SCHEDULE 1 Regulation 15

Transitional provisions and savings

PART 1

Interpretation

Interpretation

1. In this Schedule—

"competent authority" and "constituent territory" have the meanings given in Article 3A of Regulation (EC) No 396/2005 (as inserted by regulation 3(5));

"MRLs register" has the meaning given by Article 3(2)(zb) of Regulation (EC) No 396/2005 (as inserted by regulation 3(4)(c)(i)).

PART 2

Existing MRLs etc. under Regulation (EC) No 396/2005

Existing MRL product list

2. The list of products in Parts A and B of Annex 1 to Regulation (EC) No 396/2005 as it had effect immediately before exit day is taken to be the list of products established by each competent authority in relation to its constituent territory in accordance with Article 4(1) of Regulation (EC) No 396/2005.

Existing MRLs under Annexes 2 and 3 to Regulation (EC) No 396/2005

3.—(1) In accordance with this paragraph, an MRL which immediately before exit day is set out in an entry in a table in Annex 2 or 3 to Regulation (EC) No 396/2005 as it has effect immediately before exit day is taken to have been set by each competent authority in relation to its constituent territory—

(a) for an MRL set out in an entry in the table in Annex 2, under Articles 14(1B)(a) and 15(2)(b) of Regulation (EC) No 396/2005;

(b) for an MRL set out in an entry in the table in Part A of Annex 3, under Articles 14(1B)(a) and 15(2)(a)(i) of Regulation (EC) No 396/2005;

(c) for an MRL set out in an entry in the table in Part B of Annex 3, under Articles 14(1B)(a), 15(2)(a)(ii) and 16 of Regulation (EC) No 396/2005.

(2) An MRL to which sub-paragraph (1) applies is taken to apply from the date which it applied from in accordance with a regulation made under Article 14(1) of Regulation (EC) No 396/2005 as it had effect immediately before exit day, except in accordance with sub-paragraph (3).

(3) For the purposes of Article 15(3) of Regulation (EC) No 396/2005, an MRL to which sub-paragraph (1)(b) applies is taken to apply from exit day.

(4) In sub-paragraph (1), "Annex 2" and "Annex 3" means Annex 2 and 3 respectively to Regulation (EC) No 396/2005 as it has effect immediately before exit day and read in accordance with paragraph 4.

(5) Sub-paragraph (6) applies where an MRL to which sub-paragraph (1) applies is immediately before exit day subject to a requirement for the European Commission—

(a) to take specified information into account when reviewing that MRL if the information is submitted by a specified date, or the lack of that information if it is not submitted by that date; or

(b) to take into account the commercial availability of a specified reference standard when reviewing that MRL by a specified date, or the unavailability of it if it is not commercially available by that date.

(6) Where this sub-paragraph applies, on and after exit day the MRL is taken to be subject to the requirement as modified as follows—

(a) a reference in the requirement to the Commission is to be read as a reference to the competent authority;

(b) if the specified date occurred before exit day and on or before that specified date—

(i) the specified information was submitted to the European Commission, the specified date is to be read as 1st January 2020;

(ii) the specified information was not submitted to the European Commission, the requirement is to be read as a requirement on the competent authority to take the lack of that information into account when reviewing the MRL.

(7) For the purposes of sub-paragraphs (5) and (6), a date, a reference standard or information is "specified" if it is specified in the requirement.

Existing MRLs: modifications of Annexes 2 and 3 to Regulation (EC) No 396/2005

4.—(1) For the purposes of paragraph 3, Annexes 2 and 3 of Regulation (EC) No 396/2005 as they had effect immediately before exit day are to be read as follows.

(2) Annex 2 is to be read as if, in the table—

(a) in table footnote (**), for "Annex III Part B" there were substituted "Part 3 of the MRLs register";

(b) in table footnote (1)—

(i) in the first sentence, the words from ", pending" to the end were omitted;

(ii) in the second sentence, ", unless modified by a Regulation" were omitted;

(c) in table footnote (2), in the second sentence "unless modified by a Directive or a Regulation" were omitted;

(d) in table footnote (a)—

(i) after "apply" there were inserted "in relation to a constituent territory";

(ii) for "Annex I" there were substituted "Part 1 of the MRLs register";

(e) in the column for Chlormequat, in entry 0151010 (Table grapes), in table footnote (+), "by a Regulation" were omitted;

(f) in the column for Fipronil, in entry 0211000 ((a) potatoes), entry 1011020 (Fat), entry 1011030 (Liver), entry 1011040 (Kidney), entry 1011050 (Edible offal), entry 1011990 (Others), entry 1012010 (Muscle), entry 1012020 (Fat), entry 1012030 (Liver), entry 1012040 (Kidney), entry 1012050 (Edible offal), entry 1012990 (Others), entry 1013010 (Muscle), entry 1013020 (Fat), entry 1013030 (Liver), entry 1013040 (Kidney), entry 1013050 (Edible offal), entry 1013990 (Others), entry 1014010 (Muscle), entry 1014020 (Fat), entry 1014030 (Liver), entry 1014040 (Kidney), entry 1014050 (Edible offal), entry 1014990 (Others), entry 1015010 (Muscle), entry 1015020 (Fat), entry 1015030 (Liver), entry 1015040 (Kidney), entry 1015050 (Edible offal), entry 1015990 (Others), entry

1016010 (Muscle), entry 1016020 (Fat), entry 1016030 (Liver), entry 1017010 (Muscle), entry 1017020 (Fat), entry 1017030 (Liver), entry 1017040 (Kidney), entry 1017050 (Edible offal), entry 1017990 (Others), entry 1020000 ((ii) Milk), entry 1020010 (Cattle), entry 1020020 (Sheep), entry 1020030 (Goat), entry 1020040 (Horse), entry 1020990 (Others), entry 1030000 ((iii) Bird eggs), entry 1030010 (Chicken), entry 1030020 (Duck), entry 1030030 (Goose), entry 1030040 (Quail), and entry 1030990 (Others), in table footnote (+), "by a Regulation" were omitted;

(g) in the column for Metalaxyl, in entry 0820000 (Fruit spices), entry 0820010 (Allspice/pimento), entry 0820020 (Sichuan pepper), entry 0820030 (Caraway), entry 0820040 (Cardamom), entry 0820050 (Juniper berry), entry 0820060 (Peppercorn), entry 0820070 (Vanilla), entry 0820080 (Tamarind) and entry 0820090 (Others), in table footnote (+), in the second sentence, "by a Regulation" were omitted;

(h) in the column for Mepiquat—

(i) in entry 0280010 (Cultivated fungi), in table footnote (+), in the second sentence, the words from "by a Regulation" to the end were omitted;

(ii) in entry 0401090 (Cotton seeds), in table footnote (+), in the first sentence, "by a Regulation" were omitted;

(i) in the column for Hexachlorobenzene, in entry 0401100 (Pumpkin seeds), in table footnote (+), in the second sentence—

(i) for "Commission" there were substituted "competent authority";

(ii) for "within 10 years from the date of publication" there were substituted "on or before 20th October 2026";

(j) in the column for Cyantraniliprole, in entry 0153010 (Blackberries), in table footnote (+), in the first sentence, "by a Regulation" were omitted.

