(2) This Order may be cited as the Export of Goods (North Korea) (Revocation) Order, 1957, and shall come into operation on the 4th day of March, 1957.

Dated this 18th day of February, 1957.

William Hughes,
An Under-Secretary of the
Board of Trade.

EXPLANATORY NOTE

(This Note is not part of the Order, but is intended to indicate its general purport.)

This Order revokes the Export of Goods (North Korea) Order, 1950, which contains an absolute prohibition on the export of all goods to North Korea. In future the export of all goods to North Korea will be subject to licence under the Export of Goods (Control) (Consolidation) Order, 1956.

DANGEROUS DRUGS

1957 No. 704

The Dangerous Drugs Regulations, 1957

Made - - - -	18*th April*, 1957
Laid before Parliament	26*th April*, 1957
Coming into Operation	1*st May*, 1957

In pursuance of the powers conferred on me by section nine of the Dangerous Drugs Act, 1951(**a**), I hereby make the following Regulations:—

1. In sub-paragraph (*f*) of paragraph (1) of Regulation 4 and sub-paragraph (*g*) of paragraph (1) of Regulation 10 of the principal Regulations, for the references to the Food and Drugs Act, 1938(**b**), and the Food and Drugs (Adulteration) Act, 1928(**c**), there shall be substituted respectively references to the Food and Drugs Act, 1955(**d**), and the Food and Drugs (Scotland) Act, 1956(**e**).

2. For sub-paragraph (*b*) of paragraph (1) of Regulation 17 of the principal Regulations (which contains requirements as to the register of drugs required to be kept under the Regulations) there shall be substituted the following sub-paragraphs:—

"(*b*) he shall use a separate register or separate part of the register for entries made with respect to each of the classes of drugs and preparations specified in paragraph 1 of the Third Schedule to these Regulations,

(**a**) 14 & 15 Geo. 6. c. 48. (**b**) 1 & 2 Geo. 6. c. 56. (**c**) 18 & 19 Geo. 5. c. 31.
(**d**) 4 & 5 Eliz. 2. c. 16. (**e**) 4 & 5 Eliz. 2. c. 30.

and for this purpose the drugs comprised in each sub-paragraph of the said paragraph 1, other than sub-paragraph (44), shall constitute a class and each ester, ether or derivative specified in the said sub-paragraph (44) shall, together with its salts and any preparation, admixture, extract or other substance containing it or its salts, constitute a class;

(*c*) he shall use a separate register or separate part of the register for entries made with respect to any drug not specified in Part I of the said Third Schedule to which this Part of these Regulations for the time being applies (that is to say, a drug to which Part III of the Act does not apply at the date of the making of these Regulations but is subsequently applied by virtue of an Order in Council made under the Act) and for this purpose such a drug shall be deemed to comprise its salts and any preparation, admixture, extract or other substance containing it or its salts.".

3. In sub-paragraph (*a*) of paragraph (2) of Regulation 17 of the principal Regulations (which contains exceptions to the requirements as to making entries in the register of drugs required to be kept under the Regulations) after the words "particulars with respect to drugs or preparations" there shall be inserted the words "supplied by him".

4. For the Third Schedule to the principal Regulations there shall be substituted the Schedule set out in the Schedule to these Regulations.

5. In these Regulations the expression "the principal Regulations" means the Dangerous Drugs Regulations, 1953(**a**), as amended by the Dangerous Drugs Regulations, 1954(**b**).

6. The Dangerous Drugs Regulations, 1954, are hereby revoked.

7. These Regulations may be cited as the Dangerous Drugs Regulations, 1957, and shall come into operation on the first day of May, 1957.

R. A. Butler,
One of Her Majesty's Principal Secretaries of State.

Home Office,
Whitehall.
18th April, 1957.

SCHEDULE Regulation 4

"THIRD SCHEDULE Regulations 6 and 17

DRUGS TO WHICH PART II OF THESE REGULATIONS APPLIES AND CLASSES OF DRUGS FOR THE PURPOSES OF THE REGISTER REQUIRED TO BE KEPT UNDER REGULATION 17

1. Subject as provided in paragraph 2 of this Schedule, the drugs to which Part II of these Regulations applies are as follows, that is to say:—

(1) Medicinal opium;

(2) Any extract or tincture of cannabis (Indian hemp) and any preparation, not being a preparation capable of external use only, made from extract or tincture of cannabis;

(a) S.I. 1953/499 (1953 I, p. 540). (b) S.I. 1954/1047 (1954 I, p. 676).

