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NOTIFICATION OF PUBLICATION

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ACTS

LOI NO. 9 DE 1988 PORTANT MODIFICATION DU REGLEMENT CONJOINT RELATIF A LA VENTE DE MEDICAMENTS (INTERDICTION).

THE JOINT SALE OF MEDICINES (PROHIBITION) (AMENDMENT) ACT NO. 9 OF 1988.

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ORDERS

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THE SALE OF MEDICINES (REGULATION) ORDER NO. 25 OF 1988.

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THE FIREARMS REGULATIONS NO. 27 OF 1988.

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THE FIREARMS ACT NO. 7 OF 1987 - NOTICE

REPUBLIC OF VANUATU

THE JOINT SALE OF MEDICINES (PROHIBITION) (AMENDMENT) ACT No. 9 OF 1988

Arrangement of Sections

REPUBLIC OF VANUATU

Assent: 12.4.88
Commencement: 27.6.88

THE JOINT SALE OF MEDICINES (PROHIBITION) (AMENDMENT) ACT No. 9 OF 1988

An Act to amend the Joint Sale of Medicines (Prohibition) Regulation No.32 of 1966.

BE IT ENACTED by the President and Parliament as follows:-

AMENDMENT OF SECTION 2 OF THE JOINT REGULATION No. 32 OF 1966

1. Section 2 of the Joint Sale of Medicines (Prohibition) Regulation No. 32 of 1966 (in this Act referred to as the "Regulation") is amended by the repeal of subsections (3) and (4) of that section.

INSERTION OF SECTION 2A IN THE REGULATION

2. The following section is inserted after section 2 of the Regulation:

"REGULATION

2A. (1) The Minister may by order make regulations not inconsistent with the provisions of this Regulation -

(a) regulating the sale of medicines, or

(b) prohibiting the sale of any medicines except upon the prescription of a medical practitioner or dental practitioner or veterinary surgeon.

(2) Any regulation made under subsection (1) may authorize the Principal Pharmacist to make and issue notices or directives with respect to such matters prescribed by that regulation."

AMENDMENT OF SECTION 3 OF THE REGULATION

3. Section 3 of the Regulation is amended by the repeal of subsection (1) of that section and the substitution therefor of the following subsection:

"(1) No person shall sell wholesale any medicine other than those as may be prescribed under section 2A(1) to any person other than a pharmacist or druggist."

COMMENCEMENT

4. This Act shall come into force on the date of its publication in the Gazette.

REPUBLIQUE DE VANUATU

LOI NO. 9 DE 1988 PORTANT MODIFICATION DU REGLEMENT CONJOINT RELATIF A LA VENTE DE MEDICAMENTS (INTERDICTION)

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REPUBLIQUE DE VANUATU

LOI NO. 9 DE 1988 PORTANT MODIFICATION DU REGLEMENT CONJOINT RELATIF A LA VENTE DE MEDICAMENTS (INTERDICTION)

portant modification du Règlement Conjoint No. 32 de 1966 relatif à la vente de médicaments (interdiction)

Le président de la République et le Parlement promulguent le texte suivant :

MODIFICATION DE L'ARTICLE 2 DU REGLEMENT CONJOINT NO. 32 DE 1966

1. L'article 2 du Règlement Conjoint No. 32 de 1966 relatif à la vente de médicaments (interdiction) (mentionné dans la présente loi comme "Règlement") est modifié par la suppression des paragraphes (3) et (4) dudit article.

INSERTION DE L'ARTICLE 2A DANS LE REGLEMENT

2. L'article suivant est inséré après l'article 2 du Règlement :

"REGLE

2A. 1) Le Ministre peut prendre des ordonnances fixant des règles compatibles avec les dispositions du présent Règlement :

a) règlémentant la vente de médicaments, ou

b) interdisant la vente de tout médicament sauf sur ordonnance d'un médecin, d'un dentiste ou d'un chirurgien vétérinaire.

2) Toute ordonnance prise conformément au paragraphe 1) dudit article peut habiliter le pharmacien principal à rendre et à publier des notes ou directives concernant les questions prescrites par ladite règle".

MODIFICATION DE L'ARTICLE 3 DU REGLEMENT

3. L'article 3 du Règlement est modifié par la suppression du paragraphe 1) dudit article et par son remplacement par le paragraphe suivant :

"1) Nul ne peut vendre des médicaments en gros à l'exception de ceux pouvant être prescrits conformément à l'article 2A 1) à toute personne autre qu'un pharmacien ou un dépositaire de médicaments".

ENTREE EN VIGUEUR

4. La présente Loi entrera en vigueur le jour de sa publication au Journal officiel.

REPUBLIC OF VANUATU

THE SALE OF MEDICINES (REGULATION) ORDER No. 25 OF 1988

An Order to regulate the sale of medicines.

IN EXERCISE of the powers conferred by Section 2A of the Joint Sale of Medicines (Prohibition) Regulation No,32 of 1966, as amended, I hereby make the following regulations:-

INTERPRETATION

1. In these regulations unless the context otherwise requires:

"Child - resistant closure" means:

(a) a closure which is resistant to opening by children;

(b) in the case of a can fitted with a pres-on lid, a lid of the design known as "double tight" or "tripple tight",

"dosage unit" means an individual dose of a medicine and includes a tablet, capsule, cachet, single dose powder or single dose sachet of powders or granules

"internal use" means administration:

(a) orally, except for topical effect in the mouth, or

(b) for absorption and the production of a systemic effect,

(i) by way of a body orifice other than the mouth, or

(ii) parenterally, other than by application to unbroken skin.

"medicine" means any substance or preparation which is included in the Schedules to this Order;

"primary pack" means the pack in which a medicine and its immediate container or immediate wrapper or measure pack are presented for sale or supply.

(a) "immediate container" includes all forms of containers in which a medicine is directly packed but does not include any such container intended for consumption or any immediate wrapper;

(b) "immediate wrapper" means metal foil, plastic foil, waxed paper, or any such material not intended for consumption, when used as the first wrapper for a dosage unit which contains any medicine;

...2.

(c) "measure pack" means one of two or more sealed containers each of which contains a measured quantity of medicine for use on one occasion as a pesticide and which form part of a single primary pack;

"therapeutic use" means use in or in connection with,

(a) the preventing, diagnosing, curing or alleviating of a disease, ailment, defect or injury in human beings or animals;

(b) the influencing, inhibiting or modifying of a physiological process in human beings or animals; or

(c) the testing of the susceptibility of human beings or animals to a disease or ailment;

"topical use" means application of a medicine for the purpose of producing a localised effect on the surface of the organ or within the issue to which it is applied.

SALE OF MEDICINES

2. (1) Any person may sell any of the medicines specified in Schedule 1 hereto,

(2) No person, other than a pharmacist or a druggist, shall sell any of the medicines specified in Schedule 2 hereto.

(3) No person shall sell the medicines specified in Schedule 3 hereto, except upon the prescription of a medical practitioner, dental practitioner or a veterinary surgeon.

NOTICES

3. (1) The Principal Pharmacist may issue such notices as he deems fit for the proper carrying out of these regulations.

(2) Any notice issued under this regulation shall be published in the Official Gazette.

COMMENCEMENT

4. This Order shall come into force on the date of its publication in the Gazette.

MADE at Port Vila, this 28th day of June , 1988.

FRED TIMAKATA
Minister of Health

SCHEDULE 1

ACETIC ACID (excluding its salts and derivatives) and preparations containing more than 80 per cent of acetic acid, for therapeutic use.

ALOXIPRIN

AMMONIATED MERCURY

ANTAZOLINE in eye drops.

ASPIRIN except:

(a) when included in Schedule 3,

(b) in individually wrapped powders or sachets of granules each containing 650 milligrams or less of aspirin as the only therapeutically active constituent when:

(i) the pack is labelled with the warning statement:

WARNING - THIS MEDICATION MAY BE DANGEROUS WHEN USED IN LARGE AMOUNTS OR FOR A LONG PERIOD, or

CAUTION - THIS PREPARATION IS FOR THE RELIEF OF MINOR AND TEMPORARY AILMENTS AND SHOULD BE USED STRICTLY AS DIRECTED. PROLONGED USE WITHOUT MEDICAL SUPERVISION COULD BE HARMFUL; and

(ii) in a primary pack containing not more than 12 such powders or sachets of granules; or

(c) tablets or capsules each containing 325 milligrams or less of aspirin as the only therapeutically active constituent when:

(i) the pack is labelled with the warning statement:

WARNING - THIS MEDICATION MAY BE DANGEROUS WHEN USED IN LARGE AMOUNTS OR FOR A LONG PERIOD; or

CAUTION - THIS PREPARATION IS FOR THE RELIEF OF MINOR AND TEMPORARY AILMENTS AND SHOULD BE USED STRICTLY AS DIRECTED. PROLONGED USE WITHOUT MEDICAL SUPERVISION COULD BE HARMFUL; and

(ii) packed in blister or strip packaging or in containers with a child-resistant closure, and

(iii) in a primary pack containing not more than 25 such tablets or capsules.

ATROPINE, except atropine methonitrate included in Schedule 3,

(a) in preparations containing 0.25 per cent or less of atropine; or

(b) atropine sulphate, 0.6 mg tablets in packs of six, when labelled for treatment of organophosphorus poisoning.

.../2.

BELLADONNA in preparations containing 0.25 per cent or less of the alkaloids of belladonna, calculated as hyoscyamine.

BENZAMINE when included in:

(a) lozenges, pastilles, tablets or capsules containing 30mg or less of benzamine in each;

(b) suppositories or bougies containing 200mg or less of benzamine in each, or

(c) preparations for external use, other than eyedrops, containing 10 per cent or less of benzamine.

BENZOCAINE when included in:

(a) lozenges, pastilles, tablets or capsules containing 30mg or less of benzocaine in each;

(b) suppositories or bougies containing 200mg or less of benzocaine in each, or

(c) preparations for external use, other than eye drops, containing 10 per cent or less of benzocaine.

BENZOYL PEROXIDE in preparations for external human therapeutic use containing 5 per cent or less benzoyl peroxide.

BENZYDAMINE in preparations for topical use containing 3 per cent or less of benzydamine.

BROMHEXINE

BROMPHENIRAMINE when compounded with one or more of the following medicaments:

(a) an antitussive except codeine or dihydrocodeine,

(b) an expectorant, or

(c) a sympathomimetic amine,

except in preparations for the treatment of children under 2 years of age.

BUCLIZINE in primary packs of 10 doses or less, for the prevention or treatment of motion sickness.

BUTYLAMINOBENZOATE when included in:

(a) lozenges, pastilles, tablets or capsules containing 30 mg or less of butylaminobenzoate in each;

(b) suppositories or bougies containing 200mg or less of butylaminobenzoate in each, or

.../3.

(c) preparations for external use, other than eye drops, containing 10 per cent or less of butylaminobenzoate.

CARBARYL in preparations for external human therapeutic use containing 2 per cent or less of carbaryl.

CARBENOXOLONE for topical oral use.

CARBETAPENTANE except in preparations containing 0.5 per cent or less of carbetapentane.

CHLOROFORM in preparations for therapeutic use except.

(a) when included in Schedule 3, or

(b) in preparations containing 0.5 per cent or less of chloroform.

CHLORPHENIRAMINE when compounded with one or more of the following medicaments:

(a) an antitussive except codeine or dihydrocodeine,

(b) an expectorant, or

(c) a sympathomimetic amine,

except in preparations for the treatment of children under 2 years of age.

CINNAMEDRINE

CLIOQUINOL and other halogenated derivatives of 8-Hydroxyquinoline for external human use.

