
STATUTORY INSTRUMENTS

2019 No. 88

**The Genetically Modified Organisms (Amendment)
(England) (EU Exit) Regulations 2019**

PART 3

Amendments to subordinate legislation relating
to withdrawal from the European Union

The Genetically Modified Organisms (Deliberate Release) Regulations 2002

3.—(1) The Genetically Modified Organisms (Deliberate Release) Regulations 2002 are amended as follows.

(2) In regulation 2—

(a) for the definition of “approved product” substitute—

““approved product” means a product—

(a) permitted to be marketed in England by—

(i) a consent granted by the Secretary of State under section 111(1) of the Act, or

(ii) an authorisation under the Food and Feed Regulation, or

(b) which, immediately before exit day, was permitted to be marketed by a consent granted in accordance with Article 15(3), 17(6) or 18(2) of the Deliberate Release Directive or Article 13(2) or (4) of the 1990 Directive (a “pre-exit approved product”);”;

(b) omit the definition of “the Commission”;

(c) omit the definition of “the Contained Use Directive”;

(d) for the definition of “the Deliberate Release Directive” substitute—

““the Deliberate Release Directive” means Council Directive [2001/18/EC](#) on the deliberate release into the environment of genetically modified organisms^{MI} as it applied immediately before exit day;”;

(e) in the definition of “the First Simplified Procedure (crop plants) Decision”, insert at the end “ as it applies immediately before exit day ”.

(3) In regulation 9, omit the words from “the release is” to “or in which”.

(4) In regulation 11(1)(d)—

(a) omit the words from “, in the format” to “Directive,”;

(b) at the end, insert “, in the relevant format set out in the Annex to Council Decision [2002/813/EC](#) ”.

(5) In regulation 15—

(a) the existing text becomes paragraph (1);

- (b) in new paragraph (1), after sub-paragraph (a) insert—
 - “(aa) a pre-exit approved product is marketed during the relevant period for a use for which it had approval before exit day and in accordance with the limitations and conditions to which the use of that product was subject before exit day,”;
- (c) for sub-paragraphs (b) and (c) substitute—
 - “(b) genetically modified organisms are made available for activities regulated under the Genetically Modified Organisms (Contained Use) Regulations 2014^{M2},”;
- (d) for sub-paragraph (e) substitute—
 - “(e) a genetically modified organism is marketed which is contained in a medicinal product authorised under the Human Medicines Regulations 2012^{M3} or the Veterinary Medicines Regulations 2013^{M4},”;
- (e) omit sub-paragraph (g);
- (f) after new paragraph (1), insert—
 - “(2) For the purposes of paragraph (1), “the relevant period”, in relation to a pre-exit approved product, means the period beginning with exit day and ending on the day on which the consent concerned ceases to be valid.”.
- (6) In regulation 16(2)—
 - (a) in sub-paragraph (b)—
 - (i) for “European Union” substitute “ United Kingdom ”;
 - (ii) omit the words from “or to another competent authority” to the end;
 - (b) in sub-paragraph (g), after “Directive”, insert “ , as read with the guidance notes set out in Commission Decision 2002/811/EC, ”;
 - (c) in sub-paragraph (j), for the words from “established by the Commission” to the end substitute “ set out in the Annex to Commission Decision 2002/812/EC ”.
- (7) In regulation 20—
 - (a) omit sub-paragraph (c);
 - (b) in sub-paragraph (f), omit the words from “and any comments made” to the end.
- (8) In regulation 21—
 - (a) in paragraph (3), omit “and to the Commission”;
 - (b) for paragraph (6) substitute—
 - “(6) Information submitted in accordance with paragraph (5) must be provided in the format set out in the Annex to Commission Decision 2003/701/EC.”.
- (9) In regulation 23—
 - ^{F1}(a)
 - (b) omit paragraph (2);
 - (c) in paragraph (3), for “paragraphs (1) and (2)” substitute “ paragraph (1) ”;
 - ^{F2}(d)
- (10) In regulation 24—
 - ^{F3}(a)
 - (b) in paragraph (5)—
 - (i) omit “under the relevant EU provisions”;

- (ii) for the words from “an official national catalogue” to the end substitute “ a National List in accordance with regulation 3 of the Seeds (National Lists of Varieties) Regulations 2001 ^{M5} ”.
- (c) in paragraph (6), for the words from “an official national register” to the end substitute “ the National Register in accordance with regulations 6 and 7 of the Forest Reproductive Material (Great Britain) Regulations 2002 ^{M6} ”.
- (11) In regulation 25, omit paragraphs (1)(d) and (2).
- (12) In regulation 26—
 - (a) for paragraph (1) substitute—

“(1) The Secretary of State must not grant an application for the renewal of a consent under section 111(1) of the Act to market genetically modified organisms as it relates to the protection of human health without the agreement of the Health and Safety Executive.”;
 - (b) for paragraph (2) substitute—

“(2) The Secretary of State must communicate a decision on an application to renew a consent to market genetically modified organisms to the applicant as soon as possible and must include in any refusal of a consent the reasons for that decision.”.
- (13) In regulation 28(f), for the words from “the reports of” to “member States” substitute “ monitoring reports in the relevant format set out in the Annexes to Commission Decision [2009/770/EC](#) ”.
- (14) For regulation 31 substitute—

“Variation or revocation of a consent to market

31.—(1) The Secretary of State may only vary or revoke a consent to market genetically modified organisms under section 111(10) of the Act without the agreement of the holder of the consent where new information has become available which the Secretary of State considers would affect the assessment of the risk of damage being caused to the environment by the release.

(2) The Secretary of State must not revoke or vary a consent to market genetically modified organisms under section 111(10) of the Act as it relates to the protection of human health without the agreement of the Health and Safety Executive.”.

