

24 February 2021

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Committee on Import Licensing

AGREEMENT ON IMPORT LICENSING PROCEDURES

NOTIFICATION UNDER ARTICLE 5.1 TO 5.4 OF THE AGREEMENT¹

UNITED KINGDOM

The following notification dated 2 February 2021, is being circulated at the request of the delegation of the United Kingdom.

	Category	Notification details
1	Notifying Member	United Kingdom
2	Title of new legislation/procedure	 Retained EU law²: Regulation (EC) 273/2004 of the European Parliament and Council of 11 February 2004 on drug precursors. Council Regulation (EC) 111/2005 of 22 December 2004 laying down rules for the monitoring of trade between the Community and third countries in drug precursors. Commission Delegated Regulation (EU) 2015/1011 of 24 April 2015 supplementing Regulation (EC) No 273/2004 of the European Parliament and of the Council on drug precursors and Council Regulation (EC) No. 111/2005 laying down rules for the monitoring of trade between the Union and third countries in drug precursors, and repealing Commission Regulation (EC) No. 1277/2005. Commission Implementing Regulation (EU) 2015/1013 of 25 June 2015 laying down rules in respect of Regulation (EC) No. 273/2004 of the European Parliament and of the Council on drug precursors and of Council Regulation (EC) No. 111/2005 laying down rules for the monitoring of trade between the Union and third countries in drug precursors.
		 UK legislation: The Controlled Drugs (Drug Precursors) (Intra-Community Trade) Regulations 2008. The Controlled Drugs (Drug Precursors) (Community External Trade) Regulations 2008. Law Enforcement and Security (Amendment) (EU Exit) Regulations 2019 – Part 5, Chapter 1, Drug Precursors.
3	Date of Publication	1. 26 June 2018 2. 26 June 2018 3. 26 June 2018 4. 26 June 2018 5. 12 February 2008 6. 11 February 2008 7. 28 March 2019

¹ It is understood that the notifying Member has also completed its notification obligations under Article 1.4(a) and Article 8.2(b) regarding the relevant law/regulation/procedure notified for by filling this form in a full and complete manner.

² Some European Union law was retained in UK law by virtue of the European Union (Withdrawal) Act 2018 and the European Union (Withdrawal Agreement) Act 2020, with minor modifications to ensure operability.

	Category	Notification details
4	Date of entry into force	1. 31 December 2020 2. 31 December 2020
		3. 31 December 2020 4. 31 December 2020
		5. 7 March 2008
		6. 7 March 2008 7. 31 December 2020
5	Website link/Official	1. https://www.legislation.gov.uk/eur/2004/273/contents
	publication of the new regulation/procedure	2. https://www.legislation.gov.uk/eur/2005/111/contents 3. https://www.legislation.gov.uk/eur/2015/1011/contents
	regulation/procedure	4. https://www.legislation.gov.uk/eur/2015/1013/2015-06-25
		5. https://www.legislation.gov.uk/uksi/2008/295/contents/made 6. https://www.legislation.gov.uk/uksi/2008/296/contents/made
		7. http://www.legislation.gov.uk/uksi/2019/742/made
6	Have you attached a copy of the regulation (PDF) to the Secretariat	[] Yes. (<i>Please attach a copy of the regulation to the notification.</i>) [X] No.
7	Type of notification	[X] (a) New licensing regulation/procedure³; (please answer question 8 to 14)
		[] (b) Changes to a regulation/procedure which has been previously notified in document:; (please answer question 15 and 16)
8	List of products subject to licensing	Full list in the annex and can be found in the following link:
	nechanig	https://www.gov.uk/government/publications/precursor-chemical- import-and-export-authorisation-wallchart
9	Nature of licensing	Automatic: []
		Non-Automatic: [X]
10	Administrative	(a) Protect public morals;
	purpose/measure being implemented	(b) Protect human, animal or plant life and health; protect environment;
		(c) Collect trade statistics or market surveillance;
		(d) Protection of patents, trademarks and copyrights, and the prevention of deceptive practices;
		(e) Dursue obligations under the UN Charter and other international treaties (i.e. CITES, Basel Convention, Rotterdam Convention, UNSC Resolutions etc.)
		(f) Quota (including TRQ) administration;
		(g) Regulate imports of arms, ammunition or fissionable materials and safeguard national security;
1.	Administrative beduites	(h) Other: (please specify)
11	Administrative body(ies) for submission of	Ministry/authority and Department: Drugs and Firearms Licensing Unit
	applications	Address: 2 Marsham St, LONDON, SW1P 4DF, UK Website: https://www.gov.uk/guidance/precursor-chemical-licensing
		Telephone: 020 7035 6330 E-Mail: dflu.ie@homeoffice.gov.uk
12	Contact point for	Ministry/authority and Department: Drugs and Firearms Licensing
	information on eligibility	Unit Address: 2 Marsham St, LONDON, SW1P 4DF, UK
		Website: https://www.gov.uk/guidance/precursor-chemical-licensing
		Telephone: 020 7035 6330 E-Mail: dflu.ie@homeoffice.gov.uk
13	Expected duration of	Ongoing.
	licensing procedure	

 $^{^3}$ "New licensing regulation/procedure" is understood to refer to any newly introduced law, regulation or procedure, and those which are in force but being notified for the first time to the Committee.

	Category	Notification details
14	A summary of the notification in one of the WTO official languages	The Drug Precursors Licensing regime is an established regime which derives from the requirements of wider international obligations under the United Nations Convention against Illicit Traffic in Narcotic Drugs and Psychotropic Substances 1988.
		This will mean that companies trading in drug precursors will be required to apply to the Home Office Drugs and Firearms Licensing Unit for:
		 Domestic Licences and Registrations – when possessing certain categories of drug precursor chemicals. Import and Export Licences – when trading in certain categories of chemicals.
		 Pre-export notification – when trading in certain categories of chemicals.
		The full licensing requirements and processes can be found at the following link:
4.5	To the second 7/10 miles	https://www.gov.uk/quidance/precursor-chemical-licensing
15	In the case of 7(b), please indicate the type of new	(a) Termination
	change(s)	(b) ☐ Suspension (c) ☐ Modification of specific details in existing
		procedures:
		☐ Product coverage;
		☐ Administrative purpose;
		☐ Automatic or Non-automatic;
		☐ Duration of licensing;
		☐ Change the nature of quantity/value restriction;
		☐ Eligibility of applicants;
		☐ Contact information on eligibility;
		\square Administrative body(ies) for submission of application;
		Documentation requirements (including application form);
		Period for Application;
		Administrative body(ies) to issue licence;
		Processing time for issuing licence;
		Licence fee/administrative charge;
		Deposit/advance payment and relevant conditions;
		Appeal regulations/procedures;
		☐ Validity of licence; ☐ Other conditions of licence (extension, transferability.
		Other conditions of licence (extension, transferability, penalty of non-use etc.);
		Foreign exchange requirements;
		Other: (please specify).
16	Please elaborate the changes in detail (in one of the WTO official	
	languages)	