CODEINE

(a) when compounded with aspirin, paracetamol or salicylamide or any one of their derivatives, in tablets or capsules each containing 10mg or less of codeine, and no other analgesic substance, when:

(i) packed in blister or strip packaging or in containers with child-resistant closures, and

(ii) in a primary pack containing 25 or less dosage units; or

(b) when compounded with one or more other therapeutically active substances:

(i) in divided preparations containing 10mg or less per dosage unit of codeine and with a recommended dose not exceeding 15mg of codeine, or

(ii) in undivided preparations containing 0.25 per cent or less of codeine and with a recommended dose not exceeding 15mg of codeine.

.../4.

CREOSOTE, for therapeutic use, except in preparations containing 3 per cent or less of phenols included in Schedule 1.

CYANIDES - see hydrocyanic acid.

DDT - see dicophane.

DEXCHLORPHENIRAMINE when compounded with one or more of the following medicaments:

(a) an antitussive except codeine or dihydrocodeine;

(b) an expectorant; or

(c) a sympathomimetic amine;

except in preparations for the treatment of children under 2 years of age.

DEXTROMETHORPHAN when compounded with one or more other therapeutically active substances in such a way that the dextromethorphan contained therein cannot be readily extracted, when:

(a) in divided preparations containing 30mg or less per dosage unit and with a recommended dose not exceeding 30mg of dextromethorphan; or

(b) in undivided preparations containing 0.3 per cent or less of dextromethorphan with a recommended dose not exceeding 30mg of dextromethorphan.

TRANS-4-((3,5-DIBROMO-2-HYDROXYBENZYL) AMINO) CYCLOHEXANOL HYDROCHLORIDE MONOHYDRATE (Sputolysin), in oral preparations for the treatment of animals.

DICOPHANE (DDT) in preparations for human therapeutic use.

DICYCLOMINE in preparations containing 0.1 per cent or less of dicyclomine.

DIMENHYDRINATE in primary packs of 10 doses or less, for the prevention or treatment of motion sickness.

DIMETHISOQUIN in preparations for topical use.

DIPHEMANIL METHYLSULPHATE in preparations for topical use.

DIPHENHYDRAMINE

(a) in primary packs of 10 doses or less, for the prevention or treatment of motion sickness, or

(b) when compounded with one or more of the following medicaments:

(i) an antitussive except codeine or dihydrocodeine;

(ii) an expectorant, or

(iii) a sympathomimetic amine;

.../5.

except in preparations for the treatment of children under 2 years of age.

DIPHENYLPYRALINE when compounded with one or more of the following medicaments:

(a) an antitussive except codeine or dihydrocodeine;

(b) an expectorant, or

(c) a sympathomimetic amine;

except in preparations for the treatment of children under 2 years of age.

DOXYLAMINE when compounded with one or more of the following medicaments:

(a) an antitussive except codeine or dihydrocodeine;

(b) an expectorant; or

(c) a sympathomimetic amine,

except in preparations for the treatment of children under 2 years of age.

EPHEDRINE for internal use, when compounded with one or more other therapeutically active substances in preparations containing 30mg or less of ephedrine per recommended dose, other than preparations for stimulant, appetite supression or weight control purposes, except in liquid preparations containing 10mg or less of ephedrine per recommended dose.

ERYTHRITYL TETRANITRATE for therapeutic use.

ETAFEDRINE

ETHER for therapeutic use except:

(a) when included in Schedule 3; or

(b) in preparations containing 10 per cent or less of ether.

ETHOHEPTAZINE in preparatins containing 1 per cent or less of ethoheptazine.

ETHYLMORPHINE, when compounded with one or more other therapeutically active substances:

(a) in divided preparations containing 10mg or less per dosage unit and with a recommended dose not exceeding 15mg of ethylmorphine; or

(b) in undivided preparations containing 0.25 per cent or less of ethylmorphine with a recommended dose not exceeding 15mg of ethylmorphine.

FLUORIDES for human therapeutic use:

(a) sodium fluoride, in preparations for ingestion containing 2.2mg or less of sodium fluoride per dosage unit, or

(b) in preparations for topical use except:

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(i) in dentifrices containing 1000mg/kg or less of fluoride ion; or

(ii) in substances containing 15mg/kg or less of fluoride ion.

GELSEMIUM

GLUTARALDEHYDE for human therapeutic use.

GLYCERYL TRINITRATE for therapeutic use except when included in Schedule 3.

GUAIPHENESIN

(a) in liquid preparations containing 2 per cent (200 mg/10ml) or less of guaiphenesin; or

(b) in divided preparations containing 120mg or less of guaiphenesin in each dosage unit.

HEXACHLOROPHANE in preparations for human skin cleansing purposes containing 3 per cent or less of hexachlorophane except in preparations for use on infants as specified in Schedule 3.

HOMATROPINE in preparations containing 0.25 per cent or less of homatropine.

HUMAN CHORIONIC GONADOTROPHIN OR ANTIBODY in pregnancy test kits.

HYDROCYANIC ACID and CYANIDES in preparations for therapeutic use containing the equivalent of 0.15 per cent or less of hydrocyanic acid.

8-HYDROXYQUINOLINE and its non-halogenated derivatives for human therapeutic use, except in preparations for external use containing 1 per cent or less of such substances.

HYOSCINE, except hyoscine butylbromide included in Schedule 3:

(a) in preparations containing 0.25 per cent or less of hyoscine; or

(b) in transdermal applicators containing 2mg or less of hyoscine.

HYOSCYAMINE in preparations containing 0.25 per cent or less of hyoscyamine.

HYOSCYAMUS in preparations containing 0.25 per cent or less of the alkaloids of hyoscyamus calculated as hyoscyamine.

IODINE (excluding its salts, derivatives and iodophors), in prparations for human therapeutic use containing more than 2.5 per cent of available iodine.

IRON COMPOUNDS for human internal use except:

(a) when included in Schedule 3;

(b) in divided preparations containing 5 mg or less of iron per dosage unit; or

(c) in liquid oral preparations containing 0.1 per cent or less of iron.

.../7.

ISOPROPAMIDE in preparations containing 2 per cent or less of isopropamide for cutaneous use.

ISOSORBIDE DINITRATE for therapeutic use.

LIGNOCAINE when included in:

(a) lozenges, pastilles, tablets or capsules containing 30mg or less of lignocaine in each;

(b) suppositories or bougies containing 200mg or less of lignocaine in each, or

(c) preparations for external use, other than eye drops, containing 10 per cent or less of lignocaine.

LINDANE in preparations for external human therapeutic use containing 2 per cent or less of lindane.

LOBELIA in preparations containing 0.5 per cent or less of lobeline, except in preparations for smoking or burning.

LUTEINISING HORMONE ANTIBODIES in human ovulation test kits.

MALDISON in preparations for external human therapeutic use containing 2 per cent or less of maldison.

MEBENDAZOLE for human therapeutic use.

MERCURIC CHLORIDE in preparations containing 0.5 per cent or less of mercuric chloride, except when included in a notice.

MERCURIC IODIDE in preparations for therapeutic use containing 2 per cent or less of mercuric iodide.

MERCURIC NITRATE in prepartions for therapeutic use containing 5 per cent or less of mercuric nitrate.

MERCURIC OXIDE and all oxides of mercury.

MERCURIC-POTASSIUM IODIDE in preparations containing the equivalent of 2 per cent or less of mercuric iodide, in such form.

MERCURY (metallic) for therapeutic use.

MERCURY ORGANIC COMPOUNDS for topical therapeutic use in preparations containing 0.5 per cent or less of mercury.

METHOXAMINE except:

(a) preparations containing 0.5 per cent or less of methoxamine; or

(b) preparations for external use containing 1 per cent or less of methoxamine.

.../8.

METHOXYPHENAMINE

METHYLEPHEDRINE

NAPHAZOLINE

NICLOSAMIDE for human therapeutic use.

NITRIC ESTERS of polyhydric alcohols for therapeutic use except when separately specified in these Schedules.

NOSCAPINE

OXETHAZAINE in preparations for interal use only.

OXOLAMINE

OXYMETAZOLINE
PAPAVERINE

PARACETAMOL except:

(a) when included in Schedule 3.

(b) in individually wrapped powders or sachets of granules each containing 1000 milligrams or less of paracetamol as the only therapeutically active constituent when:

(i) the pack is labelled with the warning statement:

WARNING - THIS MEDICATION MAY BE DANGEROUS WHEN USED IN LARGE AMOUNTS OR FOR A LONG PERIOD, or

CAUTION - THIS PREPARATION IS FOR THE RELIEF OF MINOR AND TEMPORARY AILMENTS AND SHOULD BE USED STRICTLY AS DIRECTED. PROLONGED USE WITHOUT MEDICAL SUPERVISION COULD BE HARMFUL, and

(ii) in a primary pack containing not more than 12 such powders or sachets of granules, or

(c) tablets or capsules each containing 500 milligrams or less of paracetamol as the only therapeutically active constituent when:

(i) the pack is labelled with the warning statement:

WARNING - THIS MEDICATION MAY BE DANGEROUS WHEN USED IN LARGE AMOUNTS OR FOR A LONG PERIOD, or

CAUTION - THIS PREPARATION IS FOR THE RELIEF OF MINOR AND TEMPORARY AILMENTS AND SHOULD BE USED STRICTLY AS DIRECTED. PROLONGED USE WITHOUT MEDICAL SUPERVISION COULD BE HARMFUL, and

(ii) packed in blister or strip packaging or in containers with a child-resistant closure; and

.../9

(iii) in a primary pack containing not more than 25 such tablets or capsules.

PHEDRAZINE

PHENAMAZOLINE

PHENAZONE for external use.

PHENIRAMINE

(a) in primary packs of 10 doses or less, for the prevention or treatment of motion sickness, or

(b) when compounded with one or more of the following medicaments:

(i) an antitussive except codeine or dihydrocodeine,

(ii) an expectorant, or

(iii) a sympathomimetic amine;

except in preparations for the treatment of children under 2 years of age.

PHENOL and any homologue of phenol boiling below 220 C, for human therapeutic use, except in prepartions containing 3 per cent or less by weight of such substances.

PHENYLENEDIAMINES and alkylated phenylenediamines for therapeutic use.

PHENYLEPHRINE except:

(a) when included in Schedule 3;

(b) preparations containing 0.5 per cent or less of phenylephrine, or

(c) preparations for external use containing 1 per cent or less of phenylephrine.

PHOLCODINE, when compounded with one or more other therapeutically active substances:

(a) in divided prepartions containing 10mg or less of pholcodine per dosage unit and with a recommended dose not exceeding 25mg of pholcodine, or

(b) in undivided preparations containing 0.5 per cent or less of pholcodine and with a recommended dose not exceeding 25mg of pholcodine.

PODOPHYLLUM RESIN (podophyllin) for external human use in preparatins containing 10 per cent or less of podophyllin.

.../10.

POTASSIUM CHLORATE for therapeutic use except in preparations containing 10 per cent or less of potassium chlorate.

PRAMOXINE when included in preparations for external use, other than eye drops, containing 1 per cent or less of pramoxine.

PROCYCLIDINE in preparations containing 5 per cent or less of procyclidine for cutaneous use.

PROMETHAZINE

(a) in primary packs of 10 doses or less, for the prevention or treatment of motion sickness, or

(b) when compounded with one or more of the following medicaments:

(i) an antitussive except codeine or dihydrocodeine;

(ii) an expectorant, or

(iii) a sympathomimetic amine;

except in preparations for the treatment of children under 2 years of age.

PROPANTHELINE in preparations for topical use.

PROPYLHEXEDRINE in appliances for inhalation in which the substance is absorbed upon an inert solid material.

PSEUDOEPHEDRINE except when included in Schedule 3:

(a) in divided preparations containing 60mg or less of pseudoephedrine per recommended dosage unit, or

(b) in liquid preparations containing 60mg or less of pseudoephedrine per recommended adult dose.

PYRANTEL for human therapeutic use.

PYRITHIONE ZINC for human therapeutic use, except in preparations containing 2 per cent or less of pyrithione zinc, when:

(a) in semisolid hair preparations, or

(b) in shampoos.

