MISUSE OF DRUGS ACT 1971

IMPORTATION AND EXPORTATION LICENCE

In pursuance of section 3(2) (b) of the Misuse of Drugs Act 1971, the Secretary of State (as regards Great Britain) and the Department of Health and Social Services for Northern Ireland (as regards Northern Ireland) hereby licence, with effect from 1st March 1987 and subject to the terms and conditions specified below, the importation and exportation of:

(*a*) any controlled drug specified in Part I of the Schedule to this licence by a person who is entering or leaving the United Kingdom where that drug is intended for administration for dental or medical purposes to himself or to a member of his household who is unable to administer the drug himself and who is travelling with that person at the time of importation or exportation;

(*b*) any controlled drug specified in Part II of the Schedule to this licence by a doctor of medicine who is entering or leaving the United Kingdom —

(i) with a patient for whose treatment during that journey to or from the United Kingdom the doctor considers the drug may be necessary;

(ii) for the purpose of immediately leaving, or within the next three days entering, the United Kingdom with such a patient; or

(iii) within three days after its lawful exportation by him when leaving, or immediately after its lawful importation by him when entering, the United Kingdom with a patient for whose treatment during that journey from or to the United Kingdom the doctor considered the drug might be necessary.

The terms and conditions attached to this licence are:

1. The amount of any controlled drug imported or exported under this licence shall not exceed —

(*a*) on each occasion on which it is imported or exported, the quantity specified in respect of that dug in column 2 of the Schedule to this licence;

(*b*) in any period of thirty days, twice that quantity.

2. This licence does not apply —

(*a*) to any controlled drug which is not contained in a medicinal product within the meaning of the Medicines Act 1968;

(*b*) unless the controlled drug is under the direct personal supervision of the person importing or exporting it;

(*c*) to the exportation of any controlled drug by a person who is not lawfully in possession of the drug.

It is hereby directed that, notwithstanding any provision in Regulations under section 10 of the Misuse of Drugs Act with respect to record-keeping, no record is required to be kept of any quantity of controlled drug imported or exported under this licence.

This licence shall remain in force until revoked by Order of the Secretary of State and the Department of Health and Social Services for Northern Ireland.

W. H. Bohan

An Assistant Under-Secretary of State

Home Office
7th January 1987

Sealed with the Official Seal of the Department of Health and Social Services for Northern Ireland on 20th January 1987.

J. J. Davis

An Under-Secretary

(L.S.)

Licence to Possess

In pursuance of Regulation 5 of the Misuse of Drugs Regulations 1985, the Secretary of State hereby issues a licence under that Regulation authorising any person who imports a controlled drug under and in accordance with the above importation and exportation licence to have that drug in his possession.

W. H. Bohan

An Assistant Under-Secretary of State

Home Office
7th January 1987

In pursuance of Regulation 5 of the Misuse of Drugs (Northern Ireland) Regulations 1986, the Department of Health and Social Services for Northern Ireland hereby issues a licence under that Regulation authorising any person who imports a controlled drug under and in accordance with the above importation and exportation licence to have that drug in his possession.

Sealed with the Official Seal of the Department of Health and Social Services for Northern Ireland on 20th January 1987.

J. J. Davis

An Under-Secretary

SCHEDULE

Part I

Column 1 *Controlled drugs*	*Column 2* *Maximum quantity*
Alphaprodine	3.6g
Amphetamine	300mg
Amphetamine Phosphate	225mg
Amphetamine Sulphate	225mg
Anhydrous Morphine	360mg
Anileridine	900mg
Anilerdine Hydrochloride	900mg
Anileridine Phosphate	900mg
Benzphetamine	2.25g
Bezitramide	450mg
Chlorphentermine Hydrochloride	975mg
Cocaine Hydrochloride, Nitrate or Sulphate Eydrops	20ml at 4% 4ml maximum strength
Cocaine Hydrochloride, Nitrate or Sulphate as active ingredient in a mixture being used by patient who is terminally ill.	500mg
Dexamphetamine	300mg
Dexamphetamine Phosphate	150mg
Dexamphetamine Sulphate	900mg
Dextromoramide Tartrate	900mg
Diamorphine Hydrochloride as an ingredient in a mixture or linctus	500mg
Diamorphine Hydrochloride Tablets	450mg
Diamorphine Hydrochloride Ampoules	1.35g
Diamorphine Hydrochloride Elixir	500mg
Diethylpropion Hydrochloride	1.125g
Dihydrocodeine Tartrate	3.6g
Dihydrocodeinone 0-carboxymethyloxime	150mg
Dipipanone	600mg
Drotebanol	90mg
Ethchlorvynol	15g
Ethinamate	15g
Fentanyl Citrate	45mg
Glutethimide	7.5g
Hydrocodone Hydrochloride	675mg
Hydrocodone Phosphate	675mg
Hydrocodone Tartrate	675mg
Hydromorphone	360mg
Ketobemidone	450mg
Levorphanol Tartrate	135mg
Mazindol	45mg
Mecloqualone	4.5g
Medicinal Opium	1.8g
Mephentermine Sulphate	2.7g
Meprobamate	3.6g
Methadone Hydrochloride Ampoules	500mg
Methadone Hydrochloride Linctus	500mg
Methadone Hydrochloride Tablets	500mg
Methadyl Acetate	500mg
Methaqualone	4.5g