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POISONS ACT 1964

POISONS (SCHEDULED SUBSTANCES) AMENDMENT ORDER 1984

POISONS ACT 1964.

POISONS (SCHEDULED SUBSTANCES) AMENDMENT ORDER 1984.

MADE by His Excellency the Governor in Executive Council.

1. This Order may be cited as the Poisons (Scheduled Substances) Amendment Order 1984.

2. This Order shall come into operation 1 month after the date of publication of this Order in the *Government Gazette*.

3. Appendix "A" to the Poisons Act 1964 as amended is repealed and the following Appendix is substituted—

"APPENDIX "A"

A substance specified in a schedule, unless the contrary intention appears, includes—

(a) every salt, active principle or derivative of the substance and every salt of such an active principle or derivative;

(b) except where the substance is opium, every alkaloid of the substance and every salt of such an alkaloid;

(c) except where the substance is levomethorphan or levorphanol — every stereoisomer of the substance and every salt of such a stereoisomer;

(d) a preparation or admixture containing any proportion thereof of the substance; and

(e) in the case of an Eighth Schedule substance every ester and ether of the substance and every salt of such an ester or ether.

First Schedule

ACONITE (ROOT OF ACONITUM NAPELLUS).

ANTIMONY, compounds of, except —

(a) antimony chloride in polishes;

(b) when included in the Fourth Schedule.

ATROPINE, except —

(a) when included in the Second Schedule;

(b) atropine methonitrate.

BELLADONNA HERB, except when included in the Second Schedule.

BROMINE (excluding its salts and derivatives).

BRUCINE, except when used in concentrations of 0.02 per cent or less for the denaturation of alcohol.

CONIINE.

COTARNINE.

CROTON OIL.

CYANIDES — see hydrocyanic acid.

HOMATROPINE, except when included in the Second Schedule.

HYDROCYANIC ACID for therapeutic use except when included in the Second Schedule.

HYOSCINE, except —

(a) when included in the Second Schedule;

(b) hyoscine butylbromide.

HYOSCYAMINE, except when included in the Second Schedule.

HYOSCYAMUS, except when included in the Second Schedule.

LOBELIA, except —

(a) in preparations for smoking or burning;

(b) when included in the Second Schedule.

MERCURIC CHLORIDE, except —

(a) in batteries;

(b) when included in the Second or Seventh Schedule.

MERCURIC IODIDE, except when included in the Second or Sixth Schedule.

MERCURIC NITRATE, except when included in the Second Schedule.

MERCURIC-POTASSIUM IODIDE, except when included in the Second Schedule.

MERCURIC THIOCYANATE, except when included in the Sixth Schedule.

MERCURY, organic compounds of, except —

(a) for therapeutic use;

(b) when included in the Second, Sixth or Seventh Schedule.

NUX VOMICA.

PHOSPHORUS YELLOW (excluding its salts and derivatives), except when included in the Sixth Schedule.

SAVIN, oil of.

STRAMONIUM, except —

(a) in preparations for smoking or burning;

(b) when included in the Second Schedule.

TANSY, oil of.

VERATRUM, except for therapeutic use.

Excluding however, the substances hereinbefore mentioned when contained in the products listed as exemptions, set out after the Eighth Schedule in this Appendix.

Second Schedule.

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

ACETYLDIHYDROCODEINE when compounded with one or more other medicaments, in preparations containing 1 per cent or less of acetyldihydrocodeine.

AMMONIATED MERCURY.

ANTAZOLINE in preparations labelled and packed as eye drops or as nasal preparations for topical use.

ASPIRIN and its preparations and derivatives, except—

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

(i) the pack is labelled with either of the following warning statements—

" 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. . ";

(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;

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

(i) the pack is labelled with either of the following warning statements—

" 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. ";

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

(c) when included in the Fourth Schedule.

ATROPINE, except atropine methonitrate, in preparations containing 0.25 per cent or less of atropine and atropine sulphate, 0.6 mg tablets in packs of 6, when labelled for treatment of organophosphorus poisoning.

BAMIPINE in preparations labelled and packed as eye drops or as nasal preparations for topical use.

BELLADONNA HERB 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 and capsules containing 30 mg or less of benzamine in each;

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

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

BENZOCAINE when included in—

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

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

(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.

