EUROPEAN COMMISSION

HEALTH AND FOOD SAFETY DIRECTORATE-GENERAL

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Standing Committee on Plants, Animals, Food and Feed
Section *Phytopharmaceuticals - Pesticides Residues*
22 SEPTEMBER 2016 - 23 SEPTEMBER 2016

CIRCABC Link: *https://circabc.europa.eu/w/browse/963ff4fe-0abe-4784-8f9c-8cd8a99ddba2*

AGENDA

Section A Information and/or discussion

A.01 Procedures for routine Maximum Residue Levels (MRL) setting under Regulation (EC) No 396/2005 for Note Taking.

(A.01_SANTE_10595_2015 Rev. 2)

A.02 Amendments to the Extrapolation Guidance Document for Note Taking.

(A.02_SANCO 7525_VI_95 Rev. 10.2)

A.03 Glyphosate-residue definition for Article 12 assessment of MRLs for Note Taking, animal health mandate.

A.04 Update on chlorate.

A.05 Exchange of views of the Committee as regards maximum residue levels for bitertanol, chlormequat and tebufenpyrad (Article 12).

(A.05_SANTE_10827_2016 Rev. 1)

A.06 Exchange of views of the Committee as regards maximum residue levels for fenpyroximate, triadimenol and triadimefon (Article 12).

(A.06_SANTE_10781_2016 Rev. 0)

A.07 Article 12 of Regulation (EC) No 396/2005 procedures:

1. Priorities under Article 12
2. Handling of confirmatory data
3. Footnotes for commercial availability of analytical standards which expired in 2015: Exchange of views of the Committee as regards maximum residue levels for benthiavalicarb, chlorpropham, fenpropidin, pymetrozine and thiobencarb (SANTE//11414/2016 Rev. 0)
4. Substances for which endpoints were changed in Annex I Renewal (AIR) project process after completion of Article 12
5. Other

A.08 Specific substances:

1. Tricyclazole state of play
2. Chlorpyriphos
3. Mercury
4. Amitraz, coumaphos, flumequine, oxytetracycline, permethrin and streptomycin used in Veterinary Medicinal Products
5. New active substances currently under discussion in the Legislation Committee

A.09 Preparation CCPR 49 (2017) (Codex Committee on Pesticide Residues):

1. Priority list: priority of EU nominated substances
2. Concern form quinclorac
3. Other info; e.g. new reporting templates, comments to Codex Circular Letter CL 2016/28-PR (Cereals items)

A.10 Exchange of views of the Committee as regards maximum residue levels for fluopyram, HCH isomers, profenphos and nicotine.

A.11 Maximum residue levels for substances for which LOQs (limits of quantifications) need to be increased in line with the working document on the summing up of LOQs: Exchange of views of the Committee as regards maximum residue levels for bifenazate, daminozide and tolylfluanid.

(A.11_SANTE_11397_2016 Rev. 0)

A.12 Exchange of views of the Committee as regards maximum residues level for achrinathrin, lambda-cyahalothrin, metalaxyl (combined review) and thiabendazole. (Article 12).

(A.12_SANTE_11077_2016 Rev. 0)

A.13 Monitoring:

1. Annual Report 2014 - conclusions on risk assessment
2. Follow up on recommendations
3. Expert Group Meeting on Pesticides Residues Monitoring 2016

A.14 News from the European Food Safety Authority:

1. Progress under Article 12 of Regulation (EC) No 396/2005
2. Progress under Article 10 of Regulation (EC) No 396/2005
3. Update on Article 43 mandates of Regulation (EC) No 396/2005

A.15 Amendments to Annex I to Regulation (EC) No 396/2005 (Regulation (EU) No 752/2014) - state of play.

A.16 Honey guidance.

A.17 Screening exercise on t-MRLs in Regulation (EC) No 396/2005 that will be expiring in 2016 and beginning of 2017.

A.18 Inclusions in Annex IV of Regulation (EC) No 396/2005:

1. State of play of Annex IV inclusions
2. EFSA opinion on Bacillus thuringiensis and follow up

A.19 Commission working document on risk management aspects related to the assessment of cumulative exposure:

- Chapters 3.5.1.1, 3.5.1.2, 3.5.1.6, 3.5.1.7, 3.5.1.8, 3.5.3.3. and 3.5.3.4 for agreement of Member States to ask EFSA and RIVM to implement the described approach
- Chapters 3.1.2.1, 3.5.3.1, 3.5.3.2 and 3.5.4 for agreement of Member States to request EFSA and RIVM to test the described options

(A.19_SANTE_10216_2015 Rev. 6)

A.20 Notifications under Article 18(4) to Regulation (EC) No 396/2005.

A.21 Designation of Member States for maximum residue levels (MRL) applications.

A.22 Information on ongoing work on endocrine disruptors.

A.23 Planned evaluations of Regulation (EC) No 396/2005 and Regulation (EC) No 1107/2009 – State of play.

A.24 Official Food and Feed Control Regulation – areas where delegated/implementing acts will be needed.

A.25 Update on the state of play of MRL setting for biocides.

A.26 Guidance document extraction efficiency.

A.27 Guidance document processing factors.

A.28 AOB:

- Number of residue trials from non- EU countries (UK request)

Section B **Draft(s) presented for an opinion**

B.01 Exchange of views and possible opinion of the Committee on a draft Commission Regulation amending Annexes II, III and IV to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for aminopyralid, azoxystrobin, cyantraniliprole, cyflufenamid, cyproconazole, diethofencarb, dithiocarbamates, fluazifop-P, fluopyram, haloxyfop, isofetamid, metalaxyl, prohexadione, propaquizafop, pyrimethanil, Trichoderma atroviride strain SC1 and zoxamide in or on certain products (Article 10).

(B.01_SANTE_11309_2016 Rev. 1)

Legal Basis: Article 14(1)(a) of Regulation (EC) No 396/2005
Procedure: Regulatory procedure with scrutiny

B.02 Exchange of views and possible opinion of the Committee on a draft Commission Regulation amending Annexes II, III and V to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for bifenthrin, carbetamide, cinidon-ethyl, fenpropimorph and triflusulfuron in or on certain products (Article 12).

(B.02_SANTE 11418_2015 Rev. 1))

Legal Basis: Article 14(1)(a) and Article 49(2) of Regulation (EC) No 396/2005
Procedure: Regulatory procedure with scrutiny