Home Office Form MD 6A (S)

This copy to accompany the consignment
File No: _____
(as shown in top right corner of licence to export)

MISUSE OF DRUGS ACT 1971
EXPORT DECLARATION

This declaration must be completed for each consignment containing drug specified in Schedule 4 Part II to The Misuse of Drugs Regulations 2001.

1. Exporter
Name ……
Address from which consignment dispatched ……
……
……

2. Importer
Name ……
Address from which consignment be delivered ……
……
……

3. Name of drug(s) in consignment (see Note B overleaf) ……
……
……

4. (a) If consignment includes preparation(s) made up into distinct dosage or other units (e.g. tablets, ampoules, etc.)-

(i) what form(s) are the units in? (e.g., tablets, ampoules, etc.)
……

(ii) how many units of each form are in the consignment?
……

(iii) how much of each base substance is contained in each unit of each form? (if consignment contains any form of unit in more than one size, concentration or composition, specify separately for each such size, concentration or composition) ……
……
……

(b) If consignment consists of pure base drug or a preparation not covered in (a) above -
Specify the total quantity of each drug in the consignment
……
……

5. Date of despatch from address entered at (1) above.
……

I hereby declare that to the best of my knowledge and belief, all the above particulars are correctly and fully stated.

Signed ……) (see Note C

Status ……) overleaf)

Date ……

Notes

A. Please note that completed copies of this form must be sent to:

1. The Home Office, Drugs Licensing, Peel Building, 2 Marsham Street, London, SW1P 4DF.
2. Travel with the consignment to which it refers.
3. Be retained by the exporter.

B. The required particulars must be entered for each drug in the consignment which is specified in Schedule 4 Part II to The Misuse of Drugs Regulations 2001 (that is to say, each base substance, sterioisomeric form or salt, and each preparation or other product, which is specified in that Schedule).

The name entered for each drug must be the international non-proprietary name by which it is known; in the case of a preparation the international non-proprietary name of the drug(s) of which it is a preparation must be included.

"Preparation" includes any mixture, whether solid or liquid, containing one or more of the base substances, stereoisomeric forms or salts specified in Schedule 4 Part II to the Regulations.

C. If the exporter is an individual, he must sign personally; if it is a partnership, a partner must sign; if it is a company, a director or the secretary must sign. The status of the signatory must be indicated.