



EUROPEAN COMMISSION

HEALTH & CONSUMERS DIRECTORATE-GENERAL

Brussels,
SANCO D1 (2010) 410396

STANDING COMMITTEE ON THE FOOD CHAIN AND ANIMAL HEALTH

Section *PLANT PROTECTION PRODUCTS – LEGISLATION*

Summary Report 11-12 March 2010

President: W. Reinert

25 Member States were present. Cyprus was absent and represented by Greece, Lithuania was absent and represented by Latvia.

Qualified majority: 255 votes and 14 Member States in favour

A. Points for vote

- 1. Examination and possible opinion on a draft Commission Directive amending Council Directive 91/414/EEC to include sulfuryl fluoride as active substance.** (draft Directive SANCO/10484/2010 Rev. 2; draft review report SANCO/10567/2010 Rev. 0)

Vote postponed

- 2. Examination and possible opinion on a draft Commission Directive amending Council Directive 91/414/EEC to include flonicamid as active substance.** (draft Directive SANCO/10478/2010 Rev. 1; draft review report SANCO /10479/2010 Rev. 2)

The Committee took note of the review report outlined in document SANCO/10479/2010 Rev. 2.

Vote: Unanimous favourable opinion.

- 3. Examination and possible opinion on a draft Commission Directive amending Council Directive 91/414/EEC to include triflumizole as active substance.** (draft Directive SANCO/10456/2010 Rev. 3; draft review report SANCO/10455/2010 Rev. 0).

The Committee took note of the review report outlined in document SANCO/10455/2010 Rev. 0.

Vote: Unanimous favourable opinion.

- 4. Examination and possible opinion on a draft Commission Directive amending Annex I to Council Directive 91/414/EEC as regards the active substances clofentezine, diflubenzuron, lenacil, oxadiazon, picloram and pyriproxyfen.** (draft Directive SANCO/10491/2010 Rev. 0)

Vote postponed

- 5. Examination and possible opinion on a draft Commission Decision of recognising in principle the completeness of the dossier submitted for detailed examination in view of the possible inclusion of 1-4 Dimethylnaphtalene and cyflumetofen in Annex I to Council Directive 91/414/EEC** (SANCO/10463/2010 Rev. 0)

Vote: Unanimous favourable opinion.

- 6. Examination and possible opinion on a draft Commission Directive amending Council Directive 91/414/EEC to include metalaxyl proposal** (draft Directive SANCO/10376/2010 Rev. 1; draft review report SANCO/10476/2010 Rev. 1).

The Committee took note of the review report outlined in document SANCO/10476/2010 Rev. 1.

Vote: Favourable opinion by Qualified Majority (No votes against, 26 MS in favour).

7. Notifications under Article 8(4) of the Directive

Fipronil (NL)

Iprodion (NL)

1.3D (BE)

Imidacloprid, Pencycuron (CZ)

Beauveria brongiartii (DE)

Chlorpyrifos (DE)

Chlorpyrifod-methyl (DE)

Fipronil (DE)

1,3 D (EL) (2)

Abamectin (EL) (2)

Bacillus thuringiensis var. kurstaki (strain ABTS-351)(EL)

Bacillus thuringiensis var. kurstaki (stain PB-54)(EL)

Chlorpyrifos (EL)

Cypermethrin (EL)

Flubendiamide (EL)

Imidacloprid (EL)

Indoxacarb (EL)

Spinosad (EL)

Metaflumizone (EL)

Thiamethoxam (EL)

1,3 D (ES)

Pyraclostrobin (ES)

Piriproxifen (PT)

Clomazone, Metribuzin (PT)

Thiophanate-methyl (PT)

Magnesium phosphide (SE)

Laminarin (SK)

The Committee took note of the documents submitted by BE, CZ, DE, EL, ES, NL, PT, SE, SK.

The Commission recalls that under the provisions of article 8(4) Member States are obliged to inform the Commission and the other Member States immediately after they granted such derogation.

In addition, the Commission pointed out that if an MRL set under Reg. (EC) No 396/2005 cannot be met and a national MRL is set, nevertheless, a consumer risk assessment has to be carried out and forwarded to the Commission, the European Food Safety Authority and Member States.

Member States are reminded that they shall put in place the necessary risk mitigation measures to ensure acceptable uses for human and animal health and the environment.

8. Resubmissions - respect of deadlines for additional report

The Commission reminds Member States to submit in time (at the latest within 6 months), the additional assessment report for re-submitted active substance and, if applied, to inform the Commission and EFSA about additional time requested in case additional information is sought from applicant ("clock-stopping" mechanism). This is essential to ensure that decision making can be finalised in time.

The Commission is determined to follow up any delay in this submission with the appropriate action and will address MS directly in the near future, where necessary.

9. Fipronil – update of the Review Report (confirmatory data) (SANCO/10033/2006 final) (*point to note*)

The Committee took note of the amended review report outlined in document SANCO/10030/2006 final.

10. Triticonazole – update of the Review Report (SANCO/10442/2005 final) (confirmatory data) (*point to note*)

The Committee took note of the amended review report outlined in document SANCO/10442/2005 final.