SALICYLAMIDE except:

(a) when included in Schedule 3,

(b) in individually wrapped powders or sachets of granules each containing 1 000 milligrams or less of salicylamide as the only therapeutically active constituent when:

(i) the pack is labelled with the warning statement:

.../11.

WARNING - THIS MEDICATION MAY BE DANGEROUS WHEN USED IN LARGE AMOUNTS OR FOR A LONG PERIOD; or

CAUTION - THIS PREPARATION IS FOR THE RELIEF OF MINOR AND TEMPORARY AILMENTS AND SHOULD BE USED STRICTLY AS DIRECTED. PROLONGED USE WITHOUT MEDICAL SUPERVISION COULD BE HARMFUL, and

(ii) in a primary pack containing not more than 12 such powders or sachets of granules; or

(c) tablets or capsules each containing 500 milligrams or less of salicylamide as the only therapeutically active constituent when:

(i) the pack is labelled with the warning statement:

WARNING - THIS MEDICATION MAY BE DANGEROUS WHEN USED IN LARGE AMOUNTS OR FOR A LONG PERIOD, or

CAUTION - THIS PREPARATION IS FOR THE RELIEF OF MINOR AND TEMPORARY AILMENTS AND SHOULD BE USED STRICTLY AS DIRECTED. PROLONGED USE WITHOUT MEDICAL SUPERVISION COULD BE HARMFUL: and

(ii) packed in blister or strip packaging or in containers with a child-resistant closure, and

(iii) in a primary pack containing not more than 25 such tablets or capsules.

SILVER SALTS for therapeutic use, except

(a) chewing tablets containing 5mg or less of silver per tablet; or

(b) solutions containing 0.3 per cent or less of silver.

SODIUM NITRITE for therapeutic use.

SPUTOLYSIN - See trans -4- ((3,5-dibromo-2-hydroxybenzyl)-amino) cyclohexanol hydrochloride monohydrate.

STAPHISAGRIA except in preparations containing 0.2 per cent or less of staphisagria.

STRAMONIUM in preparations containing 0.25 per cent or less of the alkaloids calculated as hyoscyamine, except preparations for smoking or burning.

TETRAHYDROZOLINE

THENYLDIAMINE

(a) in nasal preparations for topical use; or

(b) when compounded with one or more of the following medicaments:

(i) an antitussive except codeine or dihydrocodeine,

(ii) an expectorant, or

(iii) a sympathomimetic amine,

.../12.

except in preparations for the treatment of children under 2 years of age.

TRAMAZOLINE

TRIMEPRAZINE when compounded with one or more of the following medicaments:

(a) an antitussive except codeine or dihydrocodeine;

(b) an expectorant or

(c) a sympathomimetic amine,

except in preparations for the treatment of children under 2 years of age.

TRIPROLIDINE when compounded with one or more of the following medicaments:

(a) an antitussive except codeine or dihydrocodeine,

(b) an expectorant, or

(c) a sympathomimetic amine,

except in preparations for the treatment of children under 2 years of age.

TYMAZOLINE

XYLOMETAZOLINE

SCHEDULE 2

ACEPIFYLLINE in liquid oral preparations.

ADRENALINE in preparations containing 1 per cent or less of adrenaline except in preparations containing 0.02 per cent or less of adrenaline.

AMINOPHYLLINE in liquid oral preparations.

AMYL NITRITE

BENZOYL PEROXIDE in preparations containing 10 per cent or less of benzoyl peroxide for external human therapeutic use, except when included in Schedule 1.

BROMPHENIRAMINE in oral preparations except when included in Schedule 1.

BUCLIZINE in oral preparations except when included in Schedule 1.

BUTYL NITRITE

CHLORAL HYDRATE for human internal therapeutic use in preparations containing 5 per cent or less of chloral hydrate, when packed in containers of 100 ml or less.

CHLOROFLUOROCARBONS - see FLUOROCARBONS

CHLORPHENIRAMINE in oral preparations except when included in Schedule 1.

CLEMASTINE in oral preparations.

CLOTRIMAZOLE, for human use in preparations containing 1 per cent or less of clotrimazole, for treatment of fungal infections of the skin.

CODEINE in tablets or capsules each containing 10mg or less of codeine when compounded with aspirin, paracetamol or salicylamide or any one of their derivatives and no other analgesic substance, except when included in Schedule 1.

CYPROHEPTADINE in oral preparations.

DEXCHLORPHENIRAMINE in oral preparations except when included in Schedule 1.

DIHYDROCODEINE, when compounded with one or more other therapeutically active substances when:

(a) in divided preparations containing 10 mg or less per dosage unit and with a recommended dose not exceeding 15mg of dihydrocodeine; or

(b) in undivided preparations containing 0.25 per cent or less of dihydrocodeine with a recommended dose not exceeding 15mg of dihydrocodeine.

.../2.

DIMENHYDRINATE in oral preparations except when included in Schedule 1.

DIMETHINDENE in oral preparations.

DIPHENHYDRAMINE in oral preparations except when included in Schedule 1.

DIPHENYLPYRALINE in oral preparations except when included in Schedule 1.

DITHRANOL for human therapeutic use.

DOXYLAMINE in oral preparations except when included in Schedule 1.

ECONAZOLE for human use in preparations containing 1 per cent or less of econazole for treatment of fungal infections of the skin.

EPHEDRINE for internal use when compounded with one or more other therapeutically active substances, other than preparations for stimulant, appetite supression or weight control purposes, except:

(a) when included in Schedule 1, or

(b) in liquid preparations containing 10mg or less of ephedrine per recommended dose.

FENOTEROL in metered aerosols delivering 200 micrograms or less of fenoterol per metered dose.

FLAVOXATE

FLUOROCARBONS and CHLOROFLUOROCARBONS alone or in combination with other propelants or refrigerants in liquified gas form for therapeutic use.

FOLIC ACID for human therapeutic use except in preparations containing 500 micrograms or less of folic acid per recommended daily dose.

FOLINIC ACID for human therapeutic use except in preparations containing 500 micrograms or less of folinic acid per recommended daily dose.

FOLINIC ACID for human therapeutic use except in preparations containing 500 micrograms or less of folinic acid per recommended daily dose.

HYDROCORTISONE and HYDROCORTISONE ACETATE as the only therapeutically active substances in preparations for dermal use containing 0.5 per cent or less of hydrocortisone in packs containing 30 grams or less where the labelling warns against contact with the eyes and against use for acne and on children below 2 years of age, use beyond 7 days, and use under occlusive dressings, except on medical advice.

IDOXURIDINE in preparations containing 0.5 per cent or less of idoxuridine for cutaneous use.

INSULIN

ISOCONAZOLE for human use in preparations containing 1 per cent or less of isoconazole, for treatment of fungal infections of the skin.

.../3

LOPERAMIDE in packs of 8 dosage units or less, each dosage unit containing 2mg or less of loperamide.

MEFENAMIC ACID in packs of 30 or less capsules for treatment of spasmodic dysmenorrhoea.

MEPYRAMINE in oral preparations.

METHDILAZINE in oral preparations.

MICONAZOLE for human use in preparations containing 2 per cent or les of miconazole for treatment of fungal infections of the skin.

NAPROXEN in packs of 12 or less tablets for capsules, for treatment of spasmodic dysmenorrhoea.

NITROFURAZONE in preparation for cutaneous use containing 0.2 per cent or less of nitrofurazone.

OCTYL NITRITE

PHENIRAMINE in oral preparatins except when included in Schedule 1.

PHENYLPROPANOLAMINE in preparations for relief of coughs or colds, containing 25 mg or less per dose of phenylpropanolamine.

PHENYLTOLOXAMINE in oral preparations.

PODOPHYLLUM RESIN (Podophyllin) for external human use in preparations containing 20 per cent or less of podophyllin except when included in Schedule 1.

PROMETHAZINE in oral preparations except when included in Schedule 1.

PSEUDOEPHEDRINE except when included in Schedule 1 or 3.

QUININE for human internal therapeutic use except in liquids containing 40mg/L or less of quinine.

SALBUTAMOL

(a) in metered aerosols delivering 100 micrograms or less of salbutamol per metered dose, or

(b) in capsules of dry powder for inhalation delivering 200 micrograms or less of salbutamol per dose.

SANTONIN

SODIUM CROMOGLYCATE in nasal preparations for topical use.

TERBUTALINE in metered aerosols delivering 250 micrograms or less of terbutaline per metered dose.

.../4

THENYLDIAMINE in oral preparations except when included in Schedule 1.

THEOPHYLLINE in liquid oral preparations.

TIOCONAZOLE for human use in prparations containing 1 per cent or less of tioconazole, for treatment of fungal infections of the skin.

TRETINOIN for external human therapeutic use.

TRIMEPRAZINE

(a) in solid oral preparations; or

(b) in liquid oral preparations containing 10mg or less of trimeprazine per 5ml;

except when included in Schedule 1.

TRIPROLIDINE in oral preparations except when included in Schedudle 1.

SCHEDULE 3

(Substances marked + are listed in the Notice)

ACEBUTOLOL

ACEPIFYLLINE except when included in Schedule 2.

ACEPROMAZINE

ACETANILIDE and alkyl acetanilides, for human therapeutic use.

ACETAZOLAMIDE

ACETOHEXAMIDE

ACETYLCHOLINE and other choline esters except when separately specified in this Schedule.

ACETYLCYSTEINE

ACETYLDIHYDROCODEINE, when compounded with one or more other medicaments:

(a) in divided preparations containing not more than 100mg of acetyldihydrocodeine per dosage unit; or

(b) in undivided preparations with a concentration of not more than 2.5 per cent of acetyldihydrocodeine.

ACETYLMETHYLDIMETHYLOXIMIDOPHENYLHYDRAZINE

ACYCLOVIR

ADIPHENINE

ADRENALINE except:

(a) when included in Schedule 2; or

(b) in preparations containing 0.02 per cent or less of adrenaline.

ALCURONIUM

alpha-CHLORALOSE - See chloralose

ALPHADOLONE

ALPHAXALONE

ALPRAZOLAM

ALPRENOLOL

ALPROSTADIL

AMANTADINE

AMBENONIUM

AMBUCETAMIDE

AMBUTONIUM

.../2.

AMETHOCAINE

AMIKACIN

AMILORIDE

AMINOCAPROIC ACID

AMINOGLUTETHIMIDE

AMINOMETRADINE

AMINOPHENAZONE and derivatives therefrom for the treatments of animals.

AMINOPHYLLINE except when included in Schedule 2.

AMINOPTERIN

AMINOREX

AMIODARONE

AMIPHENAZOLE

AMISOMETRADINE

AMITRIPTYLINE and other compounds structurally derived therefrom by substitution in the side chain except when separately specified in this Schedule.

AMODIAQUINE

AMOXYCILLIN

AMPHOMYCIN

AMPHOTERICIN

AMPICILLIN

AMSACRINE

AMYLOBARBITONE when packed and labelled for injection.

AMYLOCAINE

ANABOLIC STEROIDAL AGENTS except when separately specified in this Schedule.

ANGIOTENSINAMIDE

ANTAZOLINE except when included in Schedule 1.

+ANTIBIOTICS except:

.../3.

(a) when separately specified in these Schedules,

(b) avoparcin when packed and labelled for use as an animal feed additive; or

(c) nisin.

ANTIHISTAMINES

(a) when included in Schedule 1 or 2, or

(b) when separately specified in this Schedule.

ANTIMALARIAL SUBSTANCES except when separately specified in this Schedule.

ANTIMONY, organic compounds of, for therapeutic use.

ANTITUBERCULAR SUBSTANCES including isoniazid and its derivatives, para-aminosalicyclic acid and thiacetazone except when separately specified in these Schedules.

APOMORPHINE

APROTININ

ARSENIC - see THIACETARSAMIDE

ASPIRIN when combined with caffeine, paracetamol or salicylamide or any derivative of these substances.