(3) Annex 3 is to be read as if—

(a) in Part A, in the table—

(i) in table footnote (a)—

(aa) after "apply" there were inserted "in relation to a constituent territory";

(bb) for "Annex I" there were substituted "Part 1 of the MRLs register";

(ii) in table footnote (***), for "Annex III Part B" there were substituted "Part 3 of the MRLs register";

(iii) in table footnote (****), for "Annex V" there were substituted "Part 5 of the MRLs register";

(iv) in the column for Chlorantraniliprole, in entry 0700000 (HOPS), in table footnote (+), "by a Regulation" were omitted;

(v) in the column for Diphenylamine, in entry 0130010 (Apples) and entry 0130020 (Pears), in table footnote (+), in the second sentence—

(aa) for "Commission" there were substituted "competent authority";

(bb) for "within 2 years from the date of publication" there were substituted "on or before 21st January 2018";

(vi) in the column for Mercury compounds, in entry 0120000 (Tree nuts), entry 0256000 ((f) herbs and edible flowers), entry 0280010 (Cultivated funghi), entry 0280020 (Wild funghi), entry 0401000 (Oilseeds), entry 0610000 (Teas), entry 0620000 (Coffee beans), entry 0630000 (Herbal infusions from), entry 0640000 (Cocoa beans), entry 0800000 (SPICES), entry 1011000 ((a) Tissues from swine), entry 1012000 ((b) Tissues from bovine), entry 1013000 ((c) Tissues from sheep), entry

1014000 ((d) Tissues from goat), entry 1015000 ((e) Tissues from equine), entry 1016010 (Muscle (poultry)), entry 1016020 (Fat tissue (poultry)), entry 1016030 (Liver (poultry)), entry 1016040 (Kidney (poultry)), entry 1016050 (Edible offals (poultry)), entry 1017000 ((g) Tissues from other farmed terrestrial animals), entry 1020000 (Milk), entry 1040000 (Honey and other apiculture products), and entry 1070000 (Wild terrestrial vertebrate animals), in table footnote (+)—

(aa) for "Commission" there were substituted "competent authority";

(bb) for "within 10 years from the date of publication" there were substituted "on or before 17th January 2028";

(b) in Part B, in the table, in table footnote (a)—

(i) after "apply" there were inserted "in relation to a constituent territory";

(ii) for "Annex I" there were substituted "Part 1 of the MRLs register".

Existing list of active substances in Annex 4 to Regulation (EC) No 396/2005

5. The list of active substances in Annex 4 to Regulation (EC) No 396/2005 as it had effect immediately before exit day is taken to be the list of active substances established by each competent authority in relation to its constituent territory in accordance with Article 5(1) of Regulation (EC) No 396/2005.

Existing default values for active substances in Annex 5 to Regulation (EC) No 395/2005

6.—(1) In accordance with this paragraph, a default value for an active substance which immediately before exit day is set out in an entry in the table in Annex 5 is taken to have been set by each competent authority in relation to its constituent territory under Article 18A(1) of Regulation (EC) No 396/2005.

(2) Sub-paragraph (3) applies where a default value to which sub-paragraph (1) applies is immediately before exit day subject to a requirement for the European Commission to take specified information into account when reviewing that default value if the information is submitted by a specified date, or the lack of that information if it is not submitted by that date.

(3) Where this sub-paragraph applies, on and after exit day the default value is taken to be subject to the requirement as modified as follows—

(a) a reference in the requirement to the Commission is to be read as a reference to the competent authority;

(b) if the specified date occurred before exit day and on or before that specified date—

(i) the specified information was submitted to the European Commission, the specified date is to be read as 1st January 2020;

(ii) the specified information was not submitted to the European Commission, the requirement is to be read as a requirement on each competent authority to take the lack of that information into account when reviewing the default value.

(4) In sub-paragraph (1), "Annex 5" means Annex 5 to Regulation (EC) No 396/2005 as it has effect immediately before exit day, read as if, in the table—

(a) in table footnote (a)—

(i) after "apply" there were inserted "in relation to a constituent territory";

(ii) for "Annex I" there were substituted "Part 1 of the MRLs register";

(b) in table footnote (**), for "Annex III Part A" there were substituted "Part 3 of the MRLs register";

(5) In sub-paragraphs (2) and (3), "specified" means specified in the requirement.

Existing concentration or dilution factors

7. A concentration or dilution factor set out in Annex 6 to Regulation (EC) No 396/2005 as it had effect immediately before exit day is taken to be specified by each competent authority in relation to its constituent territory in accordance with Article 20(2) of Regulation (EC) No 396/2005.

Existing active substance/product combinations

8.—(1) An active substance/product combination set out in the table in Annex 7 to Regulation (EC) No 396/2005 as it had effect immediately before exit day is taken to be defined by each competent authority in relation to its constituent territory in accordance with Article 18(3A) of Regulation (EC) No 396/2005.

(2) For the purposes of sub-paragraph (1), Annex 7 to Regulation (EC) No 396/2005 as it had effect immediately before exit day is to be read as if, in the heading of the second column of the table in that Annex, the reference to Annex 1 were a reference to Part 1 of the MRLs register.

Existing transitional measures for MRLs under Article 49(2) of Regulation (EC) No 396/2005

9.—(1) Sub-paragraph (2) applies where immediately before exit day a Regulation listed in Schedule 2 includes provision which continues to apply Regulation (EC) No 396/2005 as it had effect before that Regulation applied in respect of the pesticide residue of an active substance in or on one or more products lawfully produced before a specified date (a "transitional measure").

