

Form for export authorization established by CND*

Export authorization**

Export authorization No. ...

I. On behalf of the Government of _____, the
(Name of State)

undersigned, empowered by the competent authority, pursuant to article 12, paragraph 1, of the Convention on Psychotropic Substances of 1971, to issue authorizations to export psychotropic substances listed in Schedule I and/or Schedule II of that Convention and/or preparations containing such substances, hereby authorizes with reference to import authorization No. _____,

dated _____
(Day) (Month) (Year)

and issued by _____
(Name of the agency having issued the import authorization)

of _____, which the exporter presented to the
(Name of the importing country)

undersigned, the following export:

1. *Exporter:*

Name: _____

Address: _____

2. *Importer:*

Name: _____

Address: _____

[Note: Export of consignments to a post office box is not allowed.]

3. In the case of the export of a substance or substances listed in Schedule I*** and Schedule II***:

(a) The international non-proprietary name, or, in the absence of such a name, the designation of the substance(s) in the Schedule(s):

** Established by the Commission on Narcotic Drugs in accordance with article 12, paragraph 1, of the Convention on Psychotropic Substances of 1971.

*** Delete whichever is not appropriate.

(b) The quantity of the substance(s) authorized to be exported:

4. In the case of the export of a preparation or preparations containing a substance or substances listed in Schedule I*** and Schedule II***:

(a) The international non-proprietary name(s) of the substance(s) contained therein or, in the absence of such a name, the designation of the substance(s) in the Schedule(s):

(b) The name(s) and contents of active ingredients of the preparation(s) authorized to be exported:

(c) The quantity of the preparation(s) authorized to be exported:

(d) The total quantity of each such substance contained in the total amount of the preparation(s) authorized to be exported:

(e) The pharmaceutical form(s) in which the preparation(s) is (are) authorized to be exported (e.g. ampoule, pill, powder):

II. In the case of an export related to a consignment to be placed in a bonded warehouse (prohibited with regard to substances or preparations in Schedule I), it is hereby certified that the consignment to be exported as specified in section I above shall be placed in the following bonded warehouse:

Name: _____

Address: _____

as approved by the import authorization referred to in section I above.

III. Expiration date

The present export authorization expires on _____
(Day) (Month) (Year)

(Place) (Date of issuance)

(Signature of official, name and stamp
of the competent authority)

Notes:

1. The white copy of this export authorization shall accompany the consignment. The competent authority of the Government having issued this export authorization shall send a copy to the competent authority of the Government of the importing country or region which, when the importation has been effected, shall return the export authorization, with an endorsement certifying the amount actually imported, to the competent authority of the Government of the exporting country or region.
2. The information required shall be given in such a way as to facilitate the task of the control officers to verify the identity of the substances and preparations in the shipment. With regard to the information to be given concerning preparations, the name alone is sufficient only if it can safely be expected that this name will unequivocally indicate to control officers the contents of active ingredients of the preparations in the shipment; otherwise full information on such active ingredients is required.