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(2) The import is for research, medical, pharmaceutical, or protective purposes;

(3) The import is in types and quantities strictly limited to those that can be justified for such purposes; and

(4) You have notified BIS at least 45 calendar days prior to the import, pursuant to § 712.6 of the CWCR.

NOTE 1 TO § 712.2(b): Pursuant to § 712.6, advance notifications of import of saxitoxin of 5 milligrams or less for medical/diagnostic purposes must be submitted to BIS at least 3 days prior to import.

NOTE 2 TO § 712.2(b): For specific provisions relating to the prior advance notification of exports of all Schedule 1 chemicals, see § 742.2 and § 742.18 of the EAR for Schedule 1 chemicals subject to the jurisdiction of the Department of Commerce and see the International Traffic in Arms Regulations (22 CFR parts 120 through 130) for Schedule 1 chemicals subject to the jurisdiction of the Department of State.

(c)(1) The provisions of paragraphs (a) and (b) of this section do not apply to the retention, ownership, possession, transfer, or receipt of a Schedule 1 chemical by a department, agency, or other entity of the United States, or by a person described in paragraph (c)(2) of this section, pending destruction of the Schedule 1 chemical;

(2) A person referred to in paragraph (c)(1) of this section is:

(i) Any person, including a member of the Armed Forces of the United States, who is authorized by law or by an appropriate officer of the United States to retain, own, possess, transfer, or receive the Schedule 1 chemical; or

(ii) In an emergency situation, any otherwise non-culpable person if the person is attempting to seize or destroy the Schedule 1 chemical.


§ 712.4 New Schedule 1 production facility.

(a) Establishment of a new Schedule 1 production facility. (1) If your facility has never before been declared under § 712.5 of the CWCR, or the initial declaration for your facility has been withdrawn pursuant to § 712.5(g) of the CWCR, and you intend to begin production of Schedule 1 chemicals at your facility in quantities greater than 100 grams aggregate per year for research, medical, or pharmaceutical purposes, you must provide an initial declaration (with a current detailed technical description of your facility) to BIS in no less than 200 calendar days in advance of commencing such production. Such facilities are considered to be “new Schedule 1 production facilities” and are subject to an initial inspection within 200 calendar days of submitting an initial declaration.

(2) New Schedule 1 production facilities that submit an initial declaration pursuant to paragraph (a)(1) of this section are considered approved Schedule 1 production facilities for purposes of the CWC, unless otherwise notified by BIS within 30 days of receipt by BIS of that initial declaration.

(b) Types of declaration forms required. If your new Schedule 1 production facility will produce in excess of 100 grams aggregate of Schedule 1 chemicals, you must complete the Certification Form, Form 1–1 and Form A. You must also provide a detailed technical description of the new facility or its relevant parts, and a detailed diagram of the declared areas in the facility.

(c) Two hundred days after a new Schedule 1 production facility submits its initial declaration, it is subject to the declaration requirements in § 712.5(a)(1) and (a)(2) and § 712.5(b)(1)(ii) of the CWCR.