(2) In respect of the pesticide residue and the product or products to which the transitional measure applies, paragraphs 3 to 6 of this Part apply as if a reference in those paragraphs to Regulation (EC) No 396/2005 as it had effect immediately before exit day were a reference to Regulation (EC) No 396/2005 as it had effect before the specified date.

(3) For the purposes of this paragraph, a date is "specified" if it is specified in the transitional measure.

PART 3

Ongoing applications for MRLs

Ongoing applications under Article 7 of Regulation (EC) No 396/2005

10.—(1) This paragraph applies to an application where—

(a) before exit day, that application was submitted under Article 7 of Regulation (EC) No 396/2005 as it had effect immediately before exit day to the United Kingdom for evaluation, and

(b) immediately before exit day, a regulation or decision adopted under Article 14(1) of Regulation (EC) No 396/2005 as it had effect immediately before exit day in relation to that application has not entered into force.

(2) An application to which this paragraph applies is taken as being made under Article 7 of Regulation (EC) No 396/2005 on the date on which it was made, to each competent authority in relation to its constituent territory.

(3) Anything done before exit day in relation to an application to which this paragraph applies—

(a) by the United Kingdom as the evaluating member State;

(b) by the European Food Safety Authority under Article 10 or 11 of Regulation (EC) No 396/2005 as it had effect immediately before exit day;

is taken to have been done by the competent authority.

PART 4

Ongoing MRL assessments

Ongoing EFSA assessments under Article 12 of Regulation (EC) No 396/2005

11.—(1) This paragraph applies where—

(a) before exit day, the European Food Safety Authority is required to provide a reasoned opinion in respect of an active substance, in accordance with Article 12(1) of Regulation (EC) No 396/2005 as it had effect immediately before exit day, and

(b) immediately before exit day, either—

(i) such an opinion has not been provided in accordance with Article 12(1) of Regulation (EC) No 396/2005 as it had effect immediately before exit day, or

(ii) such an opinion has been provided, but a regulation or decision made as a result of that reasoned opinion under Article 14(1) of Regulation (EC) No 396/2005 as it had effect immediately before exit day has not come into force.

(2) A competent authority may produce a reasoned opinion within a period of 36 months beginning with 1st April 2019 in respect of that active substance in relation to its constituent territory, except where sub-paragraph (3) applies.

(3) Where at the end of the 36 month period described in sub-paragraph (2) there are outstanding renewals of authorisations under Article 43 of Regulation (EC) No 1107/2009 relating to that active substance in relation to its constituent territory, a competent authority may instead produce a reasoned opinion before the end of the period of 6 months beginning with the date on which the last of those outstanding renewals is concluded.

(4) Articles 12(3) to (6) and 14 of Regulation (EC) No 396/2005 apply to a reasoned opinion under sub-paragraph (2) as they apply to a reasoned opinion under Article 12(1) of that Regulation.

(5) In providing a reasoned opinion under sub-paragraph (2), the competent authority may also consider relevant information provided by an interested person, including (but not limited to)—

(a) the GAP;

(b) evidence of an authorisation;

(c) relevant assessments undertaken in other countries;

(d) data required by regulations made under Article 8(4) of Regulation (EC) No 1107/2009, including, where appropriate, toxicological data and data on routine analytical methods for use in control laboratories, as well and plant and animal metabolism data.

(6) In sub-paragraph (5), "interested person" includes manufacturers, growers, importers and producers of products listed in a list in Part 1 of the MRLs register in relation to the competent authority's constituent territory.

(7) In this Article, "Regulation (EC) No 1107/2009" means Regulation (EC) No 1107/2009 of the European Parliament and of the Council concerning the placing of plant protection products on the market.

PART 5

Existing technical guidelines

Existing technical guidelines under Article 28(2) of Regulation (EC) No 396/2005

12.—(1) This paragraph applies to technical guidelines which—

(a) before exit day, were adopted in accordance with Article 28(2) of Regulation (EC) No 396/2005 as it had effect immediately before exit day, and

(b) immediately before exit day had not been withdrawn.

(2) Technical guidelines to which this paragraph applies are taken to have been issued by each competent authority in relation to its constituent territory in accordance with Article 46(1) of Regulation (EC) No 396/2005.

PART 6

Control programmes

Saving for existing EU control programme under Article 29 of Regulation (EC) No 396/2005

13.—(1) Despite the amendment of Article 29 of Regulation (EC) No 396/2005 by regulation 7(6), Commission Implementing Regulation (EU) 2018/555 (as amended by regulation 12) continues to have effect, and for the purposes of Sections 2 to 4 of Chapter 5 of Regulation (EC) No 396/2005 is taken to have been prepared by the competent authorities jointly as the United Kingdom control programme for 2019, 2020 and 2021.

(2) The appropriate authority may, by regulations, amend or revoke Commission Implementing Regulation (EU) 2018/555.

(3) In sub-paragraph (2), "the appropriate authority" means—

(a) for regulations applying in relation to England, the Secretary of State;

(b) for regulations applying in relation to Wales, the Welsh Ministers;

(c) for regulations applying in relation to Scotland, the Scottish Ministers;

(d) for regulations applying in relation to Northern Ireland, the Department.

(4) But the appropriate authority is the Secretary of State if consent is given by—

(a) for regulations applying in relation to Wales, the Welsh Ministers;

(b) for regulations applying in relation to Scotland, the Scottish Ministers;

(c) for regulations applying in relation to Northern Ireland, the Department.

(5) In this Part, "the Department" means the Department of Agriculture, Environment and Rural Affairs in Northern Ireland.

Regulations under paragraph 13(2)

14.—(1) Regulations made by the Secretary of State or Welsh Ministers under paragraph 13(2) are to be made by statutory instrument.

(2) For regulations made under paragraph 13(2) by the Scottish Ministers, see section 27 of the Interpretation and Legislative Reform (Scotland) Act 2010.

(3) The power to make regulations conferred on the Department under paragraph 13(2) is exercisable by statutory rule for the purposes of the Statutory Rules (Northern Ireland) Order 1979.

(4) A statutory instrument containing regulations made by the Secretary of State under paragraph 13(2) is subject to annulment in pursuance of a resolution of either House of Parliament.

