E/NL. 1961/44 - 46
30 June 1961
ENGLISH ONLY

UNITED NATIONS

LAWS AND REGULATIONS

PROMULGATED TO GIVE EFFECT TO THE PROVISIONS OF THE CONVENTION OF 13 JULY 1931 FOR LIMITING THE MANUFACTURE AND REGULATING THE DISTRIBUTION OF NARCOTIC DRUGS, AS AMENDED BY THE PROTOCOL OF 11 DECEMBER 1946

UNITED KINGDOM OF GREAT BRITAIN AND NORTHERN IRELAND

Communicated by the Government of the United Kingdom of Great Britain and Northern Ireland

NOTE BY THE SECRETARY-GENERAL -- In accordance with Article 21 of the Convention of 13 July 1931 for Limiting the Manufacture and Regulating the Distribution of Narcotic Drugs, as amended by the Protocol of 11 December 1946, the Secretary-General has the honour to communicate the following legislative texts.

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Statutory Instruments,
1961 No. 837 - Dangerous Drugs

E/NL.1961/44

THE DANGEROUS DRUGS ACT, 1951 (APPLICATION) ORDER, 1961

Made 27th April, 1961
Coming into Operation 15th May, 1961

At the Court at Buckingham Palace, the 27th day of April, 1961

Present,

The Queen's Most Excellent Majesty in Council

Whereas by subsection (2) of section ten of the Dangerous Drugs Act, 1951(a)[1/], power is conferred upon Her Majesty by order in Council to declare that Part III of that Act shall apply to any drug of whatever kind in the same manner as it applies to the drugs mentioned in subsection (1) of the said section ten if it appears to Her Majesty that the drug is, or is likely to be, productive, if improperly used, of ill effects substantially of the same character or nature as, or analogous to, those produced by morphine or cocaine, or is capable of being converted into a substance which is, or is likely to be, productive, if improperly used, of such effects:

And whereas it appears to Her Majesty that the drugs specified in the Schedule to this Order are, or are likely to be, productive, if improperly used, of effects substantially of the same character or nature as, or analogous to, those produced by morphine or cocaine:

Now, therefore, Her Majesty, in pursuance of the power conferred upon Her by subsection (2) of section ten of the Dangerous Drugs Act, 1951, is pleased, by and with the advice of Her Privy Council, to order and declare, and it is hereby ordered and declared, as follows:

(a) 14 & 15 Geo. 6. c. 48.

1/ Note by the Secretariat: E/NL.1952/8.

1. Part III of the Dangerous Drugs Act, 1951 (which provides that it shall not be lawful to import or export any drug to which that Part applies except under licence, and empowers the Secretary of State to make regulations for the purpose of preventing the improper use of such drugs), shall apply to the drugs specified in the Schedule to this Order in the same manner as the said Part III applies to the drugs mentioned in subsection (1) of section ten of that Act.

2. This Order may be cited as the Dangerous Drugs Act, 1951 (Application) Order, 1961, and shall come into operation on the fifteenth day of May, 1961.

W.G. Agnew

SCHEDULE

Allylprodine[2/] (3-allyl-1-methyl-4-phenyl-4-propionoxypiperidine), its salts and any preparation, admixture, extract or other substance containing any proportion of allylprodine;

Clonitazene (2-p-chlorobenzyl-1-(2-diethylaminoethyl)-5-nitrobenzimidazole) [2-(p-chlorbenzyl)-1-diethylaminoethyl-5-nitrobenzimidazole][3/], its salts and any preparation, admixture, extract or other substance containing any proportion of clonitazene;

Diphenoxylate (ethyl 1-(3-cyano-3:3-diphenylpropyl)-4-phenylpiperidine-4-carboxylate) [1-(3-cyano-3,3-diphenylpropyl)-4-phenylpiperidine-4-carboxylic acid ethyl ester], its salts and any preparation, admixture, extract or other substance containing any proportion of diphenoxylate;

Etonitazene (1-(2-diethylaminoethyl)-2-p-ethoxybenzyl-5-nitrobenzimidazole) [2-(p-ethoxybenzyl)-1-diethylaminoethyl-5-nitrobenzimidazole], its salts and any preparation, admixture, extract or other substance containing any proportion of etonitazene;

Hydromorphinol (14-hydroxydihydromorphine), its salts and any preparation, admixture, extract or other substance containing any proportion of hydromorphinol;

Levophenacylmorphan ((-)-3-hydroxy-N-phenacylmorphinan), its salts and any preparation, admixture, extract or other substance containing any proportion of levophenacylmorphan;

Metazocine (2^1-hydroxy-2:5:9-trimethyl-6:7-benzomorphan) [1,2,3,4,5,6,-hexahydro-8-hydroxy-3,6,11-trimethyl-2,6-methano-3-benzazocine], its salts and any preparation, admixture, extract or other substance containing any proportion of metazocine;

N-(2-(N-methylphenethylamino)propyl) propionanilide [N-(2-(methylphenethylamino)-propyl)-propionanilide (diampromide)], its salts and any preparation, admixture, extract or other substance containing any proportion of N-(2-(N-methylphenethylamino)propyl) propionanilide;