ATENOLOL

ATROPINE METHONITRATE

AURANOFIN

AZAPERONE

AZAPETINE

AZATADINE

AZLOCILLIN

AZTREONAM

BACAMPICILLIN

BACITRACIN except:

(a) when specified in the Notice

(b) in animal feeds for growth promotion containing 50mg/kg or less of antibiotic substances, or

(c) in milk replacers for calves and starter rations for pigs, containing 100mg/kg or less of antibiotic substances.

BACLOFEN

BAMIPINE

BARBITURIC ACID and its derivatives except:

(a) when included or separately specified in the Notice; or

(b) when separately specified in this Schedule.

BECLAMIDE

BEMEGRIDE

BENACTYZINE and other substances structurally derived from diphenylmethane with ataractic properties when used for therapeutic purposes.

BENDROFLUAZIDE

BENORYLATE

BENSERAZIDE

BENZAMINE, except when included in Schedule 1.

BENZHEXOL

BENZILONIUM

BENZOCAINE, except when included in Schedule 1.

BENZODIAZEPINE derivatives except when separately specified in these Schedules.

BENZOYL PEROXIDE in preparations for external human therapeutic use, except when included in Schedule 1 or 2.

BENZPHETAMINE and other substances structurally derived from beta-aminopropylbenzene or beta-aminoisopropylbenzene by substitution in the side-chain or by ring-closure therein (or by both such substitution and such closure) except:

(a) where separately specified in this or any other Schedule; or

(b) ephedrine and pseudoephedrine in preparations exempted from Schedule 1.

BENZTROPINE

BENZYDAMINE except when included in Schedule 1.

BENZYLPENICILLIN (including procaine penicillin) except when specified in the Notice.

BETAHISTINE

BETHANIDINE

BIFONAZOLE

.../5.

BIPERIDEN

BISMUTH compounds of, for human therapeutic or cosmetic use, except:

(a) bismuth citrate when incorporated in hair colourant preparations in concentrations of 0.5 per cent or less;

(b) bismuth oxychloride in cosmetics, or

(c) bismuth formic iodide or bismuth subiodide in dusting powders containing 3 per cent or less of bismuth.

BLEOMYCIN

BORON for human therapeutic use -

(a) for internal use

(b) in glycerines or honeys of borax or boric acid,

(c) in dusting powders for paediatric use; or

(d) as a therapeutically active ingredient in other preparations for dermal use except:

(i) in antifungal preparations, or

(ii) in preparations containing 0.1 per cent or less of boron.

BRETYLIUM

BROMAZEPAM

BROMIDES, inorganic, for therapeutic use.

BROMOCRIPTINE

BROMOFORM for therapeutic use.

BROMPHENIRAMINE except when included in Schedule 1 or 2.

BROMVALETONE

BUCLIZINE except when included in Schedule 1 or 2.

BUFEXAMAC except in preparations containing 5 per cent or less of bufexamac for external human therapeutic use, including suppositories.

BUMETANIDE

BUPIVACAINE

+BUPRENORPHINE

BUSPIRONE

.../6.

BUSULPHAN

BUTACAINE

BUTYLAMINOBENZOATE except when included in Schedule 1.

BUTYLCHLORAL HYDRATE

CALCITONIN

CALCITRIOL

CALCIUM CARBIMIDE for therapeutic use.

CAMPHORATED OIL excluding admixtures.

CAMPHOTAMIDE

CANDICIDIN

CANINE TICK ANTI-SERUM

CANTHARIDIN

CAPREOMYCIN

CAPTODIAME

CAPTOPRIL

CAPURIDE

CARAMIPHEN

CARBACHOL

CARBAMAZEPINE

CARBARYL for human therapeutic use except when included in Schedule 1.

CARBAZOCHROME

CARBENICILLIN

CARBENOXOLONE except when included in Schedule 1.

CARBIDOPA

CARBIMAZOLE

CARBOCROMEN

CARBOPLATIN

CARBROMAL

.../7.

CARDIAC GLYCOSIDES except when separately specified in these Schedules.

CARINDACILLIN

CARMUSTINE

+CARNIDAZOLE

CEFACLOR

CEFOPERAZONE

CEFOTAXIME

CEFOTETAN

CEFOXITIN

CEFTAZIDIME

CEFTRIAXONE

CEPHACETRILE

CEPHADROXIL, for the treatment of animals.

CEPHALEXIN

CEPHALORIDINE

CEPHALOTHIN

CEPHAMANDOLE

CEPHAPIRIN

CEPHAZOLIN

CEPHRADINE

CHENODEOXYCHOLIC ACID

CHLORAL FORMAMIDE

CHLORAL HYDRATE except:

(a) when included in Schedule 2, or

(b) in preparations for topical use containing 2 per cent or less of chloral hydrate.

CHLORALOSE except when specified in the Notice.

+CHLORAMPHENICOL

CHLORAZANIL

.../8.

CHLORBUTOL in preparations for human oral use, except in preparations containing 0.5 per cent or less of chlorbutol as a preservative.

CHLORCYCLIZINE

CHLORDIAZEPOXIDE

CHLORMEKODRIN

CHLORMETHIAZOLE

CHLORMEZANONE

CHLOROFORM for use in anaesthesia.

2-(4-CHLOROPHENYL)-1,2,4-TRIAZOLE [5,1a]-ISOQUINOLINE for the treatment of animals.

CHLOROQUINE

CHLOROTHIAZIDE

CHLORPHENIRAMINE except when included in Schedule 1 or 2.

CHLORPHENTERMINE

CHLORPROMAZINE

CHLORPROPAMIDE

CHLORTETRACYCLINE except when specified in the Notice

CHLORTHALIDONE

CHLORZOXAZONE

CHOLESTYRAMINE for human therapeutic use.

CHYMOPAPAIN, injection for human therapeutic use.

CICLACILLIN

CILASTATIN

CIMETIDINE

CINCHOCAINE

CINOXACIN

CISPLATIN

CLANOBUTIN, in injections for the treatment of animals.

CLAVULANIC ACID

CLEMASTINE except when included in Schedule 2.

CLEMIZOLE

+CLENBUTEROL

CLIDINIUM

CLINDAMYCIN

CLOBAZAM

CLOBETASONE-17-BUTYRATE

CLOFENAMIDE

CLOFIBRATE

+CLOMIPHENE

CLOMIPRAMINE

CLOMOCYCLINE

CLONAZEPAM

CLONIDINE

CLOPAMIDE

+CLOPROSTENOL

CLORAZEPATE

CLOREXOLONE

CLORPRENALINE

CLOTRIMAZOLE, except when included in Schedule 2 or in the Notice.

CLOXACILLIN

CLOZAPINE

CODEINE, except when included in Schedule 1 or 2, when compounded with one or more other therapeutically active substances:

(a) in divided preparations containing 30mg or less of codeine per dosage unit; or

(b) in undivided preparations containing 1 per cent or less of codeine.

.../10.

COLASPASE

COLCHICINE

COLESTIPOL for human therapeutic use.

COLISTIN

CORTISONE and steroid suprarenal cortical hormones, except hydrocortisone in Schedule 2.

CURARE, TUBOCURARINE, d-TUBOCURARINE, d-TUBOCURARINEDIMETHYLETHER and all synthetic quaternary ammonium compounds and other compounds having curarising properties except when separately specified in this Schedule.

CYCLANDELATE

CYCLIZINE

+CYCLOFENIL

CYCLOPENTOLATE

CYCLOPROPANE for therapeutic use.

CYCLOSERINE

CYCLOSPORIN

CYCRIMINE

CYPROHEPTADINE except when included in Schedule 2.

DACARBAZINE

DANAZOL

DANTROLENE

DAPSONE and all derivatives of 4,4-diaminodiphenylsulphone.

DEANOL

DEBRISOQUINE

DEMECARIUM BROMIDE

DEMECLOCYCLINE

DESIPRAMINE

DESMOPRESSIN (D.D.A.V.P.)

+DETOMIDINE

.../11.

DEXCHLORPHENIRAMINE except when included in Schedule 1 or 2.

DEXTROMETHORPHAN except when included in Schedule 1.

+DEXTROPROPOXYPHENE

(a) in divided preparations containing 135mg of dextropropoxyphene or less per dosage unit, or

(b) liquid preparations containing 2.5 per cent or less of dextropropoxyphene.

DEXTRORPHAN

DIAZEPAM

DIBENZEPIN

TRANS-4-((3,5-DIBROMO-2-HYDROXYBENZYL)-AMINO) CYCLOHEXANOL HYDROCHLORIDE MONOHYDRATE (Sputolysin) except when in Schedule 1.

DICHLORALPHENAZONE

DICHLORPHENAMIDE

DICLOFENAC

DICYCLOMINE except when included in Schedule 1.

DIETHAZINE

DIETHYLCARBAMAZINE for human therapeutic use.

DIETHYLPROPION

DIFENOXIN in preparations containing, per dosage unit, 0.5mg or less of difenoxin and a quantity of atropine sulphate equivalent to at least 5 per cent of the dose of difenoxin.

DIFLUNISAL

DIGITALIS and its glycosides.

DIHYDRALAZINE

DIHYDROCODEINE when compounded with one or more other medicaments:

(a) in divided preparations containing not more than 100mg of dihydrocodeine per dosage unit; or

(b) in undivided preparations with a concentration of not more than 2.5 per cent of dihydrocodeine,

except when included in Schedule 1.

.../12.

DIHYDROSTREPTOMYCIN except when specified in the Notice.

DIISOPROPYLAMINE DICHLOROACETATE

DILTIAZEM

DIMENHYDRINATE except when included in Schedule 1 or 2.

DIMETHINDENE except when included in Schedule 2.

DIMETHISOQUIN except when included in Schedule 1.

DIMETHOXANATE

DIMETHYL SULPHOXIDE for therapeutic use except when specified in the Notice.

DINITROCRESOLS for therapeutic use.

DINITRONAPHTHOLS for therapeutic use.

DINITROPHENOLS for therapeutic use.

DINITROTHYMOLS for therapeutic use.

+DINOPROST

DIPERODON

DIPHEMANIL METHYLSULPHATE except when included in Schedule 1.

DIPHENHYDRAMINE except when included in Schedule 1 or 2.

DIPHENIDOL

DIPHENOXYLATE in preparations containing per dosage unit 2.5mg or less of diphenoxylate and a quantity of atropine sulphate equivalent to at least 1 per cent of the dose of diphenoxylate.

DIPHENYLPYRALINE except when included in Schedule 1 or 2.

DIPIVEFRIN

DIPYRIDAMOLE

DISOPHENOL

DISPYRAMIDE

DISULFIRAM for therapeutic use.

DITHIAZANINE except when specified in the Notice.

DOBUTAMINE

DOMPERIDONE

DOPAMINE

DOTHIEPIN

.../13.

DOXAPRAM

DOXEPIN

DOXORUBICIN

DOXYCYCLINE

DOXYLAMINE except when included in Schedule 1 or 2.

DROPERIDOL

DROSTANOLONE

ECONAZOLE except when included in Schedule 2 or in the Notice.

EDETIC ACID for human therapeutic use in preparations for injection or infusion.

EMETINE except in preparations containing 0.2 per cent or less of emetine.

ENALAPRIL

EPHEDRINE except:

(a) when included in Schedule 1 or 2,

(b) in preparations for topical use containing 1 per cent or less of ephedrine, or

(c) when compounded with one or more other therapeutically active substances in liquid preparations for internal use containing 10mg or less of ephedrine per recommended dose, other than preparations for stimulant, appetite suppression or weight control purposes.

ENFLURANE for therapeutic use.

EPICILLIN

EPIRUBICIN

ERGOT

ERYTHROMYCIN except:

(a) when specified in the Notice,

(b) in animal feeds for growth promotion containing 50mg/kg or less of antibiotic substances, or

(c) in milk replacers for calves or starter rations for pigs, containing 100mg/kg or less of antibiotic substances.

