

2019 No. 483

EXITING THE EUROPEAN UNION

HUMAN TISSUE

**The Quality and Safety of Organs Intended for Transplantation
(Amendment) (EU Exit) Regulations 2019**

Made - - - - *5th March 2019*

Coming into force in accordance with regulation 1

The Secretary of State makes these Regulations in exercise of the powers conferred by sections 8(1) and 23(1) of the European Union (Withdrawal) Act 2018(a).

In accordance with paragraph 1(1) of Schedule 7 to that Act, a draft of this instrument has been laid before Parliament and approved by a resolution of each House of Parliament.

PART 1

Introduction

Citation and commencement

1. These Regulations may be cited as the Quality and Safety of Organs Intended for Transplantation (Amendment) (EU Exit) Regulations 2019 and come into force on exit day.

PART 2

Amendment of primary legislation

Amendment of the Human Tissue Act 2004

2.—(1) Section 32 of the Human Tissue Act 2004(b) (prohibition of commercial dealings in human material for transplantation) is amended as follows.

(2) In subsection (3A)—

(a) for “could result in the United Kingdom being in breach of” substitute “would be incompatible with the principles set out in”;

(b) at the end insert—

(a) 2018 c. 16.

(b) 2004 c. 30. Section 32 was amended by S.I. 2012/1501 and 2014/1459.

“and for the purposes of this subsection, those Articles of those Directives are to be read subject to the modifications set out in subsections (3B) and (3C).”.

(3) After subsection (3A) insert—

“(3B) Article 12 of Directive 2004/23/EC(a) is to be read as if—

(a) in paragraph 1—

(i) for the first subparagraph there were substituted—

“Donations of tissues and cells shall be voluntary and unpaid.”;

(ii) in the second subparagraph, the second sentence were omitted;

(iii) the third subparagraph were omitted;

(b) in paragraph 2, for the first subparagraph there were substituted—

“Any promotion and publicity activities in support of the donation of human tissues and cells shall comply with any directions of the Authority or any provision of any enactment which relates to such activities.”;

(c) also in paragraph 2, in the second subparagraph—

(i) “Member States shall endeavour to ensure that” were omitted;

(ii) for “is” there were substituted “shall be”.

(3C) Article 13 of Directive 2010/53/EU(b) is to be read as if—

(a) in paragraph 1—

(i) “Member States shall ensure that” were omitted; and

(ii) for “are” there were substituted “shall be”.

(b) in paragraph 2, the second sentence were omitted;

(c) in paragraph 3—

(i) “Member States shall prohibit” were omitted; and

(ii) at the end there were inserted “shall be prohibited”;

(d) in paragraph 4—

(i) “Member States shall ensure that” were omitted; and

(ii) for “is” there were substituted “shall be”.

PART 3

Amendment of subordinate legislation

Amendment of the Quality and Safety of Organs Intended for Transplantation Regulations 2012

3.—(1) The Quality and Safety of Organs Intended for Transplantation Regulations 2012(c) are amended as follows.

(2) In regulation 3 (interpretation)—

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

(b) in that paragraph (1)—

(i) omit the definition of “the Directive”;

(ii) omit the definition of “the Implementing Directive”(d);

(a) OJ No L 102, 07.04.2004, p48.

(b) OJ No L 207, 06.08.2010, p14.

(c) S.I. 2012/1501, amended by S.I. 2014/1459 and 2015/1679.

(d) The definition of “the Implementing Directive” was inserted by S.I. 2014/1459.

