A.23 Glyphosate:

State of the dossier.

A.24 Chlorpyrifos-methyl (revised review report to be noted).

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

A.26 Imidacloprid confirmatory data and review of the aquatic risk assessment (Art 21 Regulation (EC) No 1107/2009 (revised review report for discussion).

A.27 Note taking procedures.

A.28 Dialogue event on risk assessment of active substances in plant protection products.

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 concerning the non-approval of the basic substance Rheum officinale root 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, and amending the Annex to Implementing Regulation (EU) No 540/2011 (Draft Review Report Doc. SANCO/12694/201 Rev. 0)

(B.01_SANCO_12693_2014 Rev. 0)

Legal Basis: Articles 13(2) and 23(5) 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 basic substance Calcium hydroxide 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 (Draft Review Report Doc. SANCO/10148/2015 Rev. 0)

(B.02_SANCO_10147_2015 Rev. 0)

Legal Basis: Articles 13(2) and 23(5) 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 amending Regulation (EU) No 284/2013 as regards the transitional measures applying to procedures concerning plant protection products

(B.03_SANTE_10106_2015 Rev. 0)

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

Miscellaneous

M.01 News from the Food and Veterinary Office (FVO)

M.02 New scientific publications

M.03 AOB