(5) A statutory instrument containing regulations made by the Welsh Ministers under paragraph 13(2) is subject to annulment in pursuance of a resolution of the National Assembly for Wales.

(6) Regulations made by the Scottish Ministers under paragraph 13(2) are subject to the negative procedure (see section 28 of the Interpretation and Legislative Reform (Scotland) Act 2010).

(7) Regulations made by the Department under paragraph 13(2) are subject to negative resolution within the meaning of section 41(6) of the Interpretation Act (Northern Ireland) Act 1954.

(8) Such regulations may—

(a) contain consequential, incidental, supplementary, transitional or saving provision (including provision amending, repealing or revoking enactments);

(b) make different provision for different purposes.

SCHEDULE 2

Regulation 16

Revocations

1. Commission Regulation (EC) No 645/2000 setting out detailed implementing rules necessary for the proper functioning of certain provisions of Article 7 of Council Directive 86/362/EEC and of Article 4 of Council Directive 90/642/EEC concerning the arrangements for monitoring the maximum levels of pesticide residues in and on cereals and products of plant origin, including fruit and vegetables, respectively.

2. Commission Regulation (EC) No 178/2006 amending Regulation (EC) No 396/2005 of the European Parliament and of the Council to establish Annex I listing the food and feed products to which maximum levels for pesticide residues apply.

3. Commission Regulation (EC) No 149/2008 amending Regulation (EC) No 396/2005 of the European Parliament and of the Council by establishing Annexes II, III and IV setting maximum residue levels for products covered by Annex I thereto.

4. Commission Regulation (EC) No 260/2008 amending Regulation (EC) No 396/2005 of the European Parliament and of the Council by establishing Annex VII listing active substance/product combinations covered by a derogation as regards post harvest treatments with a fumigant.

5. Commission Regulation (EC) No 839/2008 amending Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards Annexes II, III and IV on maximum residue levels of pesticides in or on certain products.

6. Commission Regulation (EC) No 256/2009 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for azoxystrobin and fludioxonil in or on certain products.

7. Commission Regulation (EC) No 822/2009 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 azoxystrobin, atrazine, chlormequat, cyprodinil, dithiocarbamates, fludioxonil, fluroxypyr, indoxacarb, mandipropamid, potassium tri-iodide, spirotetramat, tetraconazole, and thiram in or on certain products.

8. Commission Regulation (EC) No 1050/2009 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for azoxystrobin, acetamiprid, clomazone, cyflufenamid, emamectin benzoate, famoxadone, fenbutatin

oxide, flufenoxuron, fluopicolide, indoxacarb, ioxynil, mepanipyrim, prothioconazole, pyridalyl, thiacloprid and trifloxystrobin in or on certain products.

9. Commission Regulation (EC) No 1097/2009 amending Annex II to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for dimethoate, ethephon, fenamiphos, fenarimol, methamidophos, methomyl, omethoate, oxydemeton-methyl, procymidone, thiodicarb and vinclozolin in or on certain products.

10. Commission Regulation (EU) No 304/2010 amending Annex II to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for 2-phenylphenol in or on certain products.

11. Commission Regulation (EU) No 459/2010 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 certain pesticides in or on certain products.

12. Commission Regulation (EU) No 600/2010 amending Annex I to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards additions and modification of the examples of related varieties or other products to which the same MRL applies.

13. Commission Regulation (EU) No 750/2010 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for certain pesticides in or on certain products.

14. Commission Regulation (EU) No 765/2010 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for chlorothalonil clothianidin, difenoconazole, fenhexamid, flubendiamide, nicotine, spirotetramat, thiacloprid and thiamethoxam in or on certain products.

15. Commission Regulation (EU) No 893/2010 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for acequinocyl, bentazone, carbendazim, cyfluthrin, fenamidone, fenazaquin, flonicamid, flutriafol, imidacloprid, ioxynil, metconazole, prothioconazole, tebufenozide and thiophanate-methyl in or on certain products.

16. Commission Regulation (EU) No 310/2011 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for aldicarb, bromopropylate, chlorfenvinphos, endosulfan, EPTC, ethion, fenthion, fomesafen, methabenzthiazuron, methidathion, simazine, tetradifon and triforine in or on certain products.

17. Commission Regulation (EU) No 460/2011 amending Annex III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards the maximum residue level for chlorantraniliprole (DPX E-2Y45) in or on carrots.

18. Commission Regulation (EU) No 508/2011 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for abamectin, acetamiprid, cyprodinil, difenoconazole, dimethomorph, fenhexamid, proquinazid, prothioconazole, pyraclostrobin, spirotetramat, thiacloprid, thiamethoxam and trifloxystrobin in or on certain products.

19. Commission Regulation (EU) No 520/2011 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for benalaxyl, boscalid, buprofezin, carbofuran, carbosulfan, cypermethrin, fluopicolide, hexythiazox, indoxacarb, metaflumizone, methoxyfenozide, paraquat, prochloraz, spirodiclofen, prothioconazole and zoxamide in or on certain products.

20. Commission Regulation (EU) No 524/2011 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for

biphenyl, deltamethrin, ethofumesate, isopyrazam, propiconazole, pymetrozine, pyrimethanil and tebuconazole in or on certain products.

21. Commission Regulation (EU) No 559/2011 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for captan, carbendazim, cyromazine, ethephon, fenamiphos, thiophanate-methyl, triasulfuron and triticonazole in or on certain products.

22. Commission Regulation (EU) No 812/2011 amending Annex III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for dimethomorph, fluopicolide, mandipropamid, metrafenone, nicotine and spirotetramat in or on certain products.

23. Commission Regulation (EU) No 813/2011 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for acequinocyl, emamectin benzoate, ethametsulfuron-methyl, flubendiamide, fludioxonil, kresoxim-methyl, methoxyfenozide, novaluron, thiacloprid and trifloxystrobin in or on certain products.

24. Commission Regulation (EU) No 978/2011 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for acetamiprid, biphenyl, captan, chlorantraniliprole, cyflufenamid, cymoxanil, dichlorprop-P, difenoconazole, dimethomorph, dithiocarbamates, epoxiconazole, ethephon, flutriafol, fluxapyroxad, isopyrazam, propamocarb, pyraclostrobin, pyrimethanil and spirotetramat in or on certain products.