- (iii) after the definition of “procurement activity” insert—
 - ““procurement organisation” means a healthcare establishment, a team or a unit of a hospital, a person, or any other body which undertakes or coordinates the procurement of organs, and is authorised to do so by the Authority;”;
- (c) after paragraph (1) insert—
 - “(2) In these Regulations, a reference to ensuring compliance with these Regulations includes a reference to ensuring compatibility with the principles set out in Article 13 of Directive 2010/53/EU of the European Parliament and of the Council on standards of quality and safety of human organs intended for transplantation as modified by section 32(3C) of the 2004 Act.”.
- (3) Omit regulation 4 (designation of the competent authority).
- (4) In regulation 5 (licensing requirement), at the end insert—
 - “(6) Schedule 1A (which specifies information to be collected in certain circumstances for the purposes of paragraph 5 of Schedule 1) has effect.”.
- (5) In regulation 6(a) (application of the 2004 Act in relation to licences under Schedule 1), for “the Directive and the Implementing Directive”, in each place where those words appear, substitute “these Regulations”.
- (6) In regulation 12(b) (guidance), in paragraph (1) for “the Directive and the Implementing Directive” substitute “these Regulations”.
- (7) In regulation 13(c) (framework and compliance with licensing conditions and directions), in paragraph (1) omit “in compliance with the Directive and the Implementing Directive”.
- (8) In regulation 18(d) (organs sent to or received from another country)—
 - (a) omit paragraphs (1), (1A) and (2);
 - (b) in paragraph (3) for “to, or received from, countries which are not in the European Union” substitute “, or received from, outside the United Kingdom”;
 - (c) in paragraph (4) for “that are not in the European Union” substitute “outside the United Kingdom”.
- (9) Omit regulation 19 (European Union network of competent authorities).
- (10) In regulation 24 (review) omit subsection (2).
- (11) After regulation 24 insert—

“PART 5A

Power to amend data sets specified in Schedule 1A

Power for appropriate authority to amend Schedule 1A

- 24A.**—(1) The appropriate authority may by regulations amend—
- (a) the minimum data set specified in Part A of Schedule 1A (organ and donor characterisation) where the appropriate authority considers, on the basis of scientific evidence, that the amendment is justified by a serious risk to human health;
 - (b) the complementary data set specified in Part B of that Schedule where the appropriate authority considers, on the basis of scientific evidence, that it is appropriate to do so.

(a) Regulation 6 was amended by S.I. 2014/1459.
 (b) Regulation 12 was amended by S.I. 2014/1459.
 (c) Regulation 13 was amended by S.I. 2014/1459.
 (d) Regulation 18 was amended by S.I. 2014/1459.

- (2) In this regulation—
- “appropriate authority” means—
- (a) in relation to England, the Secretary of State;
 - (b) in relation to Wales—
 - (i) the Welsh Ministers; or
 - (ii) the Secretary of State acting with the consent of the Welsh Ministers;
 - (c) in relation to Scotland—
 - (i) the Scottish Ministers; or
 - (ii) the Secretary of State acting with the consent of the Scottish Ministers;
 - (d) in relation to Northern Ireland—
 - (i) the Department of Health in Northern Ireland; or
 - (ii) the Secretary of State acting with the consent of that Department;
 - (e) for the whole of the United Kingdom, the Secretary of State acting with the consent of the Welsh Ministers, the Scottish Ministers and the Department of Health in Northern Ireland.

Scope and nature of powers

24B.—(1) Regulations made by the Secretary of State or the Welsh Ministers under regulation 24A are to be made by statutory instrument.

(2) For regulations made under regulation 24A by the Scottish Ministers see section 27 of the Interpretation and Legislative Reform (Scotland) Act 2010^(a) (Scottish statutory instruments).

(3) Any power of the Department of Health in Northern Ireland to make regulations under regulation 24A is exercisable by statutory rule for the purposes of the Statutory Rules (Northern Ireland) Order 1979^(b).

- (4) Any power in regulation 24A to make regulations includes power to make—
- (a) different provision for different purposes;
 - (b) consequential, supplementary, incidental, transitional, transitory or saving provision.

Scrutiny of regulations

24C.—(1) A statutory instrument containing regulations made by the Secretary of State under regulation 24A is subject to annulment in pursuance of a resolution of either House of Parliament.

(2) Regulations made under regulation 24A by the Scottish Ministers are subject to the negative procedure (see section 28 of the Interpretation and Legislative Reform (Scotland) Act 2010 (instruments subject to the negative procedure)).

(3) A statutory instrument containing regulations made by the Welsh Ministers under regulation 24A is subject to annulment in pursuance of a resolution of the National Assembly for Wales.

(4) Regulations made by the Department of Health in Northern Ireland under regulation 24A are subject to negative resolution within the meaning of section 41(6) of the Interpretation Act (Northern Ireland) 1954^(c) (definitions for parliamentary purposes) as if they were a statutory instrument within the meaning of that Act.”.