BROMHEXINE

BROMODIPHENHYDRAMINE—

(a) in preparations labelled and packed as eye drops or as nasal preparations for topical use;

(b) in oral liquid preparations when compounded with an antitussive, an expectorant or a sympathomimetic amine and when packed, labelled and sold for the relief of coughs and colds, not containing codeine and not indicating a dosage for children under 2 years of age.

BROMPHENIRAMINE—

(a) in preparations labelled and packed as eye drops or as nasal preparations for topical use;

(b) in oral liquid preparations when compounded with an antitussive, an expectorant or a sympathomimetic amine and when packed, labelled and sold for the relief of coughs and colds, not containing codeine and not indicating a dosage for children under 2 years of age.

BUCLIZINE—

(a) in preparations labelled and packed for the treatment of motion sickness in packs of 10 doses or less;

(b) in preparations labelled and packed as eye drops or as nasal preparations for topical use;

(c) in oral liquid preparations when compounded with an antitussive, an expectorant or a sympathomimetic amine and when packed, labelled and sold for the relief of coughs and colds, not containing codeine and not indicating a dosage for children under 2 years of age.

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

BUTYLAMINOBENZOATE when included in—

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

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

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

CANTHARIDIN in preparations containing 0.01 per cent or less of cantharidin.

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.

CARBINOXAMINE—

(a) in preparations labelled and packed as eye drops or as nasal preparations for topical use;

(b) in oral liquid preparations when compounded with an antitussive, an expectorant or a sympathomimetic amine and when packed, labelled and sold for the relief of coughs and colds, not containing codeine and not indicating a dosage for children under 2 years of age.

CHLOROFORM (excluding its derivatives), except—

(a) in preparations containing 10 per cent or less of chloroform where the chloroform content is declared on the label;

(b) when included in the Fourth Schedule.

CHLOROPYRILENE —

(a) in preparations labelled and packed as eye drops or as nasal preparations for topical use;

(b) in oral liquid preparations when compounded with an antitussive, an expectorant or a sympathomimetic amine and when packed, labelled and sold for the relief of coughs and colds, not containing codeine and not indicating a dosage for children under 2 years of age.

CHLORPHENIRAMINE—

(a) in preparations labelled and packed as eye drops or as nasal preparations for topical use;

(b) in oral liquid preparations when compounded with an antitussive, an expectorant or a sympathomimetic amine and when packed, labelled and sold for the relief of coughs and colds, not containing codeine and not indicating a dosage for children under 2 years of age.

CHLORPHENOXAMINE —

(a) in preparations labelled and packed as eye drops or as nasal preparations for topical use;

(b) in oral liquid preparations when compounded with an antitussive, an expectorant or a sympathomimetic amine and when packed, labelled and sold for the relief of coughs and colds, not containing codeine and not indicating a dosage for children under 2 years of age.

CINNAMEDRINE.

CINNARIZINE —

(a) in preparations labelled and packed as eye drops or as nasal preparations for topical use;

(b) in solid dose preparations labelled and packed for the treatment of motion sickness, in packs of 10 doses or less;

(c) in oral liquid preparations when compounded with an antitussive, an expectorant or a sympathomimetic amine and when packed, labelled and sold for the relief of coughs and colds, not containing codeine and not indicating a dosage for children under 2 years of age.

CLEMASTINE —

(a) in preparations labelled and packed as eye drops or as nasal preparations for topical use;

(b) in oral liquid preparations when compounded with an antitussive, an expectorant or a sympathomimetic amine and when packed, labelled and sold for the relief of coughs and colds, not containing codeine and not indicating a dosage for children under 2 years of age.

(c) in oral dosage units of 1 mg or less in packs of 50 doses or less.

CLEMIZOLE—

(a) in preparations labelled and packed as eye drops or as nasal preparations for topical use;

(b) in oral liquid preparations when compounded with an antitussive, an expectorant or a sympathomimetic amine and when packed, labelled and sold for the relief of coughs and colds, not containing codeine and not indicating a dosage for children under 2 years of age.