25. Commission Regulation (EU) No 270/2012 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for amidosulfuron, azoxystrobin, bentazone, bixafen, cyproconazole, fluopyram, imazapic, malathion, propiconazole and spinosad in or on certain products.

26. Commission Regulation (EU) No 322/2012 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for clopyralid, dimethomorph, fenpyrazamine, folpet and pendimethalin in or on certain products.

27. Commission Regulation (EU) No 441/2012 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for bifenazate, bifenthrin, boscalid, cadusafos, chlorantraniliprole, chlorothalonil, clothianidin, cyproconazole, deltamethrin, dicamba, difenoconazole, dinocap, etoxazole, fenpyroximate, flubendiamide, fludioxonil, glyphosate, metalaxyl-M, meptyldinocap, novaluron, thiamethoxam, and triazophos in or on certain products.

28. Commission Regulation (EU) No 473/2012 amending Annex III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for spinetoram (XDE-175) in or on certain products.

29. Commission Regulation (EU) No 556/2012 amending Annex III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for spinosad in or on raspberries.

30. Commission Regulation (EU) No 592/2012 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for bifenazate, captan, cyprodinil, fluopicolide, hexythiazox, isoprothiolane, metaldehyde, oxadixyl and phosmet in or on certain products.

31. Commission Regulation (EU) No 897/2012 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for acibenzolar-S-methyl, amisulbrom, cyazofamid, diflufenican, dimoxystrobin, methoxyfenozide and nicotine in or on certain products.

32. Commission Regulation (EU) No 899/2012 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for acephate, alachlor, anilazine, azocyclotin, benfuracarb, butylate, captafol, carbaryl, carbofuran, carbosulfan, chlorfenapyr, chlorthal-dimethyl, chlorthiamid, cyhexatin, diazinon, dichlobenil, dicofol, dimethipin, diniconazole, disulfoton, fenitrothion, flufenzin, furathiocarb, hexaconazole, lactofen, mepronil, methamidophos, methoprene, monocrotophos, monuron, oxycarboxin, oxydemeton-methyl, parathion-methyl, phorate, phosalone, procymidone, profenofos, propachlor, quinclorac, quintozene, tolylfluanid, trichlorfon, tridemorph and trifluralin in or on certain products and amending that Regulation by establishing Annex V listing default values.

33. Commission Regulation (EU) No 34/2013 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 2-phenylphenol, ametoctradin, *Aureobasidium pullulans* strains DSM 14940 and DSM 14941, cyproconazole, difenoconazole, dithiocarbamates, folpet, propamocarb, spinosad, spirodiclofen, tebufenpyrad and tetraconazole in or on certain products.

34. Commission Regulation (EU) No 35/2013 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for dimethomorph, indoxacarb, pyraclostrobin and trifloxystrobin in or on certain products.

35. Commission Regulation (EU) No 212/2013 replacing Annex I to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards additions and modifications with respect to the products covered by that Annex.

36. Commission Regulation (EU) No 241/2013 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for chlorantraniliprole, fludioxonil and prohexadione in or on certain products.

37. Commission Regulation (EU) No 251/2013 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for aminopyralid, bifenazate, captan, fluazinam, fluopicolide, folpet, kresoxim-methyl, penthiopyrad, proquinazid, pyridate and tembotrione in or on certain products.

38. Commission Regulation (EU) No 293/2013 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for emamectin benzoate, etofenprox, etoxazole, flutriafol, glyphosate, phosmet, pyraclostrobin, spinosad and spirotetramat in or on certain products.

39. Commission Regulation (EU) No 500/2013 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 acetamiprid, *Adoxophyes orana granulovirus* strain BV-0001, azoxystrobin, clothianidin, fenpyrazamine, heptamaloxyloglucan, metrafenone, *Paecilomyces lilacinus* strain 251, propiconazole, quizalofop-P, spiromesifen, tebuconazole, thiamethoxam and *zucchini yellow mosaik virus* - weak strain in or on certain products.

40. Commission Regulation (EU) No 668/2013 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for 2,4-DB, dimethomorph, indoxacarb, and pyraclostrobin in or on certain products.

41. Commission Regulation (EU) No 772/2013 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 diphenylamine in or on certain products.

42. Commission Regulation (EU) No 777/2013 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 clodinafop, clomazone, diuron, ethalfluralin, ioxynil, iprovalicarb, maleic hydrazide, mepanipyrim, metconazole, prosulfocarb and tepraloxydim in or on certain products.

43. Commission Regulation (EU) No 834/2013 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for acequinocyl, bixafen, diazinon, difenoconazole, etoxazole, fenhexamid, fludioxonil, isopyrazam, lambda-cyhalothrin, profenofos and prothioconazole in or on certain products.

44. Commission Regulation (EU) No 1004/2013 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for 8-hydroxyquinoline, cyproconazole, cyprodinil, fluopyram, nicotine, pendimethalin, penthiopyrad and trifloxystrobin in or on certain products.

45. Commission Regulation (EU) No 1138/2013 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 bitertanol, chlorfenvinphos, dodine and vinclozolin in or on certain products.

46. Commission Regulation (EU) No 1317/2013 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 2,4-D, beflubutamid, cyclanilide, diniconazole, florasulam, metolachlor and S-metolachlor, and milbemectin in or on certain products.

47. Commission Regulation (EU) No 36/2014 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for aminopyralid, chlorantraniliprole, cyflufenamid, mepiquat, metalaxyl-M, propamocarb, pyriofenone and quinoxyfen in or on certain products.

48. Commission Regulation (EU) No 51/2014 amending Annex II to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for dimethomorph, indoxacarb and pyraclostrobin in or on certain products.

49. Commission Regulation (EU) No 61/2014 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for cyromazine, fenpropidin, formetanate, oxamyl and tebuconazole in or on certain products.

50. Commission Regulation (EU) No 79/2014 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 bifenazate, chlorpropham, esfenvalerate, fludioxonil and thiobencarb in or on certain products.

51. Commission Regulation (EU) No 87/2014 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 acetamiprid, butralin, chlorotoluron, daminozide, isoproturon, picoxystrobin, pyrimethanil and trinexapac in or on certain products.

52. Commission Regulation (EU) No 289/2014 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 foramsulfuron, azimsulfuron, iodosulfuron, oxasulfuron, mesosulfuron, flazasulfuron, imazosulfuron, propamocarb, bifenazate, chlorpropham and thiobencarb in or on certain products.