- (15) In regulation 32—
 - (a) in paragraph (1), for “an approved” substitute “ marketing a pre-exit approved ”;
 - (b) omit paragraphs (3) to (5).
- (16) In regulation 34—
 - (a) in paragraph (3)—
 - (i) in sub-paragraph (h), after “release of” insert “ , or to market, ”;
 - (ii) after sub-paragraph (h) insert—

“(i) the summary of the information contained in the application required by regulation 11(1)(d) or, as the case may be, of the application required by regulation 16(2)(j).”;
 - (b) after paragraph (3), insert—

“(3A) Subject to paragraph (4) and to the information concerned not being confidential, in relation to an application for a consent under section 111(1) of the Act to market genetically modified organisms—

- (a) the name and address of the person who is responsible for the marketing, whether manufacturer, importer or distributor;
 - (b) the proposed commercial name of the product;
 - (c) the names of the genetically modified organisms in the product, including the scientific and common names of, where appropriate, the parental, recipient and donor organisms;
 - (d) the unique identifiers of the genetically modified organisms in the product;
 - (e) an application reference code assigned by the Secretary of State;
 - (f) the information included in the application as specified at paragraphs 3 and 7 of Schedule 3;
 - (g) information about stored samples of the genetically modified organisms, including the type of material, its genetic characterisation and stability, the amount of repository material, and the conditions of appropriate storage and shelf-life.”;
- (c) in paragraph (7), after “granted” insert “ before exit day ”;
 - (d) in paragraph (9), for “by the” substitute “ before exit day by the European ”.
- (17) In regulation 35, omit paragraphs (8) and (10).
- (18) In Schedule 3—
- (a) in paragraph 2, omit “in the European Union”;
 - (b) in paragraph 5, omit “within the European Union”;
 - ^{F4}(c)
 - (d) in paragraph 8, omit “established in the European Union”;
 - (e) in paragraph 14, for “the European Union” substitute “ England ”.
- (19) In Schedule 4, in paragraph 6, omit the words from “, and whether the views” to the end.

F1	Reg. 3(9)(a) omitted (31.12.2020 immediately before IP completion day) by virtue of The Food and Farming (Amendment) (EU Exit) Regulations 2019 (S.I. 2019/759) , regs. 1(a), 11 ; 2020 c. 1, Sch. 5 para. 1(1)
F2	Reg. 3(9)(d) omitted (31.12.2020 immediately before IP completion day) by virtue of The Food and Farming (Amendment) (EU Exit) Regulations 2019 (S.I. 2019/759) , regs. 1(a), 11 ; 2020 c. 1, Sch. 5 para. 1(1)
F3	Reg. 3(10)(a) omitted (31.12.2020 immediately before IP completion day) by virtue of The Food and Farming (Amendment) (EU Exit) Regulations 2019 (S.I. 2019/759) , regs. 1(a), 11 ; 2020 c. 1, Sch. 5 para. 1(1)
F4	Reg. 3(18)(c) omitted (29.9.2019) by virtue of The Genetically Modified Organisms (Deliberate Release) (Amendment) (England) Regulations 2019 (S.I. 2019/1252) , regs. 1(1), 9

Commencement Information

I1 Reg. 3 in force at 31.12.2020 on IP completion day (in accordance with [2020 c. 1, Sch. 5 para. 1\(1\)](#)), see [reg. 1\(2\)\(b\)](#)

Marginal Citations

- M1** OJ No L 106, 17.4.2001, p. 1, as last amended by Commission Directive (EU) 2018/350 (OJ No L 67, 9.3.2018, p. 30).
- M2** [S.I. 2014/1663](#).
- M3** [S.I. 2012/1916](#), amended by [S.I. 2013/235](#), 1855, 2593, 2014/323, 324, 490, 1878, 2015/178, 259, 354, 903, 1503, 1862, 1879, 2016/186, 190, 696, 2017/715, 1322, 2018/199, 378.

- M4** S.I. 2013/2033, amended by [S.I. 2014/599](#), 2018/761.
- M5** [S.I. 2001/3510](#), amended by [S.I. 2004/2949](#), 2011/464, 2018/942; there are other amending instruments but none is relevant.
- M6** [S.I. 2002/3026](#), to which there are amendments not relevant to these Regulations.

The Genetically Modified Organisms (Transboundary Movements) (England) Regulations 2004

4. In the Schedule to the Genetically Modified Organisms (Transboundary Movements) (England) Regulations 2004 ^{M7}—

- (a) in Part 1, in the text in the second column in the row “Article 10(3)”, for the words from “without authorisation” to the end substitute “ which are not permitted to be marketed in [^{F5}Great Britain], or without authorisation to the import having been expressly agreed by the competent authority of the importing country. ”;
- (b) in Part 2, in the text in the second column in the row “Article 6”, in the second subparagraph, omit “and to the Commission”.

- F5** Words in reg. 4(a) substituted (31.12.2020 immediately before IP completion day) by [The Genetically Modified Organisms \(Amendment\) \(EU Exit\) Regulations 2020 \(S.I. 2020/1421\)](#), regs. 1(4), 3

Commencement Information

- I2** Reg. 4 in force at 31.12.2020 on IP completion day (in accordance with [2020 c. 1](#), [Sch. 5 para. 1\(1\)](#)), see [reg. 1\(2\)\(b\)](#)

Marginal Citations

- M7** [S.I. 2004/2692](#), amended by [S.I. 2008/2598](#), 2011/1043.

Changes to legislation:

There are currently no known outstanding effects for the The Genetically Modified Organisms (Amendment) (England) (EU Exit) Regulations 2019, PART 3.