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STATUTORY INSTRUMENTS

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**2022 No. 792**

**The Russia (Sanctions) (EU Exit)  
(Amendment) (No. 11) Regulations 2022**

**Interpretation of Part 5**

3.—(1) Regulation 21 is amended as follows.

(2) In paragraph (1), insert in the appropriate places—

““defence and security goods” means—

- (a) interception and monitoring goods,
- (b) internal repression goods, and
- (c) goods relating to chemical and biological weapons;”;

““defence and security technology” means—

- (a) interception and monitoring technology,
- (b) internal repression technology, and
- (c) technology relating to chemical and biological weapons;”;

““goods relating to chemical and biological weapons” means—

- (a) any thing specified in Part 4 of Schedule 3C, other than technology relating to chemical and biological weapons (but see paragraph (4A))(1), and
- (b) any tangible storage medium on which technology relating to chemical and biological weapons is recorded or from which it can be derived;”;

““interception and monitoring goods” means any item mentioned in paragraph (a) or (b), provided that it may be used for interception and monitoring services—

- (a) a relevant Part 2 item(2),
- (b) any tangible storage medium on which interception and monitoring technology is recorded or from which it can be derived;”;

““interception and monitoring technology” means any thing—

- (a) which is described as software in paragraph 2 of Part 2 of Schedule 3C provided that it may be used for interception and monitoring services, and
- (b) which is described as other software or other technology in paragraph 3 of Part 2 of Schedule 3C (but see paragraph (4C));”;

““internal repression goods” means—

- (a) any thing specified in Part 3 of Schedule 3C, other than—
  - (i) any thing which is internal repression technology,

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(1) Inserted by paragraph (7) of this regulation.

(2) “Relevant Part 2 item” is defined in regulation 21(4B) which is inserted by paragraph (7) of this regulation.

- (ii) any thing for the time being specified in Schedule 2 to the Export Control Order 2008(3), or
- (iii) any thing for the time being specified in Annex # of the Dual-Use Regulation, and
- (b) any tangible storage medium on which internal repression technology is recorded or from which it can be derived;”;

““internal repression technology” means any thing which is described in Part 3 of Schedule 3C as software or technology;”;

““maritime goods” and “maritime technology” mean respectively any goods and technology specified in Chapter 4 (Navigation Equipment) and Chapter 5 (Radio-Communication Equipment) of Annex 1 of the Merchant Shipping Notice 1874(4) but not including any thing in those Chapters for the time being specified in—

- (a) Schedules 2 and 3 to the Export Control Order 2008,
- (b) Annex I to the Dual Use Regulation, or
- (c) Schedule 2A;”;

““medical device” means—

- (a) a medical device within the meaning given in regulation 2 of the Medical Devices Regulations 2002(5) in so far as those Regulations apply to England, Wales and Scotland, and
- (b) a medical device within the meaning given in—
  - (i) article 2 of Regulation (EU) 2017/745 of the European Parliament and of the Council of 5 April 2017 on medical devices, amending [Directive 2001/83/EC](#), Regulation (EC) No 178/2002 and Regulation (EC) No 1223/2009 and repealing Council Directives 90/385/EEC and 93/42/EEC, and
  - (ii) article 2 of Regulation (EU) 2017/746 of the European Parliament and of the Council of 5 April 2017 on in vitro diagnostic medical devices and repealing [Directive 98/79/EC](#) and Commission [Decision 2010/227/EU](#),

in so far as those Regulations apply to Northern Ireland;”;

““technology relating to chemical and biological weapons” means any thing specified as technology or software in Part 4 of Schedule 3C, other than technology which is—

- (a) the minimum necessary for—
    - (i) the installation, operation, maintenance and repair of any goods which are not subject to a prohibition under this Part, or
    - (ii) patent applications,
  - (b) in the public domain,
  - (c) a medical device, or
  - (d) used for basic scientific research;”.
- (3) In the definition of “critical-industry goods”, in paragraph (a)(ii)—
- (a) at the end of (aa) omit “or”;
  - (b) at the end of (bb) for “and” substitute “or”;
  - (c) after (bb) insert—

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(3) [S.I. 2008/3231](#).

(4) [https://assets.publishing.service.gov.uk/government/uploads/system/uploads/attachment\\_data/file/1042678/MSN\\_1874\\_Amendment\\_5.pdf](https://assets.publishing.service.gov.uk/government/uploads/system/uploads/attachment_data/file/1042678/MSN_1874_Amendment_5.pdf). Hard copies can be obtained from the Department for International Trade, Old Admiralty Building, London, SW1A 2BL.

(5) [S.I. 2002/618](#).

“(cc) Part 3 of Schedule 3C, and”.

(4) In the definition of “critical-industry technology”—

- (a) at the end of paragraph (a) omit “or”;
- (b) at the end of paragraph (b) insert “or”
- (c) after paragraph (b) insert—

“(c) Part 3 of Schedule 3C.”.

(5) At the end of the definition of “restricted goods” insert—

- “(g) defence and security goods;
- (h) maritime goods;”.

(6) At the end of the definition of “restricted technology” insert—

- “(g) defence and security technology;
- (h) maritime technology;”.

(7) After paragraph (4) insert—

“(4A) The definition of “goods relating to chemical and biological weapons” does not apply to anything specified in Part 4 of Schedule 3C—

- (a) which is—
  - (i) a pharmaceutical formulation designed for human administration in the treatment of a medical condition; and
  - (ii) pre-packaged for distribution as a medicinal product; or
- (b) which is a medical device.

(4B) For the purpose of the definition of “interception and monitoring goods” in paragraph (1) “a relevant Part 2 item” means any thing described in Part 2 of Schedule 3C, other than—

- (a) any thing which is interception and monitoring technology, or
- (b) any thing for the time being specified in—
  - (i) Schedule 2 to the Export Control Order 2008, or
  - (ii) Annex # of the Dual-Use Regulation.

(4C) The definition of “interception and monitoring technology” does not apply to software which is—

- (a) generally available to the public, or
- (b) in the public domain.

(4D) For the purposes of this Part, the following terms have the meaning given to them in the Dual-Use Regulation—

- “basic scientific research”;
- “in the public domain”.