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

CODEINE —

(a) when compounded in tablets or capsules with aspirin or paracetamol or salicylamide or one of their derivatives, and no other therapeutically active substance, and containing not more than 10 mg of codeine per tablet or capsule when—

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

(ii) in a primary pack containing not more than 25 such tablets or capsules;

(b) when compounded with aspirin or paracetamol or salicylamide in individually wrapped powders containing 10 mg or less of codeine in each individually wrapped powder where the powders are enclosed in a primary pack containing not more than 12 individually wrapped powders;

(c) when compounded with one or more other medicaments in divided preparations containing not more than 10 mg codeine per dosage unit;

(d) when compounded with one or more other medicaments in undivided preparations with a concentration of not more than 0.5 per cent of codeine.

CYANIDES — see hydrocyanic acid.

CYCLIRAMINE —

(a) in preparations labelled and packed as eye drops or as nasal preparations for topical use;

(b) in oral liquid preparations when compounded with an antitussive, an expectorant or a sympathomimetic amine and when packed, labelled and sold for the relief of coughs and colds, not containing codeine and not indicating a dosage for children under 2 years of age.

CYPROPHEPTADINE —

(a) in preparations labelled and packed as eye drops or as nasal preparations for topical use;

(b) in oral liquid preparations when compounded with an antitussive, an expectorant or a sympathomimetic amine and when packed, labelled and sold for the relief of coughs and colds, not containing codeine and not indicating a dosage for children under 2 years of age.

DDT in preparations for human therapeutic use.

DEPTROPINE —

(a) in preparations labelled and packed as eye drops or as nasal preparations for topical use;

(b) in oral liquid preparations when compounded with an antitussive, an expectorant or a sympathomimetic amine and when packed, labelled and sold for the relief of coughs and colds, not containing codeine and not indicating a dosage for children under 2 years of age.

DEXBROMPHENIRAMINE —

(a) in preparations labelled and packed as eye drops or as nasal preparations for topical use;

(b) in oral liquid preparations when compounded with an antitussive, an expectorant or a sympathomimetic amine and when packed, labelled and sold for the relief of coughs and colds, not containing codeine and not indicating a dosage for children under 2 years of age.

DEXCHLORPHENIRAMINE —

(a) in preparations labelled and packed as eye drops or as nasal preparations for topical use;

(b) in oral liquid preparations when compounded with an antitussive, an expectorant or a sympathomimetic amine and when packed, labelled and sold for the relief of coughs and colds, not containing codeine and not indicating a dosage for children under 2 years of age.

DEXTROMETHORPHAN in preparations containing 1 per cent or less of dextromethorphan when compounded with one or more other medicaments in such a way that the dextromethorphan contained therein cannot readily be extracted.

DEXTRORPHAN in preparations containing 1 per cent or less of dextrorphan.

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

DIMENHYDRINATE —

(a) in preparations labelled and packed for the treatment of motion sickness in packs of 10 doses or less;

(b) in preparations labelled and packed as eye drops or as nasal preparations for topical use;

(c) in oral liquid preparations when compounded with an antitussive, an expectorant or a sympathomimetic amine and when packed, labelled and sold for the relief of coughs and colds, not containing codeine and not indicating a dosage for children under 2 years of age.

DIMETHINDENE —

(a) in preparations labelled and packed as eye drops or as nasal preparations for topical use;

(b) in oral liquid preparations when compounded with an antitussive, an expectorant or a sympathomimetic amine and when packed, labelled and sold for the relief of coughs and colds, not containing codeine and not indicating a dosage for children under 2 years of age.

DIMETHISOQUIN in preparations for topical use.

DIMETHOTHIAZINE —

(a) in preparations labelled and packed as eye drops or as nasal preparations for topical use;

(b) in oral liquid preparations when compounded with an antitussive, an expectorant or a sympathomimetic amine and when packed, labelled and sold for the relief of coughs and colds, not containing codeine and not indicating a dosage for children under 2 years of age.

DIPHEMANIL METHYLSULPHATE in preparations for topical use.

DIPHENHYDRAMINE —

(a) in preparations labelled and packed for the treatment of motion sickness in packs of 10 doses or less;

(b) in preparations labelled and packed as eye drops or as nasal preparations for topical use;

(c) in oral liquid preparations when compounded with an antitussive, an expectorant or a sympathomimetic amine and when packed, labelled and sold for the relief of coughs and colds, not containing codeine and not indicating a dosage for children under 2 years of age.