53. Commission Regulation (EU) No 318/2014 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for fenarimol, metaflumizone and teflubenzuron in or on certain products.

54. Commission Regulation (EU) No 364/2014 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for fenpyroximate, flubendiamide, isopyrazam, kresoxim-methyl, spirotetramat and thiacloprid in or on certain products.

55. Commission Regulation (EU) No 398/2014 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for benthiavalicarb, cyazofamid, cyhalofop-butyl, forchlorfenuron, pymetrozine and silthiofam in or on certain products.

56. Commission Regulation (EU) No 491/2014 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for ametoctradin, azoxystrobin, cycloxydim, cyfluthrin, dinotefuran, fenbuconazole, fenvalerate, fludioxonil, fluopyram, flutriafol, fluxapyroxad, glufosinate-ammonium, imidacloprid, indoxacarb, MCPA, methoxyfenozide, penthiopyrad, spinetoram and trifloxystrobin in or on certain products.

57. Commission Regulation (EU) No 588/2014 amending Annexes III and IV to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for orange oil, *Phlebiopsis gigantea*, gibberellic acid, *Paecilomyces fumosoroseus* strain FE 9901, *Spodoptera littoralis* nucleopolyhedrovirus, *Spodoptera exigua* nuclear polyhedrosis virus, *Bacillus firmus* I-1582, s-abscisic acid, L-ascorbic acid and *Helicoverpa armigera* nucleopolyhedrovirus in or on certain products.

58. Commission Regulation (EU) No 617/2014 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for ethoxysulfuron, metsulfuron-methyl, nicosulfuron, prosulfuron, rimsulfuron, sulfosulfuron and thifensulfuron-methyl in or on certain products.

59. Commission Regulation (EU) No 703/2014 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 acibenzolar-S-methyl, ethoxyquin, flusilazole, isoxaflutole, molinate, propoxycarbazone, pyraflufen-ethyl, quinoclamine and warfarin in or on certain products.

60. Commission Regulation (EU) No 737/2014 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for 2-phenylphenol, chlormequat, cyflufenamid, cyfluthrin, dicamba, fluopicolide, flutriafol, fosetyl, indoxacarb, isoprothiolane, mandipropamid, metaldehyde, metconazole, phosmet, picloram, propyzamide, pyriproxyfen, saflufenacil, spinosad and trifloxystrobin in or on certain products.

61. Commission Regulation (EU) No 752/2014 replacing Annex I to Regulation (EC) No 396/2005 of the European Parliament and of the Council.

62. Commission Regulation (EU) No 991/2014 amending Annex III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for fosetyl in or on certain products.

63. Commission Regulation (EU) No 1096/2014 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 carbaryl, procymidone and profenofos in or on certain products.

64. Commission Regulation (EU) No 1119/2014 amending Annex III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for benzalkonium chloride and didecyldimethylammonium chloride in or on certain products.

65. Commission Regulation (EU) No 1126/2014 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 asulam, cyanamide, dicloran, flumioxazin, flupyrsulfuron-methyl, picolinafen and propisochlor in or on certain products.

66. Commission Regulation (EU) No 1127/2014 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for amitrole, dinocap, fipronil, flufenacet, pendimethalin, propyzamide, and pyridate in or on certain products.

67. Commission Regulation (EU) No 1146/2014 amending Annexes II, III, IV and V to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for anthraquinone, benfluralin, bentazone, bromoxynil, chlorothalonil, famoxadone, imazamox, methyl bromide, propanil and sulphuric acid in or on certain products.

68. Commission Regulation (EU) 2015/165 amending Annex IV to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for lactic acid, *Lecanicillium muscarium* strain Ve6, chitosan hydrochloride and *Equisetum arvense* L. in or on certain products.

69. Commission Regulation (EU) 2015/399 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 1,4-dimethylnaphthalene, benfuracarb, carbofuran, carbosulfan, ethephon, fenamidone, fenvalerate, fenhexamid, furathiocarb, imazapyr, malathion, picoxystrobin, spirotetramat, tepraloxydim and trifloxystrobin in or on certain products.

70. Commission Regulation (EU) 2015/400 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 bone oil, carbon monoxide, cyprodinil, dodemorph, iprodione, metaldehyde, metazachlor, paraffin oil (CAS 64742-54-7), petroleum oils (CAS 92062-35-6) and propargite in or on certain products.

71. Commission Regulation (EU) 2015/401 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for acetamiprid, chromafenozide, cyazofamid, dicamba, difenoconazole, fenpyrazamine, fluazinam, formetanate, nicotine, penconazole, pymetrozine, pyraclostrobin, tau-fluvalinate and tebuconazole in or on certain products.

72. Commission Regulation (EU) 2015/552 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 1,3-dichloropropene, bifenox, dimethenamid-P, prohexadione, tolylfluanid and trifluralin in or on certain products.

73. Commission Regulation (EU) 2015/603 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 2-naphthyloxyacetic acid, acetochlor, chloropicrin, diflufenican, flurprimidol, flutolanil and spinosad in or on certain products.

74. Commission Regulation (EU) 2015/845 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for azoxystrobin, chlorantraniliprole, cyantraniliprole, dicamba, difenoconazole, fenpyroximate, fludioxonil, glufosinate-ammonium, imazapic, imazapyr, indoxacarb, isoxaflutole, mandipropamid, penthiopyrad, propiconazole, pyrimethanil, spirotetramat and trinexapac in or on certain products.

75. Commission Regulation (EU) 2015/846 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for acetamiprid, ametoctradin, amisulbrom, bupirimate, clofentezine, ethephon, ethirimol, fluopicolide, imazapic, propamocarb, pyraclostrobin and tau-fluvalinate in or on certain products.

76. Commission Regulation (EU) 2015/868 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 2,4,5-T, barban, binapacryl, bromophos-ethyl, camphechlor (toxaphene), chlorbufam, chloroxuron, chlozolinate, DNOC, di-allate, dinoseb, dinoterb, dioxathion, ethylene oxide, fentin acetate, fentin hydroxide, flucycloxuron, flucythrinate, formothion, mecarbam, methacrifos, monolinuron, phenothrin, propham, pyrazophos, quinalphos, resmethrin, tecnazene and vinclozolin in or on certain products.