ETHACRYNIC ACID

ETHAMBUTOL

.../14.

ETHAMIVAN

ETHCHLORVYNOL

ETHER for use in anaesthesia.

ETHINAMATE

ETHOGLUCID

ETHOHEPTAZINE except when included in Schedule 1.

ETHOPROPAZINE

ETHOXZOLAMIDE

ETHYL CHLORIDE for inhalation anaesthesia.

ETHYLMORPHINE when compounded with one or more other medicaments:

(a) in divided preparations containing not more than 100mg of ethylmorphine per dosage unit, or

(b) in undivided preparations with a concentration of not more than 2.5 per cent of ethylmorphine;

except when included in Schedule 1.

ETHYLOESTRENOL

ETIDOCAINE

ETIDRONATE except in tooth pastes and gels containing 1 per cent or less of etidronate.

ETILEFRIN HYDROCHLORIDE

ETOPOSIDE

+ETRETINATE

FELYPRESSIN

FENCAMFAMIN

FENFLURAMINE

FENOPROFEN

FENOTEROL except when included in Schedule 2.

FENPIPRAMIDE

FENPIPRANE

+FENPROSTALENE

.../15.

FLAVOPHOSPHOLIPOL except:

(a) when specified in the Notice, or

(b) in animal feeds for growth pronotion containing 50mg/kg or less of antibiotic substances.

FLECAINIDE

FLUCLOXACILLIN

FLUCYTOSINE

FLUFENAMIC ACID

FLUNISOLIDE

FLUNITRAZEPAM

FLUNIXIN MEGLUMINE for the treatment of animals.

FLUORIDES in preparations for human ingestion except when included in Schedule 1.

FLUOROURACIL and other substances structurally derived from uracil with cytotoxic properties when used for therapeutic purposes.

FLUOXYMESTERONE

FLUPHENAZINE

+FLUPROSTENOL

FLURAZEPAM

FLUROXENE for inhalation anaesthesia

FLUSPIRILENE

+FOLLICLE STIMULATING HORMONE (See also gonadotrophins)

FRAMYCETIN

FRUSEMIDE

FUSIDIC ACID

GALANTHAMINE

GALLAMINE

GEMEPROST

GENTAMCIN

.../16.

GLIBENCLAMIDE

GLIBORNURIDE

GLICLAZIDE

GLUCAGON

+GLUTETHIMIDE

GLYCERYL TRINITRATE in preparations for injection.

GLYCOPYRROLATE

GLYMIDINE

GONADORELIN

GONADOTROPHINS except when included in Schedule 1.

GRAMICIDIN

GRISEOFULVIN

GROWTH HORMONE

GUAIPHENESIN except when included in Schedule 1.

GUANABENZ

GUANACLINE

GUANETHIDINE

HALCINONIDE

HALOPERIDOL and other substances structurally derived from butyrophenone with ataractic properties when used for therapeutic purposes, except when separately specified in this Schedule.

HALOTHANE for therapeutic use.

HEPARIN for internal therapeutic use.

HETACILLIN

HEXACHLOROPHANE

(a) in preparations for use on infants, or

(b) in other preparations except when included in Schedule 1 or specified in the Notice.

.../17.

HEXAMETHONIUM

HEXOCYCLIUM

HYALURONIC ACID in preparations for injection.

HYDRALAZINE

HYDROFLUMETHIAZIDE

HYDROQUINONE for human therapeutic use except in preparations containing 2 per cent or less of hydroquinone.

HYDROXYCHLOROQUINE

1-HYDROXYPYRIDO (3,2,a)-5-PHENOXAZONE-3-CARBOXYLIC ACID

HYDROXYUREA

HYDROXYZINE

HYGROMYCIN except:

(a) when specified in the Notice, or

(b) in preparations in concentrations of 50mg/kg or less of antibiotic substances.

HYOSCINE BUTYLBROMIDE

HYPOTHALAMIC RELEASING FACTORS except when separately specified in this Schedule.

IBUFENAC

IBUPROFEN

IDOXURIDINE except when included in Schedule 2.

IMIPENEM

IMIPRAMINE

INDAPAMIDE

INDOMETHACIN

INOSITOL NICOTINATE, for internal use.

ION-EXCHANGE RESINS, anionic and cationic, for interal use in humans except when separately specified in this Schedule.

IOPAMIDOL

.../18.

IPRATROPIUM

IRON compounds in injectable preparations for human therapeutic use.

ISOAMINILE

ISOCONAZOLE except when included in Schedule 2 or specified in the Notice.

ISOETHARINE

ISOFLURANE

ISOMETHEPTENE

ISOPRENALINE

ISOPROPAMIDE except when included in Schedule 1.

+ISOTRETINOIN

ISOXUPRINE

KANAMYCIN

KETAMINE

KETOCONAZOLE

KETOPROFEN
KHELLIN

KITASAMYCIN except:

(a) when specified in the Notice; or

(b) in anima feeds for growth promotion containing 100mg/kg or less of antibiotic substances.

LABETALOL

LATAMOXEF

LAUDEXIUM METHYLSULPHATE

LEAD COMPOUNDS for human therapeutic use.

LEFETAMINE

LEPTAZOL

LEUPRORELIN

LEVALLORPHAN

.../19.

LEVAMISOLE

(a) for human therapeutic use, or

(b) in preparations for the prevention or treatment of heartworm in dogs.

LEVODOPA

LIDOFLAZINE

LIGNOCAINE except when included in Schedule 1.

LINCOMYCIN

LINDANE for human therapeutic use except when included in Schedule 1.

LIOTHYRONINE SODIUM (Triiodothyronine).

LITHIUM salts for therapeutic use, except in preparations containing 0.01 per cent or less of litnium.

LOPERAMIDE except when included in Schedule 2.

LORAZEPAM

LOXAPINE

+LUTEINISING HORMONE (See also gonadotrophins).

LYMECYCLINE

MAFENIDE

MALDISON for human therapeutic use except when included in Schedule 1.

MAPROTILINE

MAZINDOL

MEBEVERINE

MEBHYDROLIN

MECAMYLAMINE

MECLOFENOXATE

MECLOZINE

MEDAZEPAM

MEFENAMIC ACID except when incuded in Schedule 2.

MEFLOQUINE

.../20.

MEFRUSIDE

MEPACRINE

MEPENZOLATE

MEPHENESIN and its derivatives except guaiphenesin where specified in Schedule 1 or 3.

MEPHENTERMINE

MEPIVACAINE

MEPROBAMATE

MEPYRAMINE except when included in Schedule 2.

MERCAPTOPURINE and other substances structurally derived therefrom with cytotoxic properties when used for therapeutic purposes.

MERCUROUS CHLORIDE for internal therapeutic use.

MERCURY ORGANIC COMPOUNDS for therapeutic use, except when included in Schedule 1.

METARAMINOL

METFORMIN

METHACYCLINE

METHANDIENONE

METHANDRIOL

METHANTHELINIUM

METHAZOLAMIDE

METHDILAZINE except when included in Schedule 2.

METHENOLONE

METHICILLIN

METHIMAZOLE

METHIXENE

METHOCARBAMOL

METHOTREXATE

METHOXSALEN

METHOXYFLURANE for therapeutic use.

METHYLANDROSTANOLONE

.../21.

METHYCLOTHIAZIDE

METHYLDOPA

METHYLPENTYNOL and other substituted alkynes for internal use.

METHYPRYLONE

METOCLOPRAMIDE

METOLAZONE

METOPROLOL

METRIZAMIDE

METRONIDAZOLE including benzoylmetronidazole

METYRAPONE

MEXILETINE

MEZLOCILLIN

MIANSERIN

MIBOLERONE

MICONAZOLE, except when included in Schedule 2 or in the Notice.

MIDAZOLAM

MINOCYCLINE

MINOXIDIL

MISOPROSTOL

MITHRAMYCIN

MITOBRONITOL

MITOMYCIN

MITOZANTRONE

MONENSIN except:

(a) when specified in the Notice, or

(b) in animal feeds containing 33mg/kg or less of antibiotic substances.

MONOAMINE OXIDASE INHIBITORS, including iproniazid, isocarboxazid, nialamide, phenelzine, pheniprazine and other preparations for which monoamine oxidase inhibition is claimed, except triparanol.

.../22.

MONOBENZONE for human therapeutic use except in preparations containing 2 per cent or less of monobenzone.

MOPERONE

MUPIROCIN

MUSTINE and other substances structurally derived therefrom with cytotoxic properties, when used for therapeutic purposes except when specified in this Schedule.

NADOLOL

+NALBUPHINE

NALIDIXIC ACID except when specified in the Notice.

NALORPHINE

NALOXONE

NANDROLONE

NAPROXEN except when included in Schedule 2.

NARASIN except:

(a) when specified in the Notice, or

(b) in animal feeds containing 100mg/kg or less of narasin.

NATAMYCIN

NEOMYCIN except when specified in the Notice.

NEOSTIGMINE

NETILMICIN

NICOCODINE when compounded with one or more other medicaments:

(a) in divided preparations containing not more than 100mg of nicocodine per dosage unit, or

(b) in undivided preparations with a concentration of not more than 2.5 per cent of nicocodine.

NICODICODINE when compounded lwith one or more other medicaments:

(a) in divided preparations containing not more than 100mg of nicodicodine per dosage unit, or

(b) in undivided preparations with a concentration of not more than 2.5 per cent of nicodicodine.

.../23.

NICOTINE in chewing tablets containing 4mg or less of nicotine per tablet for use as an aid in withdrawal from tabacco smoking.

NICOTINIC ACID for human therapeutic use except in preparations containing 250mg or less of nicotinic acid per recommended daily dose.

NICOTINYL ALCOHOL for internal use.

NICOUMALONE for interal therapeutic use.

NIFEDIPINE

NIFENAZONE

NIKETHAMIDE

NIRIDAZOLE

NITRAZEPAM

NITROFURAN and its derivatives for human therapeutic use except when included in Schedule 2.

NITROUS OXIDE for therapeutic use.

NOMIFENSINE

NORADRENALINE (excluding its derivatives)

NORCODEINE when compounded with one or more other medicaments:

(a) in divided preparations containing not more than 100mg of norcodeine per dosage unit, or

(b) in undivided preparations with a concentration of not more than 2.5 per cent of norcodeine,

except when included in Schedule 1.

NORETHANDROLONE

NORFLOXACIN

NORTRIPTYLINE

NOVOBIOCIN except when specified in the Notice.

NYSTATIN

OCTAMYLAMINE

OCTATROPINE

OLEANDOMYCIN except:

.../24.

(a) when specified in the Notice, or

(b) in animal feeds for growth promotion containing 50mg/kg or less of antibiotic substances.

OPIPRAMOL

ORCIPRENALINE

ORGANOPHOSPHORUS COMPOUNDS with anticholinesterase activity for human therapeutic use except:

(a) when included in Schedule 1; or

(b) when separately specified in this Schedule.

ORNIDAZOLE

ORNIPRESSIN

ORPHENADRINE

ORTHOCAINE

ORTHOPTERIN

OXACILLIN

OXANDROLONE

OXAZEPAM

OXPRENOLOL

OXYBUPROCAINE

OXYMESTERONE

OXYMETHOLONE

OXPENTIFYLLINE

OXYPHENBUTAZONE

OXYPHENCYCLIMINE

OXYPHENONIUM

OXYTETRACYCLINE except when specified in the Notice.

OXYTOCIN

PAMAQUINE

PANCURONIUM

PARACETAMOL when combined with aspirin, caffeine or salicylamide or any derivative of these substances.

.../25.

PARALBEHYDE

PAROMOMYCIN

PEMOLINE

PEMPIDINE

d-PENICILLAMINE

PENTAMETHONIUM

PENTHIENATE

PENTOBARBITONE when packed and labelled for injection.

PENTOLINIUM

PERHEXILENE

PERICYAZINE

PERPHENAZINE

PHENACETIN for therapeutic use.