DIPHENYLPYRALINE —

(a) in preparations labelled and packed as eye drops or as nasal preparations for topical use;

(b) in oral liquid preparations when compounded with an antitussive, an expectorant or a sympathomimetic amine and when packed, labelled and sold for the relief of coughs and colds, not containing codeine and not indicating a dosage for children under 2 years of age.

DOXYLAMINE—

(a) in preparations labelled and packed as eye drops or as nasal preparations for topical use;

(b) in oral liquid preparations when compounded with an antitussive, an expectorant or a sympathomimetic amine and when packed, labelled and sold for the relief of coughs and colds, not containing codeine and not indicating a dosage for children under 2 years of age.

EMBRAMINE in preparations labelled and packed as eye drops, or as nasal preparations for topical use.

EPHEDRINE AND PSEUDOEPHEDRINE, except—

(a) in preparations containing 10 mg or less per dosage unit of ephedrine or pseudoephedrine;

(b) in preparations for external use containing 1 per cent or less of ephedrine or pseudoephedrine;

(c) when included in the Third Schedule.

ERYTHRITYL TETRANITRATE and other nitric esters of polyhydric alcohols.

ETAFEDRINE.

ETHER (excluding its derivatives) except—

(a) in preparations containing 10 per cent or less of ether;

(b) when included in the Fourth, Fifth or Sixth Schedule.

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

ETHYLMORPHINE when compounded with one or more other medicaments, in preparations containing 1 per cent or less of ethylmorphine.

FERROUS SULPHATE and other iron preparations for human internal use, except in preparations containing 5 per cent or less of iron.

FLUORIDES in—

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

(b) other metallic fluoride substances, including ammonium flouride when intended for therapeutic purposes, except—

(i) in dentrifices containing 1 000 mg/kg or less of flouride ion;

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

GELSEMIUM.

GLUTARALDEHYDE for human therapeutic use.

GLYCERYL TRINITRATE except when included in the Fourth Schedule.

GUAIPHENESIN—

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

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

HALOPYRAMINE—

(a) in preparations labelled and packed as eye drops or as nasal preparations for topical use;

(b) in oral liquid preparations when compounded with an antitussive, an expectorant or a sympathomimetic amine and when packed, labelled and sold for the relief of coughs and colds, not containing codeine and not indicating a dosage for children under 2 years of age.

HEXACHLOROPHANE in preparations for skin cleansing purposes containing 3 per cent or less of hexachlorophane, except—

(a) in preparations for use on infants;

(b) in preparations for the treatment of animals;

HISTAPYRRODINE—

(a) in preparations labelled and packed as eye drops or as nasal preparations for topical use;

(b) in oral liquid preparations when compounded with an antitussive, an expectorant or a sympathomimetic amine and when packed, labelled and sold for the relief of coughs and colds, not containing codeine and not indicating a dosage for children under 2 years of age.

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

HYDROCYANIC ACID in preparations containing the equivalent of 0.15 per cent or less of hydrocyanic acid.

8-HYROXYQUINOLINE non-halogenated derivatives, for human therapeutic use, except substances containing 1 per cent or less for external use.

HYOSCINE in preparations containing 0.25 per cent or less of hyoscine, except hyoscine butylbromide.

HYOSCYAMINE in preparations containing 0.25 per cent or less hyoscyamine.

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

IODINE (excluding its salts and derivatives), except—

(a) when included in the Sixth Schedule;

(b) in iodophors in preparations containing 1.5 per cent or less of available iodine;

(c) in solid or semi-solid preparations containing 2.5 per cent or less of available iodine;

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

ISOSORBIDE DINITRATE.

LEAD SALTS and compounds of lead when prepared for medical use.

LIGNOCAINE when included in—

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

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

(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 the alkaloids of lobelia, except preparations for smoking or burning.

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

MEBENDAZOLE for human therapeutic use.

MEPYRAMINE—

(a) in preparations labelled and packed as eye drops or as nasal preparations for topical use;

(b) in oral liquid preparations when compounded with an antitussive, an expectorant or a sympathomimetic amine and when packed, labelled and sold for the relief of coughs and colds, not containing codeine and not indicating a dosage for children under 2 years of age.

MERCURIC CHLORIDE in preparations containing 0.5 per cent or less of mercuric chloride, except—

(a) in batteries;

(b) when included in the Seventh Schedule.