77. Commission Regulation (EU) 2015/896 amending Annex IV to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for *Trichoderma polysporum* strain IMI 206039, *Trichoderma asperellum* (formerly *T. harzianum*) strains ICC012, T25 and TV1, *Trichoderma atroviride* (formerly *T. harzianum*) strains IMI 206040 and T11, *Trichoderma harzianum* strains T-22 and ITEM 908, *Trichoderma gamsii* (formerly *T. viride*) strain ICC080, *Trichoderma asperellum* (strain T34), *Trichoderma atroviride* strain I-1237,

geraniol, thymol, sucrose, ferric sulphate (iron (III) sulphate), ferrous sulphate (iron (II) sulphate) and folic acid in or on certain products.

78. Commission Regulation (EU) 2015/1040 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 azoxystrobin, dimoxystrobin, fluroxypyr, methoxyfenozide, metrafenone, oxadiargyl and tribenuron in or on certain products.

79. Commission Regulation (EU) 2015/1101 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for difenoconazole, fluopicolide, fluopyram, isopyrazam and pendimethalin in or on certain products.

80. Commission Regulation (EU) 2015/1200 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for amidosulfuron, fenhexamid, kresoxim-methyl, thiacloprid and trifloxystrobin in or on certain products.

81. Commission Regulation (EU) 2015/1608 amending Annex IV to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for capric acid, paraffin oil (CAS 64742-46-7), paraffin oil (CAS 72623-86-0), paraffin oil (CAS 8042-47-5), paraffin oil (CAS 97862-82-3), lime sulphur and urea in or on certain products.

82. Commission Regulation (EU) 2015/1910 amending Annexes III and V to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for guazatine in or on certain products.

83. Commission Regulation (EU) 2015/2075 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for abamectin, desmedipham, dichlorprop-P, haloxyfop-P, oryzalin and phenmedipham in or on certain products.

84. Commission Regulation (EU) 2016/1 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for bifenazate, boscalid, cyazofamid, cyromazine, dazomet, dithiocarbamates, fluazifop-P, mepanipyrim, metrafenone, picloram, propamocarb, pyridaben, pyriofenone, sulfoxaflor, tebuconazole, tebufenpyrad and thiram in or on certain products.

85. Commission Regulation (EU) 2016/46 amending Annex III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for oxadixyl and spinetoram in or on certain products.

86. Commission Regulation (EU) 2016/53 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for diethofencarb, mesotrione, metosulam and pirimiphos-methyl in or on certain products.

87. Commission Regulation (EU) 2016/60 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for chlorpyrifos in or on certain products.

88. Commission Regulation (EU) 2016/67 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 ametoctradin, chlorothalonil, diphenylamine, flonicamid, fluazinam, fluoxastrobin, halauxifen-methyl, propamocarb, prothioconazole, thiacloprid and trifloxystrobin in or on certain products.

89. Commission Regulation (EU) 2016/71 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 1-methylcyclopropene, flonicamid, flutriafol, indolylacetic acid, indolylbutyric acid, pethoxamid, pirimicarb, prothioconazole and teflubenzuron in or on certain products.

90. Commission Regulation (EU) 2016/75 amending Annex III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for fosetyl in or on certain products.

91. Commission Regulation (EU) 2016/143 amending Annex IV to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards COS-OGA, cerevisane, calcium hydroxide, lecithins, *Salix* spp cortex, vinegar, fructose, *Pepino mosaic* virus strain CH2 isolate 1906, *Verticillium albo-atrum* isolate WCS850 and *Bacillus amyloliquefaciens* subsp. *plantarum* strain D747.

92. Commission Regulation (EU) 2016/156 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for boscalid, clothianidin, thiamethoxam, folpet and tolclofos-methyl in or on certain products.

93. Commission Regulation (EU) 2016/439 amending Annex IV to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards *Cydia pomonella* Granulovirus (CpGV), calcium carbide, potassium iodide, sodium hydrogen carbonate, rescalure and *Beauveria bassiana* strain ATCC 74040 and *Beauveria bassiana* strain GHA.

94. Commission Regulation (EU) 2016/440 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 atrazine in or on certain products.

95. Commission Regulation (EU) 2016/452 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for captan, propiconazole and spiroxamine in or on certain products.

96. Commission Regulation (EU) 2016/486 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for cyazofamid, cycloxydim, difluoroacetic acid, fenoxycarb, flumetralin, fluopicolide, flupyradifurone, fluxapyroxad, kresoxim-methyl, mandestrobin, mepanipyrim, metalaxyl-M, pendimethalin and tefluthrin in or on certain products.

97. Commission Regulation (EU) 2016/567 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for chlorantraniliprole, cyflumetofen, cyprodinil, dimethomorph, dithiocarbamates, fenamidone, fluopyram, flutolanil, imazamox, metrafenone, myclobutanil, propiconazole, sedaxane and spirodiclofen in or on certain products.

98. Commission Regulation (EU) 2016/805 amending Annex IV to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards *Streptomyces* K61 (formerly *S. griseoviridis), Candida oleophila* strain O, FEN 560 (also called fenugreek or fenugreek seed powder), methyl decanoate (CAS 110-42-9), methyl octanoate (CAS 111-11-5) and terpenoid blend QRD 460.

99. Commission Regulation (EU) 2016/1002 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 AMTT, diquat, dodine, glufosinate and tritosulfuron in or on certain products.

100. Commission Regulation (EU) 2016/1003 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for abamectin, acequinocyl, acetamiprid, benzovindiflupyr, bromoxynil, fludioxonil, fluopicolide, fosetyl, mepiquat, proquinazid, propamocarb, prohexadione and tebuconazole in or on certain products.

101. Commission Regulation (EU) 2016/1015 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for

1-naphthylacetamide, 1-naphthylacetic acid, chloridazon, fluazifop-P, fuberidazole, mepiquat and tralkoxydim in or on certain products.

102. Commission Regulation (EU) 2016/1016 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for ethofumesate, etoxazole, fenamidone, fluoxastrobin and flurtamone in or on certain products.

103. Commission Regulation (EU) 2016/1355 amending Annex II to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards thiacloprid.

104. Commission Regulation (EU) 2016/1726 amending Annex IV to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards carvone, diammonium phosphate, *Saccharomyces cerevisiae* strain LAS02 and whey.