PHENACEMIDE

PHENAZONE except when included in Schedule 1.

PHENAZOPYRIDINE

PHENETHICILLIN except when specified in the Notice.

PHENFORMIN

PHENGLUTARIMIDE

PHENINDIONE for internal therapeutic use.

PHENIRAMINE except when included in Schedule 1 or 2.

PHENOXYBENZAMINE

PHENOXYMETHYLPENICILLIN except when specified in the Notice.

PHENSUXIMIDE and other substances structurally derived from succinamide with anticonvulsant properties when used for therapeutic purposes.

+PHENTERMINE

PHENTHIMENTONIUM

PHENTOLAMINE

PHENYAPIN

PHENYLBUTAZONE

PHENYLEPHRINE in preparations for human ophthalmic use containing 5 per cent or more of phenylephrine.

PHENYLPROPANOLAMINE except when included in Schedule 2.

.../25.

PHENYLTOLOXAMINE except when included in Schedule 2.

PHENYTOIN and other substances structurally derived from hydantoin with anticonvulsant properties when used for therapeutic purposes.

PHOLCODINE when compounded with one or more other medicaments:

(a) in divided preparations containing not more than 100mg of pholcodine per dosage unit; or

(b) in undivided preparations with a concentration of not more than 2.5 per cent of pholcodine,

except when included in Schedule 1.

PHYSOSTIGMINE

PICROTOXIN

PILOCARPINE except in preparations containing 0.025 per cent or less of pilocarpine.

PIMOZIDE

PINDOLOL

PIPENZOLATE

PIPERACILLIN

PIPERIDOLATE

PIPOBROMAN

PIPRADROL

PIRENZEPINE

PIROXICAM

PITUITARY, its extracts and active principles or their synthetic substitutes except when separately specified in this Schedule.

PIZOTIFEN

PODOPHYLLUM RESIN (Podophyllin) for human therapeutic use except when included in Schedule 1 or 2.

POLYMETHYLENE BISTRIMETHYL AMMONIUM COMPOUNDS

POLYMYXIN

POLYSULPHATED GLYCOSAMINOGLYCANS in preparations for injection, except where otherwise specified in this Schedule.

POTASSIUM PERCHLORATE for therapeutic use.

PRACTOLOL

.../27.

PRAMOXINE except when included in Schedule 1.

PRAZEPAM

PRAZOSIN

PREGNENOLONE ACETATE except in preparations for topical use.

PRENYLAMINE

PRILOCAINE

PRIMAQUINE

PRIMIDONE

PROBENECID

PROBUCOL

PROCAINAMIDE

PROCAINE

PROCARBAZINE

PROCHLORPERAZINE

PROCYCLIDINE except when included in Schedule 1.

PROGUANIL

PROLINTANE

PROMAZINE

PROMETHAZINE except when included in Schedule 1 or 2.

PROPANIDID

PROPANTHELINE except when included in Schedule 1.

PROPRANOLOL

PROPOFOL

PROPOXUR for human therapeutic use.

PROPYLHEXEDRINE except when included in Schedule 1.

PROPYHENAZONE

PROQUAZONE

PROSTAGLANDINS except where separately specified in this Schedule.

.../28.

+PROSTIANOL

PROTHIONAMIDE

PROTIRELIN (thyrotrophin releasing factor).

PROTRIPTYLINE

PROXYMETACAINE

PSEUDOEPHEDRINE in preparations for stimulant, appetite suppression or weight control purposes.

PYRIDOSTIGMINE

PYRIDOXINE HYDROCHLORIDE in preparations for human use containing more than 50mg of pyridoxine per recommended daily dose unless labelled with the warning statement "WARNING - THIS MEDICATION MAY BE DANGEROUS WHEN USED IN LARGE AMOUNTS OR FOR A LONG PERIOD".

PYRIMETHAMINE

QUINETHAZONE

QUINIDINE

RANITIDINE

RAUWOLFIA SERPENTINA

RIFAMPICIN

RITODRINE

ROLITETRACYCLINE

ROSOXACIN

SALBUTAMOL except when included in Schedule 2.

SALICYLAMIDE when combined with aspirin, caffeine or paracetamol or any derivative of these substances.

SALINOMYCIN except:

(a) when specified in the Notice, or

(b) in animal feeds containing 60mg/kg or less of antibiotic substances.

SELENIUM except:

(a) when specified in the Notice,

(b) as selenium arsenide in photocopier drums,

(c) in animal feeds containing 0.1g/tonne or less of selenium,

(d) in compressed pellets for control of selenium responsive conditions in sheep or cattle, or

(e) in fertilizers containing 200g/tonne or less of selenium.

SEX HORMONES and all substances having sex hormonal activity except when separately specified in these Schedules.

+SILVER SULPHADIAZINE

SISOMYCIN

SODIUM CELLULOSE PHOSPHATE for human internal use.

SODIUM CROMOGLYCATE except when included in Schedule 2.

SODIUM NITROPRUSSIDE for human therapeutic use.

SODIUM VALPROATE

SONTOQUINE

SOTALOL

SPARTEINE

SPECTINOMYCIN

SPIRAMYCIN except:

(a) when specified in the Notice; or

(b) in animal feeds for growth promotion in pigs or poultry containing 50mg/kg or less of antibiotic substances.

SPIRONOLACTONE

SPUTOLYSIN - See trans-4-(3,5-bibroma-2-hydroxybenzyl)-amino cyclohexanol hydrochloride monohydrate.

STANOLONE

STANOZOLOL

STREPTOMYCIN except when specified in the Notice.

STROPHANTHUS and its glycosides.

STRYCHNINE in preparations containing 1.5 per cent or less of strychnine for the treatment of animals.

SULFAMETROLE

SULINDAC

SULPHANILAMIDE and its derivatives except:

(a) when specified in the Notice;

(b) when separately specified in this Schedule;

../30.

(c) oryzalin;

(d) sulphaquinoxaline in animal feeds containing 200mg/kg or less of sulphaquinoxaline, or

(e) sulphaquinoxaline when incorporated in baits for the destruction of vermin.

+SULPHATROXAZOLE

SULPHINPYRAZONE

SULPHOMYXIN

SULPHONAL and alkyl sulphonals.

SULTHIAME

SUXAMETHONIUM

TACRINE

TAMOXIFEN

TEMAZEPAM

TENIPOSIDE

TERBUTALINE except when included in Schedule 2.

TERFENADINE

TEROPTERIN

TETRABENAZINE

TETRACOSACTRIN

TETRACYCLINE except when specified in the Notice.

+THALIDOMIDE

THENYLDIAMINE except when included in Schedule 1 or 2.

THEOPHYLLINE except when included in Schedule 2.

THIACETARSAMIDE, in preparations for the prevention or treatment of heart worm in dogs.

THIACETAZONE

THIAMBUTOSINE

THIAZOSULPHONE

THIETHYLPERAZINE

THIOPROPAZATE

THIORIDAZINE

THIOTEPA and other substances structurally derived therefrom with cytotoxic properties when used for therapeutic purposes.

THIOTHIXENE

THIOURACIL and substances structurally derived therefrom with antithyroid properties when used for therapeutic purposes.

THIOUREA for therapeutic use.

THYROID and extracts, and its active principles except when separately specified in this Schedule.

THYROTROPHIN (T.S.H.)

THYROXINE SODIUM

TIAMULIN except:

(a) when specified in the Notice.

(b) in prepared animal feeds.

TICARCILLIN

TIEMONIUM

TIGLOIDINE

TIMOLOL

TINIDAZOLE

TIOCONAZOLE except when in Schedule 2.

TIPEPIDINE

TOBRAMYCIN

TOCAINIDE

TOLAZAMIDE

TOLAZOLINE for internal use.

TOLBUTAMIDE

TOLPROPAMINE

TRANEXAMIC ACID

TRETAMINE

TRIAMTERENE

../32.

TRIAZIQUONE

TRIAZOLAM

TRICHLOROETHYLENE for therapeutic use.

TRICLOFOS

TRICYCLAMOL

TRIDIHEXETHYL

TRIFLUOPERAZINE

TRIFLUPERIDOL

TRIMEPRAZINE except when included in Schedule 1 or 2.

TRIMETAPHAN

TRIMETHOPRIM

TRIMIPRAMINE

TRIMUSTINE

TRIOXYSALEN

TRIPELENNAMINE

TRIPROLIDINE except when included in Schedule 1 or 2.

TROXIDONE and other substances structurally derived from oxazolidinone with anticonvulsant properties when used for therapeutic purposes.

TYLOSIN except:

(a) when included in the Notice,

(b) in animal feeds for growth promotion containing 50mg/kg or less of antibiotic substances, or

(c) in milk replacers for calves or starter rations for pigs, containing 100mg/kg or less of antibiotic substances.

URETHANE (excluding its derivatives) for therapeutic use.

.../33.

URETHANES AND UREIDES having or purporting to have soporific, hypnotic or narcotic properties except when separately specified in these Schedule.

VACCINES, sera, toxoids, and antigens for human parenteral use.

VACCINES, veterinary live virus except:

(a) poultry vaccines,

(b) pigeon pox vaccine, or

(c) scabby mouth vaccine.

VALNOCTAMIDE

VASOPRESSIN

VERAPAMIL

VERATRUM for therapeutic use.

VECURONIUM

VIDARABINE

VINCA ALKALOIDS including semi-synthetic derivatives.

VIPRYNIUM

VIRGINIAMYCIN except:

(a) when specified in the Notice, or

(b) in animal feeds for growth promotion containing 50mg/kg or less of antibiotic substances.

VISNADINE

VITAMIN A for human therapeutic use, except in preparation containing 10 000 I.U. or less of vitamin A per recommended daily dosage.

VITAMIN D for human therapeutic use except in preparations containing 25 micrograms or less of vitamin D per recommended daily dosage.

WARFARIN for internal therapeutic use.

XANTHINE OXIDASE INHIBITORS including allopurinol.

XANTHINOL NICOTINATE

XYLAZINE

YOHIMBINE

ZERANOL except when specified in the Notice.

REPUBLIC OF VANUATU

THE FIREARMS REGULATIONS No. 27 OF 1988

Arrangement of Sections

.../2.

REPUBLIC OF VANUATU

THE FIREARMS REGULATIONS No. 27 OF 1988

To provide for the forms, fees and other matters connected therewith for the purposes of the Firearms Act No.7 of 1987.

IN EXERCISE of the powers conferred by section 42 of the Firearms Act No.7 of 1987, I HEREBY make the following regulations:-

INTERPRETATION

1. In these regulations, unless the context otherwise requires:

"Act" means the Firearms Act No.7 of 1987;

"Licensing Officer" means a licensing officer appointed under section 2 of the Act.

FORM OF APPLICATION FOR FIREARM LICENCE

2. (1) An application for the grant of a firearm licence under subsection (1) of section 9 of the Act shall be made to the licensing officer of the area in which the applicant resides in Form 1 in Schedule 1 hereto.

(2) An application for the renewal of a firearm licence under the Act shall be made to the licensing officer in Form 1 in Schedule 1 hereto.

FORM AND CONDITIONS OF FIREARM LICENCE

3. (1) Every firearm licence granted under subsection (4) of section 9 of the Act -

(a) shall be in Form 2 set out in Schedule 1 hereto;

(b) shall contain the certificate in Form 3 set out in Schedule 1 hereto; and

shall be subject to the following conditions:-

.../2.