MERCURIC IODIDE in preparations containing 2 per cent or less of mercuric iodide, except when included in the Sixth Schedule.

MERCURIC NITRATE in preparations containing the equivalent of 3 per cent or less of mercury (Hg), in such form.

MERCURIC OXIDE and all oxides of mercury.

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

MERCURY (METALLIC) (excluding its salts and derivatives) except in scientific instruments.

MERCURY organic compounds of, in preparations containing the equivalent of 0.5 per cent or less of mercury (Hg) except—

(a) when included in the Sixth or Seventh Schedule;

(b) as a preservative in substances containing 0.01 per cent or less of mercury.

METHDILAZINE —

(a) in preparations labelled and packed as eye drops or as nasal preparations for topical use;

(b) in oral liquid preparations when compounded with an antitussive, an expectorant or a sympathomimetic amine and when packed, labelled and sold for the relief of coughs and colds, not containing codeine and not indicating a dosage for children under 2 years of age.

METHOXAMINE, except—

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

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

METHOXYPHENAMINE.

METHYLEPHEDRINE.

NAPHAZOLINE.

NICLOSAMIDE for human therapeutic use.

NICOCODINE when compounded with one or more other medicaments, in preparations containing 1 per cent or less of nicocodine.

NICODICODINE when compounded with one or more other medicaments in preparations containing 1 per cent or less of nicodicodine.

NORCODEINE when compounded with one or more other medicaments, in preparations containing 1 per cent or less of norcodeine.

OXETHAZAINE in preparations for internal use only.

OXOLAMINE.

OXYMETAZOLINE.

PAPAVERINE.

PARACETAMOL and its preparations and derivatives, except—

(a) tablets or capsules each containing 500 mg or less of paracetamol as the only therapeutically active constituent when—

(i) the pack is labelled with either of the following warning statements—

" 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. ";

(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;

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

(i) the pack is labelled with either of the following warning statements—

" 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. ";

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

(c) when included in the Fourth Schedule.

PHEDRAZINE.

PHENAMAZOLINE.

PHENAZONE for external use.

PHENINDAMINE —

(a) in preparations labelled and packed as eye drops or as nasal preparations for topical use;

(b) in oral liquid preparations when compounded with an antitussive, an expectorant or a sympathomimetic amine and when packed, labelled and sold for the relief of coughs and colds, not containing codeine and not indicating a dosage for children under 2 years of age.

PHENIRAMINE—

(a) in preparations labelled and packed for the treatment of motion sickness in packs of 10 doses or less;

(b) in preparations labelled and packed as eye drops or as nasal preparations for topical use;

(c) in oral liquid preparations when compounded with an antitussive, an expectorant or a sympathomimetic amine and when packed, labelled and sold for the relief of coughs and colds, not containing codeine and not indicating a dosage for children under 2 years of age.

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

PHENYLENE DIAMINES, toluene and all other alkylated benzene diamine derivatives, except when included in the Sixth Schedule.

PHENYLEPHRINE, except—

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

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

PHENYLPROPANOLAMINE in tablets or capsules containing not more than 30 mg of phenylpropanolamine per tablet or capsule when not in sustained release form, and not more than 50 mg in each tablet or capsule when in sustained release form.

PHENYLTOLOXAMINE—

(a) in preparations labelled and packed as eye drops or as nasal preparations for topical use;

(b) in oral liquid preparations when compounded with an antitussive, an expectorant or a sympathomimetic amine and when packed, labelled and sold for the relief of coughs and colds, not containing codeine and not indicating a dosage for children under 2 years of age.

PHOLCODINE when compounded with one or more other medicaments, in preparations containing 1 per cent or less of pholcodine.

POTASSIUM CHLORATE, 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 preparations labelled and packed for the treatment of motion sickness in packs of 10 doses or less;

(b) in preparations labelled and packed as eye drops or as nasal preparations for topical use;

(c) in oral liquid preparations when compounded with an antitussive, an expectorant or a sympathomimetic amine and when packed, labelled and sold for the relief of coughs and colds, not containing codeine and not indicating a dosage for children under 2 years of age.

PROPANTHELINE in preparations for topical use.

PROPOXUR in preparations for external human therapeutic use containing 0.2 per cent or less of propoxur.