105. Commission Regulation (EU) 2016/1785 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for cymoxanil, phosphane and phosphide salts and sodium 5-nitroguaiacolate, sodium o-nitrophenolate and sodium p-nitrophenolate in or on certain products.

106. Commission Regulation (EU) 2016/1822 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 aclonifen, deltamethrin, fluazinam, methomyl, sulcotrione and thiodicarb in or on certain products.

107. Commission Regulation (EU) 2016/1866 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 3-decen-2-one, acibenzolar-S-methyl and hexachlorobenzene in or on certain products.

108. Commission Regulation (EU) 2016/1902 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for acetamiprid, ametoctradin, azoxystrobin, cyfluthrin, difluoroacetic acid, dimethomorph, fenpyrazamine, flonicamid, fluazinam, fludioxonil, flupyradifurone, flutriafol, fluxapyroxad, metconazole, proquinazid, prothioconazole, pyriproxyfen, spirodiclofen and trifloxystrobin in or on certain products.

109. Commission Regulation (EU) 2017/170 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.

110. Commission Regulation (EU) 2017/171 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.

111. Commission Regulation (EU) 2017/405 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for sulfoxaflor in or on certain products.

112. Commission Regulation (EU) 2017/623 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for acequinocyl, amitraz, coumaphos, diflufenican, flumequine, metribuzin, permethrin, pyraclostrobin and streptomycin in or on certain products.

113. Commission Regulation (EU) 2017/624 amending Annexes II and V to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for bifenazate, daminozide and tolylfluanid in or on certain products.

114. Commission Regulation (EU) 2017/626 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for acetamiprid, cyantraniliprole, cypermethrin, cyprodinil, difenoconazole, ethephon, fluopyram, flutriafol, fluxapyroxad, imazapic, imazapyr, lambda-cyhalothrin, mesotrione, profenofos, propiconazole, pyrimethanil, spirotetramat, tebuconazole, triazophos and trifloxystrobin in or on certain products.

115. Commission Regulation (EU) 2017/627 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 fenpyroximate, triadimenol and triadimefon in or on certain products.

116. Commission Implementing Regulation (EU) 2017/660 concerning a coordinated multiannual control programme of the Union for 2018, 2019 and 2020 to ensure compliance with maximum residue levels of pesticides and to assess the consumer exposure to pesticide residues in and on food of plant and animal origin.

117. Commission Regulation (EU) 2017/671 amending Annex II to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for clothianidin and thiamethoxam in or on certain products.

118. Commission Regulation (EU) 2017/693 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 bitertanol, chlormequat and tebufenpyrad in or on certain products.

119. Commission Regulation (EU) 2017/978 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 fluopyram; hexachlorocyclohexane (HCH), alpha-isomer; hexachlorocyclohexane (HCH), beta-isomer; hexachlorocyclohexane (HCH), sum of isomers, except the gamma isomer; lindane (hexachlorocyclohexane (HCH), gamma-isomer); nicotine and profenofos in or on certain products.

120. Commission Regulation (EU) 2017/983 amending Annexes III and V to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for tricyclazole in or on certain products.

121. Commission Regulation (EU) 2017/1016 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 benzovindiflupyr, chlorantraniliprole, deltamethrin, ethofumesate, haloxyfop, Mild Pepino Mosaic Virus isolate VC1, Mild Pepino Mosaic Virus isolate VX1, oxathiapiprolin, penthiopyrad, pyraclostrobin, spirotetramat, sunflower oil, tolclofos-methyl and trinexapac in or on certain products.

122. Commission Regulation (EU) 2017/1135 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for dimethoate and omethoate in or on certain products.

123. Commission Regulation (EU) 2017/1164 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for acrinathrin, metalaxyl and thiabendazole in or on certain products.

124. Commission Regulation (EU) 2017/1777 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 *Bacillus amyloliquefaciens* strain FZB24, *Bacillus amyloliquefaciens* strain MBI 600, clayed charcoal, dichlorprop-P, ethephon, etridiazole, flonicamid, fluazifop-P, hydrogen peroxide, metaldehyde, penconazole, spinetoram, tau-fluvalinate and *Urtica* spp. in or on certain products.

125. Commision Regulation (EU) 2018/62 replacing Annex I to Regulation (EC) No 396/2005 of the European Parliament and of the Council.

126. Commission Regulation (EU) 2018/70 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 ametoctradin, chlorpyrifos-methyl, cyproconazole, difenoconazole, fluazinam, flutriafol, prohexadione and sodium chloride in or on certain products.

127. Commission Regulation (EU) 2018/73 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for mercury compounds in or on certain products.

128. Commission Regulation (EU) 2018/78 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for 2-phenylphenol, bensulfuron-methyl, dimethachlor and lufenuron in or on certain products.

129. Commission Regulation (EU) 2018/685 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 abamectin, beer, fluopyram, fluxapyroxad, maleic hydrazide, mustard seeds powder and tefluthrin in or on certain products.

130. Commission Regulation (EU) 2018/686 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for chlorpyrifos, chlorpyrifos-methyl and triclopyr in or on certain products.

131. Commission Regulation (EU) 2018/687 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for acibenzolar-S-methyl, benzovindiflupyr, bifenthrin, bixafen, chlorantraniliprole, deltamethrin, flonicamid, fluazifop-P, isofetamid, metrafenone, pendimethalin and teflubenzuron in or on certain products.

132. Commission Regulation (EU) 2018/832 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 cyantraniliprole, cymoxanil, deltamethrin, difenoconazole, fenamidone, flubendiamide, fluopicolide, folpet, fosetyl, mandestrobin, mepiquat, metazachlor, propamocarb, propargite, pyrimethanil, sulfoxaflor and trifloxystrobin in or on certain products.

133. Commission Regulation (EU) 2018/960 amending Annexes II and III to Regulation (EC) No 396/2005 of the European Parliament and of the Council as regards maximum residue levels for lambda-cyhalothrin in or on certain products.

134. Commission Regulation (EU) 2018/1049 amending Annex I to Regulation (EC) No 396/2005 of the European Parliament and of the Council.

135. Commission Regulation (EU) 2018/1514 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 abamectin, acibenzolar-S-methyl, clopyralid, emamectin, fenhexamid, fenpyrazamine, fluazifop-P, isofetamid, *Pasteuria nishizawae* Pn1, talc E553B and tebuconazole in or on certain products.