(a) 2010 asp 10.

(b) S.I. 1979/1573 (N.I. 12).

(c) 1954 c. 33.

(12) In Schedule 1 (licences)—

(a) in paragraph 3(a) omit “European Union,”;

(b) in paragraph 5(b)—

(i) in paragraph (i), for “the Annex to the Directive” substitute “Schedule 1A”;

(ii) in paragraph (ii), for the words from “the Annex” to the end substitute “Schedule 1A”;

(c) in paragraph 7 for “the Annex to the Directive” substitute “Schedule 1A”.

(13) After Schedule 1 insert—

“SCHEDULE 1A

Regulation 5

Organ and Donor Characterisation

PART A

Minimum data set

1. The information to be collected pursuant to paragraph 5(b)(i) of Schedule 1 for organ and donor characterisation is the following (the “minimum data set”)—

- (a) the establishment where the procurement takes place and other general data;
- (b) type of donor;
- (c) blood group;
- (d) gender;
- (e) cause of death;
- (f) date of death;
- (g) date of birth or estimated age;
- (h) weight;
- (i) height;
- (j) past or present history of IV drug abuse;
- (k) past or present history of malignant neoplasia;
- (l) present history of other transmissible disease;
- (m) HIV, HCV, HBV tests;
- (n) basic information to evaluate the function of the donated organ.

PART B

Complementary data set

2. The information to be collected pursuant to paragraph 5(b)(ii) of Schedule 1 for organ and donor characterisation is the following (the “complementary data set”)—

General data

- (a) Contact details of the procurement organisation and (if different) the establishment where the procurement takes place necessary for coordination, allocation and traceability of the organs from donors to recipients and vice versa.

Donor data

- (b) Demographic and anthropometrical data required in order to guarantee an appropriate matching between the donor or organ and the recipient.

Donor medical history

- (c) Medical history of the donor, in particular the conditions which might affect the suitability of the organs for transplantation and imply the risk of disease transmission.

Physical and clinical data

- (d) Data from clinical examination which are necessary for the evaluation of the physiological maintenance of the potential donor as well as any finding revealing conditions which remained undetected during the examination of the donor's medical history and which might affect the suitability of the organs for transplantation or might imply the risk of disease transmission.

Laboratory parameters

- (e) Data needed for the assessment of the functional characterisation of the organs and for the detection of potentially transmissible diseases and of possible contraindications with respect to organ donation.

Image tests

- (f) Image explorations necessary for the assessment of the anatomical status of the organs for transplantation.

Therapy

- (g) Treatments administered to the donor and relevant for the assessment of the functional status of the organs and the suitability for organ donation, in particular the use of antibiotics, inotropic support or transfusion therapy.”

(14) In Schedule 2 (directions of the Authority)—

- (a) in paragraph 1, in sub-paragraph (e) omit “European Union,”;
- (b) omit paragraph 3(a).

Signed by authority of the Secretary of State for Health and Social Care.

Jackie Doyle-Price
Parliamentary Under-Secretary of State,
Department of Health and Social Care

5th March 2019

EXPLANATORY NOTE

(This note is not part of the Regulations)

These Regulations are made in exercise of the powers in sections 8(1) and 23(1) of the European Union (Withdrawal) Act 2018 (c. 16) in order to address failures of retained EU law to operate effectively and other deficiencies (in particular under section 8(2)(a), (f) and (g)) arising from the withdrawal of the UK from the European Union.

These Regulations make amendments to legislation in the field of procedures to be followed and information to be transmitted in connection with ensuring the quality and safety of organs intended for transplantation.

Part 2 amends primary legislation. Part 3 amends subordinate legislation, including to confer a power for the appropriate authority to make regulations in connection with the information to be collected concerning the characterisation of organs and donors.

An impact assessment has not been produced for this instrument as no, or no significant, impact on the private or voluntary sector is foreseen.

(a) Paragraph 3 was inserted by S.I. 2014/1459.

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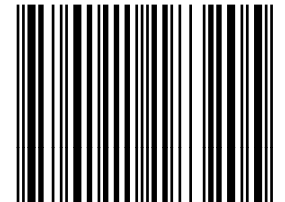
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