United Nations

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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

NEW HEBRIDES

COMMUNICATED BY THE GOVERNMENT OF THE

UNITED KINGDOM OF GREAT BRITAIN AND NORTHERN IRELAND

Lake Success,
New York, 1950

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 hereafter the text of a regulation.

Original: English

NEW HEBRIDES CONDOMINIUM

A JOINT REGULATION

No. 5 of 1950.

To amend the New Hebrides Dangerous Drugs Joint Regulation No. 12 of 1939.

Made by the Resident Commissioners under the provisions of Article 7 of the Anglo-French Protocol of the 6th August, 1914.

Amendment.

1. Article 2 of the New Hebrides Dangerous Drugs Joint Regulation No. 12 of 1939, hereinafter called the principal regulation, is hereby amended by the addition of the following substances and materials:-

"(15) Methyldihydromorphinone (commonly known as Metopon), its salts and any preparation, admixture, extract or other substance containing any proportion of methyldihydromorphinone.

(16) Alphaprodine (α -4-Propionoxy-4-phenyl-1:3-dimethyl-4-piperidine), its salts and any preparation, admixture, extract or other substance containing any proportion of alphaprodine.

(17) Amidone (6-Dimethylamino-4:4-diphenylheptan-3-one), its salts and any preparation, admixture, extract or other substance containing any proportion of amidone.

(18) Betaprodine (β -4-Propionoxy-4-phenyl-1:3-dimethyl-4-piperidine), its salts and any preparation, admixture, extract or other substance containing any proportion of betaprodine.

(19) Hydroxypethidine (Ethyl 4-m-hydroxyphenyl-1-methylpiperidine-4-carboxylate), its salts and any preparation, admixture, extract or other substance containing any proportion of hydroxypethidine.

(20) Isoamidone (6-Dimethylamino-4:4-diphenyl-5-methylhexan-3-one), its salts and any preparation, admixture, extract or other substance containing any proportion of isoamidone.

(21) Ketobemidone (4-Propionyl-4-m-hydroxyphenyl-1-methylpiperidine), its salts and any preparation, admixture, extract or other substance containing any proportion of ketobemidone.

(22) Methadol (6-Dimethylamino-4:4-diphenylheptan-3-ol, its salts and any preparation, admixture, extract or other substance containing any proportion of methadol.

(23) Methadyl acetate (6-Dimethylamino-4:4 diphenyl-3-heptyl acetate), its salts and any preparation, admixture, extract or other substance

containing any proportion of methadylacetate.

(24) Phenadoxone (6-Morpholino-4:4 diphenylheptan-3-one), its salts and any preparation, admixture, extract or other substance containing any proportion of phenadoxone.".

Amendment

2. Article 7 of the principal regulation is hereby amended by the addition of the following sub-article:-

Conditions of possession and use by authorized persons.

"(c) Any of the substances or materials specified in Article 2 hereof, except prepared opium and the pipes and utensils mentioned in Article 2 (14), supplied by a qualified medical practitioner to certain authorized persons working under the medical supervision of the said practitioner and used by such authorized persons for medical purposes. The names of such authorized persons shall first be approved by and registered with either Resident Commissioner who shall communicate such registrations to the other Resident Commissioner. Such authorized persons shall record in duplicate

(i) every dose of any of the substances or materials specified in Article 2 hereof and used under the provisions of this sub-article for medical purposes,

(ii) the name of the patient, and

(iii) the condition of the patient which necessitated the use of the substance or material and shall forward one copy of each such record to the supervising medical practitioner concerned at intervals of not more than three months.".

3. This Regulation which may be cited as the New Hebrides Dangerous Drugs (Amendment) Joint Regulation No. 5 of 1950 shall be read as one with the principal regulation and shall come into force on the day of the date hereof.

DATED AT VILA THIS TWENTY-FIRST DAY OF JULY, 1950.

The Resident Commissioner
for the French Republic.

His Britannic Majesty's
Resident Commissioner.

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