(a) the firearm and ammunition in respect of which the licence is granted shall at all times when not in actual use be kept in a secure place with a view to prevent use or access to them by any person not lawfully entitled to use them;

(b) the loss or theft of any firearm to which the licence relates shall be at once reported to the nearest police station or licensing officer;

(c) except as the licensing officer otherwise permits, the firearm and ammunition shall be kept at the permanent address of the holder of the firearm licence and the licensing officer shall be notified in writing within 14 days of any change of such address and of any change in the address at which the holder of the licence is otherwise permitted to keep the firearm and ammunition;

(d) the holder of the licence shall obey all lawful orders of any police officer or licensing officer relating to -

(i) the licence;
(ii) any firearm or ammunition possessed or acquired under the licence,

and shall produce such licence, firearm or ammunition to such officer on demand;

(e) the firearm or ammunition in respect of which the firearm licence is granted shall not be used for any unlawful purpose.

(2) Every certificate given to the holder of a firearm licence shall be retained by him for production to the licenced dealer or, if required, to a police or licensing officer.

FORM OF APPLICATION FOR A FIREARM DEALER'S LICENCE

4. An application for the grant of a firearm dealer's licence under subsection (2) of section 9 of the Act shall be made to the licensing officer in Form 4 in Schedule 1 hereto.

.../3.

FORM OF FIREARM DEALER'S LICENCE

5. Every firearm dealer's licence granted under section 11 of the Act shall be in Form 5 in Schedule 1 hereto and shall be subject to the following conditions:-

(a) that the holder of a firearm dealer's licence or his servants employed in the ordinary course of business of such holder as a licensed firearms and ammunition dealer, shall observe the provisions of the Act and of any regulations made hereunder;

(b) that the holder of a firearm dealer's licence or his servants employed as aforesaid shall not part with possession of any firearm or ammunition except to a person lawfully entitled to possess the same;

(c) that the holder of a firearm dealer's licence or his servants employed as aforesaid shall obey all lawful orders of any police or licensing officer relating to any firearm or ammunition in their possession;

(d) that any firearm and ammunition in the possession of the holder of a firearm dealer's licence shall not be used for any unlawful purpose;

(e) that the holder of such licence shall ensure that the proper records required under the Act and the Regulation are maintained by him in the prescribed form.

PARTICULARS TO BE ENTERED IN THE REGISTER OF THE LICENCED FIREARMS DEALER

6. (1) Every licenced firearms dealer shall enter or cause to be entered in the register of transactions required to be kept under section 11(6) of the Act, the following particulars:-

(a) the quantities and description of firearms and ammunition purchased or acquired with the names and addresses of the sellers or transferors and the dates of the several transactions;

(b) the quantities and description of firearms and ammunition sold or transferred with the names and addresses of the -

.../4.

purchasers and transferees (except in cases where the purchasers are transferees), the areas in which firearms were sold or transferred and the dates of the several transactions;

(c) the quantities and description of firearms and ammunition accepted for sale, repair, test, proof, cleaning, storage, or other purpose, with the names and addresses of the transferors and the dates of the several transaction;

(d) the quantities and description of firearms and ammunition in possession for sale or transfer at the date of the last stock-taking or such other date in each year as may be specified in the register.

(2) The register shall be kept in three books and shall be in Form 6 in Schedule 1 hereto.

(3) Every licenced firearms dealer shall, between the 1st and 7th of each month furnish particulars under section 11(8) of the Act, to the licensing officer in Form 7 in Schedule 1 hereto.

REQUIREMENTS FOR A FIREARMS DEALER'S LICENCE

7. Every licenced firearms dealer selling or parting with possession of a firearm or ammunition to the holder of a firearm licence shall -

(a) fill and sign the certificate of transaction in Form 3 in Schedule 1 hereto;

(b) within 48 hours from the transaction, send a copy of the certificate of the transaction to the licensing officer;

(c) in any case report in writing 48 hours to the licensing officer any circumstances attending the transaction which appear to require investigation.

FORM OF APPLICATION FOR A FIREARMS IMPORT LICENCE

8. An application for the grant of a firearm import licence under subsection (1) of section 14 of the Act shall be made to the licensing officer in Form 8 in Schedule 1 hereto.

.../5.

FORM OF FIREARMS IMPORT LICENCE

9. Every firearm import licence granted under section 14 shall be in Form 9 in Schedule 1 hereto.

REGISTER OF FIREARM IMPORTERS

10. A licensing officer shall, under subsection (2) of section 14 of the Act, keep and maintain a register which shall be in Form 10 in Schedule 1 hereto.

LOST LICENCES

11. (1) Any person who having obtained a replacement licence under section 15 of the Act, in place of the licence he has lost, finds such lost licence shall within 10 days of finding the licence, surrender it to the licensing officer or a police officer.

(2) Any person who contravenes this regulation shall be guilty of an offence and liable to a fine not exceeding 10,000 vatu or to imprisonment for a period not exceeding two months or to both such fine and imprisonment.

DISPOSAL OF DEPOSITED, DETAINED AND FORFEITED FIREARM OR AMMUNITION

12. (1) Any firearm or ammunition detained under the provisions of section 20 of the Act shall, at the expiry of the period of detention, unless previously brought before any court, be returned to the persons lawfully entitled to receive the same. If there is no such person, such firearm or ammunition may be sold or disposed of as the Commissioner of Police deems fit.

(2) Any firearm or ammunition deposited at a police station or police armoury in accordance with the provisions of subsection (1) of section 37 of the Act, if not taken possession of by the lawful owner in accordance with the provisions of sub-section (2) of section 37 and not disposed of in the manner specified by that subsection, shall be disposed of as the Commissioner of Police deems fit.

(3) Where any firearm or ammunition is detained under the provisions of section 36 of the Act and at the expiry of the period of detention, such firearm or ammunition cannot be returned to the person lawfully authorized to receive the same in accordance with section 38 as such person cannot be found,

.../6.1

such firearm or ammunition may be sold or disposed of as the Commissioner of Police deems fit.

(4) Firearms and ammunition forfeited in accordance with the provisions of section 39 of the Act may be sold or disposed of as the Commissioner of Police deems fit.

(5) If any police officer in charge of a police armoury considers any firearm or ammunition deposited or detained therein in accordance with the Act or these Regulations to be in an unsafe condition, he shall report the unsafe condition of such firearm or ammunition to the Commissioner of Police who may cause such firearm or ammunition to be destroyed in such manner as he deems fit.

(6) The proceeds from the sale of any firearm or ammuntiion referred to in this regulation, other than the proceeds of sale under section 37, shall be paid into the Treasury.

VOLUNTARY DEPOSIT OF FIREARMS AND AMMUNITION IN POLICE ARMOURIES

13. (1) Any person in lawfull possession of a firearm or ammunition may deposit such firearm or ammunition in a police armoury for safe custody.

(2) It shall be the duty of the officer in charge of the armoury to take reasonable care of all firearms and ammunition so deposited, but notwithstanding any breach of duty or the negligence of any police officer or otherwise, no action shall lie against the Government or any police officer in respect of the loss of or damage to any firearm or ammuniton so deposited.

FEES

14. (1) The fee payable in respect of any licence referred to in the Act, and in respect of any deposition of firearm in police armoury or of variation of conditions in firearm licences shall be the fees respectively specified in Schedule 2 and shall be paid to the licensing officer.

(2) Any fee payable on the grant of a licence, renewal of a licence, variation of conditions in a licence or deposit of firearm in police armoury shall be paid before the applicant -

.../7.

is granted a licence, renewal, variation of conditions or deposit of firearms, as the case maybe.

COMMENCEMENT

15. These regulations shall come into force on the date of their publication in the Gazette.

MADE at Port Vila, this 1st day of July , 1988.

IOLU J ABBIL
Minister of Home Affairs

SCHEDULE I

FORMS

FORM 1 (regulation 2)

THE FIREARMS REGULATIONS

FORM OF APPLICATION FOR A FIREARM LICENCE

1. Name in full.....

2. Full postal address - Village/Town.....

   Island.....

3. Age.....4. Nationality.....

5. Occupation.....6. Employer.....

7. Business address.....
   .....

8. Description of firearm for which licence required.....

   .....

9. (a) Details of firearms possessed at date of application.....

   .....

   (b) Ammunition -

	Quantity	Calibre and Type
(i) Possessed at date of application.....		.....

.../2.

(ii) Total amount desired to be purchased

or acquired in one year ....

(iii) Maximum amount desired to be possessed

or acquired at any one time ....

10. Whether a licence has previously been held, or applied for by applicant. If so, give details of all previous applications.....

.....

11. Date and place of issue of any licence to possess firearms or ammunition held during previous 3 years.....

.....

12. Do you suffer from any form of mental disorder or defect?

YES/NO. If YES, give details:.....

13. Have you been convicted of any offence, other than minor traffic offences. YES/NO.

If YES, give details:.....

14. Reasons for requiring each of the firearms and ammunition specified:.....

15. Where do you intend to use each of the firearms specified?

16. Where, if a firearm licence is granted will each of the firearms and the ammunition specified be kept when not in use?

DECLARATION

I hereby apply for a licence in respect of the firearm(s) and ammunition specified above, and I declare that the statements made above are true and complete in all respects.

.../3.

Date..... Signature.....

For Official Use

This application was lodged at.....(office)

on (date).....

RECOMMENDED/NOT RECOMMENDED

Reasons for non-recommendation .....

.....

LICENCE Approved/Not Approved
AMMUNITION -

(i) maximum amount authorised to be possessed at any one time*-

.....

(ii) total amount to be purchased in each year*-

.....
*State quantity, type and calibre

Date ..... Signature.....
Licensing Officer

FORM 2 (regulation 3)

THE FIREARMS REGULATIONS

FIREARM LICENCE

LICENCE NO.....

Full Name.....
(Block Letters)

Village/Town .....

Island.....
is hereby authorised to possess the firearm(s) and ammunition specified hereunder -

(a) Firearm -

(i) Type .....

(ii) Calibre.....

(iii) Maker's Name.....

(iv) Maker's Number.....

(b) Ammunition -

(i) maximum amount authorised to be possessed at any one time.*.....

(ii) total amount authorised to be purchased in each year, commencing , 19....*.....
.....

* State quantity, type and calibre.

The following conditions shall be observed by the holder of this licence:-

.../2.

(1) the firearm(s) and ammunition to which this licence relates shall at all times when not in actual use be kept in a secure place with a view to preventing use or access to them by persons not entitled to use them;

(2) the loss or theft of any of the firearm(s) or ammunition to which the licence relates shall be reported at once to the nearest police station or licensing officer;

(3) except the licensing officer otherwise permits, the firearm(s) and ammunition shall be kept at the permanent address of the holder of this licence and the licensing officer shall be notified in writing within.....of any change of such address and of any change in the address at which the holder of this licence is otherwise permitted to keep the firearm(s) or ammunition;

(4) the holder of this licence shall take all necessary steps to secure that regulation 5 of the Firearms Regulation (which relates to instructions concerning the sale or transfer of firarms and ammunition by licensed dealers) is complied with;

(5) the holder of this licence shall obey all lawful orders of any police officer or licensing officer relating to the licence, or any firearm or ammunition possessed or acquired under this licence and shall produce such licence firearm or ammunition to such officer on demand;

(6) that such firearm or ammunition shall not be used for any unlawful purposes.

The licensing officer may insert any other condition herebelow:

This licence shall reamin in force until.....
unless previously revoked or cancelled.

Date:..... Signature:.....
(Licensing Officer)

T.R. No.....

.../3.

FORM 3 (regulations 4)

THE FIREARMS REGULATIONS

CERTIFICATE

TABLE 1 (FIREARMS)

A	B	C	D Certificate
Date of sale or transfer	Name and address of person selling or transferring firearm	Quantity, Calibre, make and type of firearm identification number or other mark	I certify that the entries in columns A to C are correct and relate to a transaction and I have satisfied myself that the transaction will not place the holder of the licence in possession of firearms in excess of or otherwise than as authorised by the licence. Signature..... Date:.....

.../2.

RENEWALS

This licence is hereby renewed and is valid until.....19...,
unless previously revoked or cancelled.

Date..... Signature .....
Licensing Officer

T.R. No.....