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

PROPYPHENAZONE.

PYRANTEL for human therapeutic use.

PYRITHIONE ZINC, except in preparations containing 2 per cent or less of pyrithione zinc.

PYRROBUTAMINE—

(a) in preparations labelled and packed as eye drops or as nasal preparations for topical use;

(b) in oral liquid preparations when compounded with an antitussive, an expectorant or a sympathomimetic amine and when packed, labelled and sold for the relief of coughs and colds, not containing codeine and not indicating a dosage for children under 2 years of age.

SALICYLAMIDE and its preparations and derivatives except—

(a) tablets or capsules each containing 500 mg or less of salicylamide as the only therapeutically active constituent when—

(i) the pack is labelled with either of the following warning statements—

" 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. ";

(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;

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

(i) the pack is labelled with either of the following warning statements—

" 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. ";

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

(c) when included in the Fourth Schedule.

SILVER NITRATE.

SODIUM NITRITE for therapeutic use.

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.

THENALIDINE—

(a) in preparations labelled and packed as eye drops or as nasal preparations for topical use;

(b) in oral liquid preparations when compounded with an antitussive, an expectorant or a sympathomimetic amine and when packed, labelled and sold for the relief of coughs and colds, not containing codeine and not indicating a dosage for children under 2 years of age.

THENYLDIAMINE—

(a) in preparations labelled and packed as eye drops or as nasal preparations for topical use;

(b) in oral liquid preparations when compounded with an antitussive, an expectorant or a sympathomimetic amine and when packed, labelled and sold for the relief of coughs and colds, not containing codeine and not indicating a dosage for children under 2 years of age.

TOLPROPAMINE—

(a) in preparations labelled and packed as eye drops or as nasal preparations for topical use;

(b) in oral liquid preparations when compounded with an antitussive, an expectorant or a sympathomimetic amine and when packed, labelled and sold for the relief of coughs and colds, not containing codeine and not indicating a dosage for children under 2 years of age.

TRAMAZOLINE.

TRIMEPRAZINE—

(a) in preparations labelled and packed as eye drops or as nasal preparations for topical use;

(b) in oral liquid preparations when compounded with an antitussive, an expectorant or a sympathomimetic amine and when packed, labelled and sold for the relief of coughs and colds, not containing codeine and not indicating a dosage for children under 2 years of age.

TRIMETHOBENZAMIDE—

(a) in preparations labelled and packed as eye drops or as nasal preparations for topical use;

(b) in oral liquid preparations when compounded with an antitussive, an expectorant or a sympathomimetic amine and when packed, labelled and sold for the relief of coughs and colds, not containing codeine and not indicating a dosage for children under 2 years of age.

TRIMIZOLINE.

TRIPELENNAMINE—

(a) in preparations labelled and packed as eye drops or as nasal preparations for topical use;

(b) in oral liquid preparations when compounded with an antitussive, an expectorant or a sympathomimetic amine and when packed, labelled and sold for the relief of coughs and colds, not containing codeine and not indicating a dosage for children under 2 years of age.

TRIPROLIDINE—

(a) in preparations labelled and packed as eye drops or as nasal preparations for topical use;

(b) in oral liquid preparations when compounded with an antitussive, an expectorant or a sympathomimetic amine and when packed, labelled and sold for the relief of coughs and colds, not containing codeine and not indicating a dosage for children under 2 years of age.

TYMAZOLINE.

XYLOMETAZOLINE.

Excluding however, the substances hereinbefore mentioned when contained in the products listed as exemptions, set out after the Eighth Schedule in this Appendix.

Third Schedule

ADRENALINE in preparations containing 1 per cent or less of adrenaline, except in preparations containing 0.01 per cent or less of adrenaline.

AMYL NITRITE.

ANTAZOLINE—

(a) in oral solid preparations containing 30 dosage units or less, except when included in the Second Schedule;

(b) in oral liquid preparations, except when included in the Second Schedule.

BAMIPINE—

(a) in oral solid preparations containing 30 dosage units or less, except when included in the Second Schedule;

(b) in oral liquid preparations, except when included in the Second Schedule.

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

BROMIDES, inorganic, in extemporaneous preparations for therapeutic use.