(3) Morphine and its salts, and any solution or dilution of morphine or its salts in an inert substance whether liquid or solid containing any proportion of morphine, and any preparation, admixture, extract or other substance (not being such a solution or dilution as aforesaid) containing not less than one-fifth of one per cent. of morphine (calculated in respect of anhydrous morphine);

(4) Cocaine (including synthetic cocaine) and ecgonine and their respective salts, and any solution or dilution of cocaine or its salts in an inert substance, whether liquid or solid, containing any proportion of cocaine, and any preparation, admixture, extract or other substance (not being such a solution or dilution as aforesaid) containing not less than one-tenth of one per cent. of cocaine or any proportion of ecgonine;

(5) Acetyldihydrocodeine;

(6) Acetyldihydrocodeinone;

(7) Alphameprodine;

(8) Alphaprodine;

(9) Benzylmorphine;

(10) Betameprodine;

(11) Betaprodine;

(12) Diacetylmorphine (also known as diamorphine or heroin);

(13) Diethylthiambutene;

(14) Dihydrodesoxymorphine (also known as desomorphine);

(15) Dihydromorphine;

(16) 1 : 3-Dimethyl-4-phenyl-4-propionyloxyhexamethyleneimine;

(17) Dimethylthiambutene;

(18) Dioxaphetyl butyrate (4-morpholino-2 : 2-diphenyl ethyl butyrate);

(19) Dipipanone;

(20) Ethylmethylthiambutene;

(21) Hydrocodone (also known as dihydrocodeinone or dicodide);

(22) Hydromorphone (also known as dihydromorphinone or dilaudide);

(23) Hydroxypethidine;

(24) Isomethadone (also known as isoamidone);

(25) Ketobemidone;

(26) Levomethorphan;

(27) Levorphanol;

(28) Methadol;

(29) Methadone (also known as amidone);

(30) Methadyl acetate;

(31) Methyldesomorphine (6-methyl-Δ^6-desoxymorphine);

(32) 1-Methyl-4-phenylpiperidine-4-carboxylic acid isopropyl ester;

(33) Metopon (also known as methyldihydromorphinone);

(34) Morphine-N-oxide (also known as genomorphine);

(35) Normethadone;

(36) Oxycodone (also known as dihydrohydroxycodeinone or eucodal);

(37) Pethidine;

(38) Phenadoxone;

(39) Phenomorphan (3-hydroxy-N-phenethylmorphinan);

(40) Propoxyphene (4-dimethylamino-1 : 2-diphenyl-3-methyl-2-propionyloxybutane);

(41) Racemethorphan;

(42) Racemorphan;

(43) Thebaine;

(44) The esters of morphine (other than diacetylmorphine), ecgonine, oxycodone, hydrocodone, hydromorphone, acetyldihydrocodeinone and dihydromorphine; the ethers of morphine (other than benzylmorphine, codeine, ethylmorphine and pholcodine); the morphine-N-oxide derivatives, and any other pentavalent nitrogen morphine derivatives.

2.—(1) In this Schedule—

(*a*) "ecgonine" means laevo-ecgonine and includes any derivatives of ecgonine from which it may be recovered industrially;

(*b*) each drug specified in a sub-paragraph of paragraph 1 of this Schedule which follows sub-paragraph (4) shall be deemed to comprise its salts and any preparation, admixture, extract or other substance containing it or its salts.

(2) This Schedule shall not apply to any drug when that drug is contained in—

(i) the preparations, admixtures, extracts or other substances specified in the Second Schedule to these Regulations, or

(ii) the preparations specified in the Schedule to the Order in Council dated the thirteenth day of April, 1937(a), and by virtue of subsection (2) of section twenty-five of the Act, having effect under subsection (4) of section ten thereof, excluding certain preparations containing morphine, cocaine, etc., from the provisions of Part III of the Act.".

EXPLANATORY NOTE

(This Note is not part of the Regulations, but is intended to indicate their general purport.)

Part II of the Dangerous Drugs Regulations, 1953, makes provision for the control of the manufacture, sale, possession and distribution of the drugs to which it applies; the Third Schedule to the Regulations as amended by the Dangerous Drugs Regulations, 1954, sets out the list of these drugs as at the date of the Regulations of 1954. The principal purpose of these Regulations is to bring the list up to date, a number of new drugs having been brought under control by Orders in Council made under the Dangerous Drugs Act, 1951, since the date of the Regulations of 1954, and several changes in the names of drugs having been made. This is effected by Regulation 4 and the Schedule. The revocation of the Dangerous Drugs Regulations, 1954, is consequential. Regulation 2 is also mainly consequential, but the new sub-paragraph (*c*) of Regulation 17 (1) of the Regulations of 1953 is included to make it clear that a separate register or part of the register must be used for entries made with respect to each new drug brought under control after the date of these Regulations, and revised wording has been used in Regulation 17 (1) (*b*) in order to make it clear that a separate register or part of a register need be kept only with respect to drugs actually obtained or supplied. Certain relaxations in the requirements as to making entries in the register are allowed under the Regulations of 1953 to medical practitioners and authorised sellers of poisons who make certain other specified records regarding drugs supplied by them, and the purpose of Regulation 3 is to make it clear that the relaxations are confined to drugs supplied by them. Regulation 1, bringing statutory references up to date, is consequential on the passing of the Food and Drugs Act, 1955, and the Food and Drugs (Scotland) Act, 1956.

(a) S.R. & O. 1937/327 (Rev. V, p. 622: 1937 p. 623).