Norlevorphanol ((-)-3-hydroxymorphinan), its salts and any preparation, admixture, extract or other substance containing any proportion of norlevorphanol;

Phenampromide (N-(1-methyl-2-piperidinoethyl)propionanilide) [N-(2-(1-methylpiperid-2'yl)-ethyl)-propionanilide], its salts and any preparation, admixture, extract or other substance containing any proportion of phenampromide;

Phenoperidine (ethyl 1(3-hydroxy-3-phenylpropyl)-4-phenylpiperidine-4-carboxylate) [1-(3-hydroxy-3-phenylpropyl)-4-phenylpiperidine-4-carboxylic acid ethyl ester], its salts and any preparation, admixture, extract or other substance containing any proportion of phenoperidine;

Piminodine (ethyl 1-(3-anilinopropyl)-4-phenylpiperidine-4-carboxylate) [1-(3-phenylaminopropyl)-4-phenylpiperidine-4-carboxylic acid ethyl ester], its salts and any preparation, admixture, extract or other substance containing any proportion of piminodine.

2/ Note by the Secretariat: Proposed or recommended international non-proprietary names of drugs are underlined.

3/ Note by the Secretariat: The words in square brackets have been inserted by the Secretariat.

Statutory Instruments,
1961 No. 838 - Dangerous Drugs

E/NL.1961/45

THE DANGEROUS DRUGS ACT, 1951 (EXEMPTED PREPARATIONS) DECLARATION, 1961

Made 27th April, 1961

At the Court at Buckingham Palace, the 27th day of April, 1961

Present,

The Queen's Most Excellent Majesty in Council

Whereas it is enacted by subsection (4) of section ten of the Dangerous Drugs Act, 1951(a)[1/], that if Her Majesty in Council thinks fit to declare that a finding with respect to a preparation containing any of the drugs to which Part III of that Act applies has, in pursuance of Article 8 of the International Opium Convention signed at Geneva on the nineteenth day of February, nineteen hundred and twenty-five, been communicated by the Economic and Social Council of the United Nations to the parties to the said Convention, the provisions of the said Part III shall, as from such date as may be specified in the Declaration, cease to apply to the preparation specified therein:

Now, therefore, Her Majesty is pleased, by and with the advice of Her Privy Council, to declare, and it is hereby declared as follows, that is to say:

1. Findings with respect to the preparations specified in the Schedule hereto have in pursuance of Article 8 of the said Convention been communicated by the Economic and Social Council of the United Nations to the parties to the said Convention and the date from which the provisions of Part III of the Dangerous Drugs Act, 1951, shall cease to apply to the said preparations shall be the fifteenth day of May, 1961.

2. This Order may be cited as the Dangerous Drugs Act, 1951 (Exempted Preparations) Declaration, 1961.

W.G. Agnew

SCHEDULE

1. Tablets each weighing 0.8 grammes and containing 2.5 milligrammes of diphenoxylate hydrochloride and 0.025 milligrammes of atropine sulphate.

2. Preparations containing 2.5 milligrammes of diphenoxylate hydrochloride, 0.025 milligrammes of atropine sulphate, 85 milligrammes of lactose, 7 milligrammes of sugar, 21.6 milligrammes of starch, 3 milligrammes of talc, 1 milligramme of magnesium stearate and 0.7 milligrammes of tartrazine.

Statutory Instruments,
1961 No. 839 - Dangerous Drugs

E/NL.1961/46

THE DANGEROUS DRUGS ACT, 1951 (RELAXATION) ORDER, 1961

Made 27th April, 1961
Coming into Operation 15th May, 1961

At the Court at Buckingham Palace, the 27th day of April, 1961

Present,

The Queen's Most Excellent Majesty in Council

Whereas by subsection (1) of section eleven of the Dangerous Drugs Act, 1951(a)[1], it is provided that it shall not be lawful for any person in the United Kingdom to trade in or manufacture for the purpose of trade any of the products obtained from any of the phenanthrene alkaloids of opium or from the ecgonine alkaloids of the coca leaf, not being a product which was on the thirteenth day of July, nineteen hundred and thirty-one, being used for medical or scientific purposes, and that Her Majesty may, if satisfied that any such product is of medical or scientific value, by Order in Council, direct that the subsection shall cease to apply to that product:

And whereas Her Majesty is satisfied that the drug specified in the Schedule to this Order, being a product obtained from thebaine, one of the phenanthrene alkaloids of opium, which was not on the said date being used for medical or scientific purposes, is of medical value:

Now, therefore, Her Majesty, in pursuance of the power conferred upon Her by subsection (1) of section eleven of the Dangerous Drugs Act, 1951, is pleased, by and with the advice of Her Privy Council, to order, and it is hereby ordered, as follows:

1. Subsection (1) of section eleven of the Dangerous Drugs Act, 1951, shall cease to apply to the drug specified in the Schedule to this Order.

2. This Order may be cited as the Dangerous Drugs Act, 1951 (Relaxation) Order, 1961, and shall come into operation on the fifteenth day of May, 1961.

W.G. Agnew

SCHEDULE

Hydromophinol[2] (14-hydroxydihydromorphine).