BROMODIPHENHYDRAMINE—

(a) in oral solid preparations containing 30 dosage units or less, except when included in the Second Schedule;

(b) in oral liquid preparations, except when included in the Second Schedule.

BROMPHENIRAMINE—

(a) in oral solid preparations containing 30 dosage units or less, except when included in the Second Schedule;

(b) in oral liquid preparations, except when included in the Second Schedule.

BUCLIZINE—

(a) in oral solid preparations containing 30 dosage units or less, except when included in the Second Schedule;

(b) in oral liquid preparations, except when included in the Second Schedule.

BUTYL NITRITE.

CARBINOXAMINE—

(a) in oral solid preparations containing 30 dosage units or less, except when included in the Second Schedule;

(b) in oral liquid preparations, except when included in the Second Schedule.

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

CHLOROPYRILENE—

(a) in oral solid preparations containing 30 dosage units or less, except when included in the Second Schedule;

(b) in oral liquid preparations, except when included in the Second Schedule.

CHLORPHENIRAMINE—

(a) in oral solid preparations containing 30 dosage units or less, except when included in the Second Schedule;

(b) in oral liquid preparations, except when included in the Second Schedule.

CHLORPHENOXAMINE—

(a) in oral solid preparations containing 30 dosage units or less, except when included in the Second Schedule;

(b) in oral liquid preparations, except when included in the Second Schedule.

CINNARIZINE—

(a) in oral solid preparations containing 30 dosage units or less, except when included in the Second Schedule;

(b) in oral liquid preparations, except when included in the Second Schedule.

CLEMASTINE—

(a) in oral solid preparations containing 30 dosage units or less, except when included in the Second Schedule;

(b) in oral liquid preparations, except when included in the Second Schedule.

CLEMIZOLE—

(a) in oral solid preparations containing 30 dosage units or less, except when included in the Second Schedule;

(b) in oral liquid preparations, except when included in the Second Schedule.

CODEINE when compounded with aspirin or paracetamol or salicylamide in tablets or capsules containing 10 mg or less of codeine in each tablet or capsule or in individually wrapped powders containing 10 mg or less of codeine in each individually wrapped powder, except when included in the Second Schedule.

CYCLIRAMINE—

(a) in oral solid preparations containing 30 dosage units or less, except when included in the Second Schedule;

(b) in oral liquid preparations, except when included in the Second Schedule.

CYPROHEPTADINE—

(a) in oral solid preparations containing 30 dosage units or less, except when included in the Second Schedule;

(b) in oral liquid preparations, except when included in the Second Schedule.

DEPTROPINE—

(a) in oral solid preparations containing 30 dosage units or less, except when included in the Second Schedule;

(b) in oral liquid preparations, except when included in the Second Schedule.

DEXBROMPHENIRAMINE—

(a) in oral solid preparations containing 30 dosage units or less, except when included in the Second Schedule;

(b) in oral liquid preparations, except when included in the Second Schedule.

DEXCHLORPHENIRAMINE—

(a) in oral solid preparations containing 30 dosage units or less, except when included in the Second Schedule;

(b) in oral liquid preparations, except when included in the Second Schedule.

5,5 DIBROMO-O-CRESOLSULFONPHTHALEIN in solutions for testing for pregnancy.

DIHYDROCODEINE when compounded with one or more therapeutically active medicaments in substances containing 1 per cent or less of dihydrocodeine.

DIMENHYDRINATE—

(a) in oral solid preparations containing 30 dosage units or less, except when included in the Second Schedule;

(b) in oral liquid preparations, except when included in the Second Schedule.

DIMETHINDENE—

(a) in oral solid preparations containing 30 dosage units or less, except when included in the Second Schedule;

(b) in oral liquid preparations, except when included in the Second Schedule.

DIMETHOTHIAZINE—

(a) in oral solid preparations containing 30 dosage units or less, except when included in the Second Schedule;

(b) in oral liquid preparations, except when included in the Second Schedule.

DIPHENHYDRAMINE—

(a) in oral solid preparations containing 30 dosage units or less, except when included in the Second Schedule;

(b) in oral liquid preparations, except when included in the Second Schedule.

DIPHENYLPYRALINE—

(a) in oral solid preparations containing 30 dosage units or less, except when included in the Second Schedule;

(b) in oral liquid preparations, except when included in the Second Schedule.

DITHRANOL for human therapeutic use.