FORM 4 (regulation 4)

THE FIREARMS REGULATIONS

FORM OF APPLICATION FOR A FIREARMS DEALER'S LICENCE

1. Name in Full.....

2. Address.....

3. Intended Place of Business.....

4. Description of Firearms and Ammunition for which a licence is required -

(a) Firearms - State Make, Type, Calibre and Maximum Quantity to be held at any one time -.....

.....

.....

(b) Ammunition - State Make, Type, Calibre and Maximum Qunatity to be held at any one time - .....

.....

.....

5. (a) State whether a licence has previously been held or applied for. If so, give dates of all previous applications.

.....

.....

(b) Date and place of issue of any licence to deal in firearms and ammunition held during previous 12 months.....

.....

.../2.

TABLE 2 (AMMUNITION)

A	B	C	D	E Certificate
Date of sale or transfer	Name and address of person selling or transferring ammunition	Quantity of ammunition	Calibre and description of ammunition	I certify that the entries in columns A to D are correct and relate to a transaction with the holder of this licence, that I have inspected this licence and the record of previous transactions and I have satisfied myself that the transaction will not place the holder of the licence in possession of ammunition in excess of or otherwise than as authorised by the licence. Signature:..... Date:.....

FORM 5 (regulation 5)

THE FIREARM REGULATIONS

FIREARM DEALER'S LICENCE

LICENCE No:.....

This licence authorises.....

of .....in the Republic of Vanuatu or his servants in the ordinary course of his business as a licensed firearms and ammunition dealer -

(a) to assemble or disassemble, clean, repair, test or approve any firearm or ammunition;

(b) to manufacture any component part of a firearm or ammunition;

(c) to sell, transfer or expose for sale any firearm or ammunition;

(d) to keep or have in their possession any firearm or ammunition for any of the aforesaid purposes at the following place(s) - .....

.....

This licence is subject to the following conditions -

(i) that the holder of such licence or his servants employed in the ordinary course of business of such holder as a licensed firearms and ammunition dealer, shall observe the provisions of the Act and of any regulations made hereunder;

(ii) that the holder of such licence or his servants employed as aforesaid shall not part with possession of any firearm or ammunition except to a person lawfully entitled to possess the same;

(iii) that the holder of such licence or his servants employed as aforesaid shall obey all lawful orders of any police or licensing officer relating to any firearm or ammunition in their possession;

.../2.

DECLARATION

I hereby apply for a licence to deal in the firearms and ammunition specified above and I declare that the statements made above are true and complete in all respects.

Date.....Signature of Applicant.....

For official Use

This application was lodged at .....(office)

on (date).....

RECOMMENDED/NOT RECOMMENDED

Reasons for non-recommendation.....

.....

LICENCE Approved/Not Approved

Date..... Signature.....
Licensing Officer

FORM 6 (regulation 6)

FIREARMS REGULATIONS

DEALER'S REGISTER OF TRANSACTIONS

TABLE 1 - FIREARMS

(A separate folio to be used for each type of firearms)

Type.....

ACQUISITION

Date	Source (in case of import state country), and full name and address of firm or person from whom acquired	Make, calibre and identity No.	No. and date of import licence	Quantity

DISPOSAL

Date	Name and address of person to whom sold or supplied	Make, calibre and and quantity sold or disposed of and identity No.	Balance in stock

.../2.

(iv) that any firearm and ammunition in the possession of the holder of such licence shall not be used for any unlawful purpose;

(v) that the holder of such licence shall ensure that the proper records required under the Act and the Regulation are maintained by him in the prescribed form;

(vi) Additional conditions .....

.....

This licence shall continue in force until ....., 19.... unless previously revoked or cancelled.

Date..... Signature.....
Licensing Officer

T.R. No.....

RENEWALS

This licence is hereby renewed and shall remain in force until.....

..... 19...., unless previously revoked or cancelled.

Date:..... Signature:.....
(Licensing Officer)

T.R. No:.....

TABLE 2 - AMMUNITION

Calibre .....(A separate folio to be used for each calibre of Ammunition)

ACQUISITION

Date	Source (in case of import state country), and full name and address of firm or person from whom acquired	Make, type and quantity received	Number and date of Import Licence

DISPOSAL

Date	Make, type and quantity sold	Name and address of person to whom sold	Balance in Stock

.../3.

TABLE 3 - REPAIRS, ALTERATIONS, STORAGE, TESTING, ETC.

1	2	3	4	5
Date	Full name and address of depositor	Reason for depositing	Make, type, calibre and identity markings of firearms or component part deposited	Quantity and calibre of ammunition deposited

6	7	8	9
Date Removed	By whom removed (full name and address)	Firearms licence No. date and station of issue produced by person removing	Remarks

FORM 7 (regulaltion 6(3))

THE FIREAMRS REGULATIONS

RETURN OF FIREARMS AND AMMUNITION PURCHASED, IMPORTED AND SOLD

(To be forwarded by licensed firearm's dealers to the licensing officer between the 1st and 7th of each month)

TABLE 1 – FIREARMS

Month of..... 19.....	.22 Rifles	12g. Shotguns	.410 Shotguns	Other Shotguns
Stock in hand on				
Purchased or imported* on.....				
" "				
" "				
Sold on				
" "				
" "				
Balance in Stock on				

*Where imported also include Import Licence Number.

.../2.

TABLE 2 – AMMUNITION

Month of..... 19.....	.22 Rifles	12g Shotguns	.140 Shotguns	Other Shotguns
Stock in hand on				
Purchased or imported* on.....				
" "				
" "				
Sold on				
" "				
" "				
Balance in stock on				

*Where imported also record Import Licence Number

FORM 8 (regulation 8)

THE FIREARMS REGULATIONS

FORM OF APPLICATION FOR A FIREARM IMPORT LICENCE

1. Name in Full .....

2. Address.....

3. Place of Business .....

4. Description of Firearms and Ammunition desired to be imported -

   (a) Firearms - State make, type, calibre and quantity of firearms to be imported.....

   .....

   .....

   (b) Ammunition - State make, type, calibre and quantity to be imported.....

   .....

   .....

5. (a) State whether a licence has previously been held or applied for. If so, give dates of all previous applications.....

   .....

   .....

   (b) Date and place of issue of any licence to import firearms and ammunition granted during last 12 months.....
   .....

6. Details of firearms and ammunition possessed at date of this application. (State quantity, make, calibre, type, serial No. or other distinguishing mark, and exporter). If none, say NONE.

.../2.

7. Reasons for requiring the firearms and ammunition specified above:

.....

.....

DECLARATION

I hereby apply for a licence to import the firearms and ammunition specified above. I am the holder of a Firearm Dealer's Licence No.1* I declare that the statements made above are true and complete in all respects.

*Delete if inapplicable.

Date..... Signature of Applicant.....

For Official Use

This application was lodged at.....(office)

on (date).....

RECOMMENDED/NOT RECOMMENDED

Reasons for non-recommendation.....

.....

LICENCE Approved/Not approved

Date..... Signature:.....
Licensing Officer

REPUBLIC OF VANUATU

FORM 9 (regulation 9)

THE FIREARMS REGULATIONS

FIREARMS IMPORT LICENCE

LICENCE No.....

Full Name.....

Address.....

is hereby licensed to import into the Republic of Vanuatu firearms and ammunition of the number and description hereunder specified -

Quantity.	Description	Calibre	Remarks

This licence shall continue in force until .....19..... unless previously revoked or cancelled.

Date ..... Signature.....
Licensing Officer

T.R. No.....

FORM 10 (regulation 10)

FIREARMS REGULATIONS

REGISTER OF FIREARMS IMPORTERS

Name and place of business	Date application received	Date application approved/ not approved	Date licence issued	Licence No:	T.R.No.	Date Renewed

SCHEDULE 2 (regulation 14)

FEES

Particulars	Fees
1. Grant or renewal of firearm licence under section 9(2) of the Act.	3,000 vatu.
2. Variation of conditions in firearm licence under section 9(8) of the Act.	1,000 vatu.
3. Grant or renewal of firearm dealer's licence under section 11(3) of the Act.	30,000 vatu
4. Grant or renewal of firearm import licence under section 14(3) of the Act	500 vatu per firearm.
5. Issue of replacement firearm licence under section 15 of the Act	1,000 vatu.
6. Deposit of firearm in police armoury under section 37(1) of the Act.	1,000 vatu per firearm

REPUBLIC OF VANUATU

THE FIREARMS ACT No. 7 OF 1987

NOTICE

IN EXERCISE of the powers conferred by section 17 of the Firearms Act No.7 of 1987, I HEREBY DELCARE -

(1) That no firearms or ammunition whatsoever shall be imported into Vanuatu except at the Overseas wharf in Port Vila and Bauerfield Airport in Port Vila;

(2) That the following firearms and ammunition shall not be imported into Vanuatu:-

(a) automatic firearms, that is firearms which are so designed or adapted that if pressure is applied to the trigger, missiles continue to be discharged until pressure is removed from the trigger or the magazine containing the missiles is empty;

(b) semi-automatic firearms, that is firearms other than automatic firearms as specified in paragraph (a), which when fired eject the spent round and refill the breach;

(c) pistols and revolvers of all types;

(d) firearms and ammunition of the following calibres:-

.300, .303, .38 and .45 inches, and 7.62 and 9 millimetres;

(e) any weapon of whatever description designed or adapted for the discharge of any noxious liquid, gas or other thing; and

(f) any ammunition containing or designed or adapted to contain any such noxious thing.

This NOTICE shall come into force on the date of its publication in the Gazette.

.../2.

MADE at Port Vila this 1st day of July, 1988.

IOLU J ABBIL
Minister of Home Affairs

REPUBLIQUE DE VANUATU

ARRETE NO. 14 DE 1988 SUR LES TELECOMMUNICATIONS (TAXES ET REDEVANCES) (MODIFICATION)

portant modification de l'arrêté No. 19 de 1983 sur les télécommunications (taxes et redevances).

LE MINISTRE DE L'AVIATION CIVILE, DES COMMUNICATIONS, DE LA SYLVICULTURE ET DE L'ENERGIE

VU les disposition de l'article 26 de la loi No. 26 de 1982 sur les télécommunications, .

A R R E T E :

MODIFICATION DE L'ARRETE NO. 19 DE 1983

1. L'arrêté No. 19 de 1983 sur les télécommunications (taxes et redevances) tel que modifié est à nouveau modifié comme suit :

(i) Au titre D de l'annexe 1 en ajoutant le nouvel article qui suit immédiatement après le paragraphe 2(b) :

"2A Pour un abonné résidant en dehors des zones urbaines de la municipalité de Port-Vila ou de Luganville ou pour les usagers de son service téléphonique, la taxe maximum exigible :

(a) des communications entre les abonnés reliés au même central téléphonique :

(i) pour les trois premières minutes, ou pour toute fraction de ce temps ..... 120 VT ;

(ii) par minute supplémentaire ou pour toute fraction de ce temps ..... 40 VT ;

(b) des communications avec un abonné relié à un central téléphonique différent, à l'intérieur de Vanuatu :

(i) pour les trois premières minutes ou pour toute fraction de ce temps ..... 300 VT ;

(ii) par minute supplémentaire ou pour toute fraction de ce temps ..... 20 VT,

étant entendu que pour les communications destinées à une personne autre que l'abonné, par le service téléphonique de ce dernier, aucune taxe ou redevance ne peut être imposée à l'abonné.

(ii) A l'annexe 2, en ajoutant le nouveau paragraphe qui suit immédiatement après le paragraphe (g)(iv)

"(v) pour tous les émetteurs VHF mobiles, portatifs et à poignée utilisant un relais des Postes et Télécommunications ..... 8.000 VT.

ENTREE EN VIGUEUR

2. Le présent arrêté entrera en vigueur le jour de sa publication au Journal officiel.

FAIT à Port-Vila, le 18 avril 1988.

H.C. QUALAO
Ministre de l'Aviation civile, des Communications, de la Sylviculture et de l'Energie.