
STATUTORY INSTRUMENTS

2019 No. 676

The Veterinary Medicines and Animals and Animal Products (Examination of Residues and Maximum Residue Limits) (Amendment etc.) (EU Exit) Regulations 2019

PART 4

Amendment and revocation of retained direct EU legislation

Commission Regulation (EU) 2018/782

10.—(1) Commission Regulation (EU) 2018/782 establishing the methodological principles for the risk assessment and risk management recommendations referred to in Regulation (EC) No 470/2009 is amended as follows.

- (2) In Article 1—
- (a) for “Agency” substitute “appropriate authority”;
 - (b) for “preparing opinions on the” substitute “producing an assessment report in respect of”.
- (3) After Article 3, omit the words from “This Regulation” to “Member States”.
- (4) In Annex 1—
- (a) in paragraph 1.1, for the words from “provisions related” to “Council” substitute “the Good Laboratory Practice Regulations 1999(1)”;
 - (b) in paragraph 1.2, for “Directive 2010/63/EU of the European Parliament and of the Council” substitute “the Animals (Scientific Procedures) Act 1986(2)”;
 - (c) in paragraph 1.7, in the second subparagraph—
 - (i) for “European Medicines Agency (‘Agency’)” substitute “appropriate authority”;
 - (ii) for “the Agency” substitute “the appropriate authority”;
 - (d) in paragraph 1.8, for the words from “the Agency’s” to the end substitute “guidance issued by the appropriate authority”;
 - (e) in paragraph 2.4.2(l), omit “Agency and other”;
 - (f) in paragraph 2.4.2(m), for “Directive 2010/63/EU” substitute “the Animals (Scientific Procedures) Act 1986”;
 - (g) in paragraph 2.6.1.3, for “Agency” substitute “appropriate authority”;
 - (h) in paragraph 3.2.2(j), omit “Agency and other”;
 - (i) in paragraph 3.2.2(k), for “Directive 2010/63/EU” substitute “the Animals (Scientific Procedures) Act 1986”;
 - (j) in paragraph 3.5.5—

(1) S.I. 1999/3106, amended by S.I. 2004/994, 2005/2114, 2011/1043, 2018/378.

(2) 1986 c.14.

- (i) for “Agency” substitute “appropriate authority”;
 - (ii) for “European Reference Laboratory” substitute “relevant national reference laboratories”;
 - (k) in paragraph 3.5.6—
 - (i) for “Agency’s” substitute “appropriate authority’s”;
 - (ii) omit “other EU and”;
 - (l) in paragraph 3.6.3 for “Agency’s” substitute “the appropriate authority’s”.
- (5) In Annex 2—
- (a) in paragraph 2.1—
 - (i) in the second sentence, for the words from “the Agency’s” to the end substitute “guidance issued by the appropriate authority”;
 - (ii) in the third sentence, omit the words from “as defined” to the end;
 - (b) in paragraph 2.7.2, in the first sentence, for “Agency” substitute “appropriate authority”.