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Health and Food Safety Directorate General

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Standing Committee on Plants, Animals, Food and Feed Section *Phytopharmaceuticals - Residues* 28 - 29 September 2020

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AGENDA

Section A <u>Information and/or discussion</u>

- A.01 Art. 12 and Article 10 of Regulation (EC) No 396/2005 procedures:
 - 1. Priorities under Art. 12 updated table
 - 2. Confirmatory data Art. 12 follow-up
 - a) Outcome of several confirmatory data evaluations by EFSA and proposed follow-up
 - b) Follow up on Article 12 data gaps that were not filled
 - 3. Residue definition for risk assessment
 - 4. Added: amended WD for drafting Art. 12 proposals for Note Taking
 - 5. Overview on import tolerance requests since 2009
 - 6. OECD calculator

A.02 Feedback from Legislation Committee:

1. New active substances currently under discussion in the Legislation Committee.

A.03 Specific substances:

- 1. Glufosinate ammonium
- 2. Glyphosate
- 3. Indolylacetic acid
- 4. Mancozeb
- 5. Imazamox
- 6. Kresosim-methyl
- 7. Chlorothalonil

- 8. Ethirimol correction of MRL
- 9. Carbendazim, benomyl, and thiophanate-methyl: Discussion under agenda item B.1 of the agenda of the section Phytopharmaceuticals, Legislation
- A.04 News from and files related to the European Food Safety Authority:
 - 1. Progress under Article 10 of Regulation (EC) No 396/2005
 - 2. Progress under Article 12 of Regulation (EC) No 396/2005
 - 3. Update on Art. 43 mandates of Regulation (EC) No 396/2005
 - 4. Implementation of the EFSA GD on stereoisomers
 - 5. Discussion on rotational crops (Implementation of OECD Guidelines)
 - 6. Other
- A.05 New Transparency rules.
- A.06 Update of the Communications on data requirements.
- A.07 Monitoring of pesticides residues.
- A.08 Foods for infants and young children.
- A.09 Next steps for cumulative risk assessment.
- A.10 Project on data collection dithiocarbamates.
- A.11 New Official Control Regulation delegated and implementing acts.
- A.12 Screening exercise on temporary MRLs in Regulation (EC) No 396/2005 that will expire in 2020-2021.
- A.13 International Matters:
 - 1. OECD Guidance document on the definition for risk assessment
 - 2. OECD Honey Guidelines
 - 3. Codex Alimentarius/JMPR issues- future work organisation
 - CCPR 2021- working groups and substances
- A.14 SANTE extrapolation guidelines (SANTE-2019-12750), replacement of existing guidance document SANCO 7525/VI/95 Rev. 10.3.
- A.15 Revision of GD SANCO/3029/99 rev. 4 and SANCO/825/00 rev. 8.1 Analytical guidances.
- A.16 Notifications under Article 18(4) to Reg. (EC) No 396/2005.
- A.17 Designation of Member States for maximum residue levels (MRL) applications.

- A.18 Farm to Fork Strategy/REFIT.
- A.19 Overview on substances in forthcoming Art. 12 draft Regulations.
- A.20 Other Information points:
 - Classification of coffee leaves as (novel food) under Annex 1 to Reg. (EC) No 396/2005
 - Update on measures voted in February 2020
 - Readiness and preparedness for the end of the transition period of the UK Withdrawal agreement

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 (EU) No .../...amending Annexes II and IV to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for aclonifen, boscalid, etofenprox, ferric pyrophosphate, L-cysteine, lambda-cyhalothrin, maleic hydrazide, mefentrifluconazole, cow milk, sodium 5-nitroguaiacolate, sodium o-nitrophenolate, sodium p-nitrophenolate and triclopyr in or on certain products (Art. 10).

(SANTE/11426/2020)

Legal Basis: Regulation (EC) No 396/2005 - Articles 5 and 14(1)(a)

Procedure: Regulatory procedure with scrutiny

B.02 Exchange of views and possible opinion of the Committee on a draft Commission Regulation (EU) No .../...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 diclofop, fluopyram, ipconazole and terbuthylazine in or on certain products (Art. 12).

(SANTE/10044/2020)

Legal Basis: Regulation (EC) No 396/2005 - Articles 14(1)(a) and 49(2)

Procedure: Regulatory procedure with scrutiny

B.03 Exchange of views and possible opinion of the Committee on a draft Commission Regulation (EU) No .../...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 fluxapyroxad, hymexazol, metamitron, penflufen and spirotetramat (Art. 12).

(SANTE/10032/2020)

Legal Basis: Regulation (EC) No 396/2005 - Articles 14(1)(a) and 49(2)

Procedure: Regulatory procedure with scrutiny

B.04 Exchange of views and possible opinion of the Committee on a draft Commission Regulation (EU) No .../...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 benalaxyl, benalaxyl-M, dichlobenil, fluopicolide, proquinazid, and pyrdalyl (Art. 12).

(SANTE/10034/2020)

Legal Basis: Regulation (EC) No 396/2005 - Articles 14(1)(a),49(2) and 18(1)(b)

Procedure: Regulatory procedure with scrutiny

B.05 Exchange of views and possible opinion of the Committee on a draft Commission Regulation (EU) No .../...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 carbon tetrachloride, chlorothalonil, chlorpropham, dimethoate, ethoprophos, fenamidone, methiocarb, omethoate, propiconazole and pymetrozine in or on certain products.

(SANTE/10482/2020)

Legal Basis: Regulation (EC) No 396/2005 - Articles 14(1)(a) and 18(1)(b)

Procedure: Regulatory procedure with scrutiny

B.06 Exchange of views and possible opinion of the Committee on a draft Commission Regulation (EU) No .../...amending Annex III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for chlordecone in or on certain products.

(SANTE/12510/2019)

Legal Basis: Regulation (EC) No 396/2005 - Articles 14(1)(a) and 16(1)(a)

Procedure: Regulatory procedure with scrutiny

Section C <u>Draft(s) presented for discussion</u>

C.01 Exchange of views of the Committee on a draft Commission Regulation as regards maximum residue levels for metam, dazomet, hexythiazox, clethodim and sethoxydim (Art. 12).

(SANTE/11220/2019)

Legal Basis: Regulation (EC) No 396/2005 - Articles 14(1)(a), 49(2) and 18 (1)(b)

Procedure: Regulatory procedure with scrutiny

C.02 Exchange of views of the Committee on a draft Commission Regulation as regards maximum residue levels for ametoctradin, bixafen, fenazaquin, spinetoram, tefluthrin and thiencarbazone-methyl in or on certain products (Art. 12).

(SANTE/10706/2020)

Legal Basis: Regulation (EC) No 396/2005 - Articles 14(1)(a) and 49(2)

Procedure: Regulatory procedure with scrutiny

C.03 Exchange of views of the Committee on a draft Commission Implementing Regulation adapting the coordinated multiannual programme for pesticides residues for the years 2021, 2022 and 2023 in view of the withdrawal of the United Kingdom from the Union – adaptation of minimum sample numbers to be taken and analysed by Member State.

(SANTE/11932/2020)

Legal Basis: Regulation (EC) No 396/2005 - Article 29

Procedure: Examination procedure