



28 September 2021

(21-7203)

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Committee on Import Licensing

Original: English

AGREEMENT ON IMPORT LICENSING PROCEDURES

NOTIFICATION UNDER ARTICLE 5.1 TO 5.4 OF THE AGREEMENT¹

UNITED KINGDOM

The following notification dated 27 September 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	1. The Misuse of Drugs Act 1971 2. The Misuse of Drugs Regulations 2001 3. The Misuse of Drugs Regulations (Northern Ireland) 2002
3	Date of Publication	1. 27 May 1971 2. 14 December 2001 3. 2 January 2002
4	Date of entry into force	1. 1 July 1971 2. 1 February 2002 3. 1 February 2002
5	Website link/Official publication of the new regulation/procedure	1. https://www.legislation.gov.uk/ukpga/1971/38/contents 2. https://www.legislation.gov.uk/uksi/2001/3998/contents/made 3. https://www.legislation.gov.uk/nisr/2002/1/contents/made
6	Have you attached a copy of the regulation (PDF) to the Secretariat	<input type="checkbox"/> Yes. (Please attach a copy of the regulation to the notification.) <input checked="" type="checkbox"/> No.
7	Type of notification	<input checked="" type="checkbox"/> (a) New licensing regulation/procedure ² ; (please answer question 8 to 14) <input type="checkbox"/> (b) Changes to a regulation/procedure which has been previously notified in document: _____; (please answer question 15 and 16)

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 "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
8	List of products subject to licensing	<p>Please provide the HS codes and detailed descriptions of the products. In case of a long list, please attach the list as an Annex in MS WORD document.</p> <p>Substances are listed in Parts I, II, III and IV of Schedule 2 to the Misuse of Drugs Act 1971 and in Schedules 1 to 5 to the Misuse of Drugs Regulations 2001, and the Misuse of Drugs (Northern Ireland) Regulations 2002. A list of the most commonly encountered drugs currently controlled under those provisions can be found in the following link:</p> <p>https://www.gov.uk/government/publications/controlled-drugs-list-2/list-of-most-commonly-encountered-drugs-currently-controlled-under-the-misuse-of-drugs-legislation</p>
9	Nature of licensing	<p>Automatic: []</p> <p>Non-Automatic: [X]</p>
10	Administrative purpose/measure being implemented	<p>(a) <input type="checkbox"/> Protect public morals;</p> <p>(b) <input checked="" type="checkbox"/> Protect human, animal or plant life and health; protect environment;</p> <p>(c) <input type="checkbox"/> Collect trade statistics or market surveillance;</p> <p>(d) <input type="checkbox"/> Protection of patents, trademarks and copyrights, and the prevention of deceptive practices;</p> <p>(e) <input checked="" type="checkbox"/> Pursue obligations under the UN Charter and other international treaties (i.e. CITES, Basel Convention, Rotterdam Convention, UNSC Resolutions etc.)</p> <p>(f) <input type="checkbox"/> Quota (including TRQ) administration;</p> <p>(g) <input type="checkbox"/> Regulate imports of arms, ammunition or fissionable materials and safeguard national security;</p> <p>(h) <input type="checkbox"/> Other: _____ (please specify)</p>
11	Administrative body(ies) for submission of applications	<p>Ministry/authority and Department: Drugs and Firearms Licensing Unit</p> <p>Address: PO Box 2164, Croydon, CR90 9TB</p> <p>Website: https://www.gov.uk/guidance/controlled-drugs-domestic-licences https://www.gov.uk/guidance/controlled-drugs-import-and-export-licences</p> <p>Telephone:</p> <p>E-Mail: dflu.ie@homeoffice.gov.uk</p>
12	Contact point for information on eligibility	<p>Ministry/authority and Department: Drugs and Firearms Licensing Unit</p> <p>Address: PO Box 2164, Croydon, CR90 9TB</p> <p>Website: https://www.gov.uk/guidance/controlled-drugs-domestic-licences https://www.gov.uk/guidance/controlled-drugs-import-and-export-licences</p> <p>Telephone:</p> <p>E-Mail: dflu.ie@homeoffice.gov.uk</p>
13	Expected duration of licensing procedure	Ongoing.

	Category	Notification details
14	A summary of the notification in one of the WTO official languages	<p>The Misuse of Drugs Act 1971 is an Act of the Parliament of the United Kingdom. This Act, and the regulations made under it, represent action in line with treaty commitments under the Single Convention on Narcotic Drugs the Convention on Psychotropic Substances, and the United Nations Convention Against Illicit Traffic in Narcotic Drugs and Psychotropic Substances.</p> <p>This will mean that companies trading in Controlled Drugs will be required to apply to the Home Office Drugs and Firearms Licensing Unit for:</p> <ul style="list-style-type: none"> • Import and Export Licences – when trading in certain categories Controlled Drugs (S1-4part 2). <p>The full licensing requirements and processes can be found at the following link:</p> <p>https://www.gov.uk/guidance/controlled-drugs-import-and-export-licences</p>
15	In the case of 7(b), please indicate the type of new change(s)	<p>(a) <input type="checkbox"/> Termination</p> <p>(b) <input type="checkbox"/> Suspension</p> <p>(c) <input type="checkbox"/> Modification of specific details in existing procedures:</p> <p><input type="checkbox"/> Product coverage;</p> <p><input type="checkbox"/> Administrative purpose;</p> <p><input type="checkbox"/> Automatic or Non-automatic;</p> <p><input type="checkbox"/> Duration of licensing;</p> <p><input type="checkbox"/> Change the nature of quantity/value restriction;</p> <p><input type="checkbox"/> Eligibility of applicants;</p> <p><input type="checkbox"/> Contact information on eligibility;</p> <p><input type="checkbox"/> Administrative body(ies) for submission of application;</p> <p><input type="checkbox"/> Documentation requirements (including application form);</p> <p><input type="checkbox"/> Period for Application;</p> <p><input type="checkbox"/> Administrative body(ies) to issue licence;</p> <p><input type="checkbox"/> Processing time for issuing licence;</p> <p><input type="checkbox"/> Licence fee/administrative charge;</p> <p><input type="checkbox"/> Deposit/advance payment and relevant conditions;</p> <p><input type="checkbox"/> Appeal regulations/procedures;</p> <p><input type="checkbox"/> Validity of licence;</p> <p><input type="checkbox"/> Other conditions of licence (extension, transferability, penalty of non-use etc.);</p> <p><input type="checkbox"/> Foreign exchange requirements;</p> <p><input type="checkbox"/> Other: _____ (please specify).</p>
16	Please elaborate the changes in detail (in one of the WTO official languages)	