DOXYLAMINE—

(a) in oral solid preparations containing 30 dosage units or less, except when included in the Second Schedule;

(b) in oral liquid preparations, except when included in the Second Schedule.

ECONAZOLE for cutaneous use in preparations containing 1 per cent or less of econazole.

EMBRAMINE—

(a) in oral solid preparations containing 30 dosage units or less, except when included in the Second Schedule;

(b) in oral liquid preparations, except when included in the Second Schedule.

EPHEDRINE in uncompounded divided preparations for internal use.

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

FLAVOXATE.

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.

HALOPYRAMINE—

(a) in oral solid preparations containing 30 dosage units or less, except when included in the Second Schedule;

(b) in oral liquid preparations, except when included in the Second Schedule.

HISTAPYRRODINE—
(a) in oral solid preparations containing 30 dosage units or less, except when included in the Second Schedule;
(b) in oral liquid preparations, except when included in the Second Schedule.

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

INSULIN and preparations containing the specific hypoglycaemic principle of the pancreas.

MEFENAMIC ACID in packs of 30 capsules or less when labelled for treatment of spasmodic dysmenorrhea.

MEPYRAMINE—
(a) in oral solid preparations containing 30 dosage units or less, except when included in the Second Schedule;
(b) in oral liquid preparations, except when included in the Second Schedule.

METHDILAZINE—
(a) in oral solid preparations containing 30 dosage units or less, except when included in the Second Schedule;
(b) in oral liquid preparations, except when included in the Second Schedule.

MICONAZOLE, in preparations containing 2 per cent or less for skin applications in humans.

NAPROXEN, in packs of 12 individual dosage units, tablets or capsules, for treatment of spasmodic dysmenorrhea.

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

NOSCAPINE.

OCTYL NITRITE.

PHENINDAMINE—
(a) in oral solid preparations containing 30 dosage units or less, except when included in the Second Schedule;
(b) in oral liquid preparations, except when included in the Second Schedule.

PHENIRAMINE—
(a) in oral solid preparations containing 30 dosage units or less, except when included in the Second Schedule;
(b) in oral liquid preparations, except when included in the Second Schedule.

PHENYLTOLOXAMINE—
(a) in oral solid preparations containing 30 dosage units or less, except when included in the Second Schedule;
(b) in oral liquid preparations, except when included in the Second Schedule.

PODOPHYLLIN for therapeutic internal use.

PREGNANCY TESTING KITS, and preparations and solutions for testing for pregnancy containing Human Chorionic Gonadotrophin bound to red blood cells and Human Chorionic Gonadotrophin antiserum, packed and labelled for use on one occasion only, and sold under the brand names "PREDICTOR", "EPT", and "DISCOVER 2".

PROMETHAZINE—

(a) in oral solid preparations containing 30 dosage units or less, except when included in the Second Schedule;

(b) in oral liquid preparations, except when included in the Second Schedule.

PYRROBUTAMINE—

(a) in oral solid preparations containing 30 dosage units or less, except when included in the Second Schedule;

(b) in oral liquid preparations, except when included in the Second Schedule.

QUININE for human therapeutic use.

SALBUTAMOL—

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

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

SANTONIN.

SODIUM CROMOGLYCATE in nasal preparations, topically applied.

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

THENALIDINE—

(a) in oral solid preparations containing 30 dosage units or less, except when included in the Second Schedule;

(b) in oral liquid preparations, except when included in the Second Schedule.

THENYLDIAMINE—

(a) in oral solid preparations containing 30 dosage units or less, except when included in the Second Schedule;

(b) in oral liquid preparations, except when included in the Second Schedule.

THEOPHYLLINE and derivatives therefrom in oral liquid preparations.

TOLPROPAMINE—

(a) in oral solid preparations containing 30 dosage units or less, except when included in the Second Schedule;

(b) in oral liquid preparations, except when included in the Second Schedule.

TRETINOIN for external human therapeutic use.

TRIMEPRAZINE—

(a) in oral solid preparations containing 30 dosage units or less, except when included in the Second Schedule;

(b) in oral liquid preparations, except when included in the Second Schedule.

TRIMETHOBENZAMIDE—

(a) in oral solid preparations containing 30 dosage units or less, except when included in the Second Schedule;

(b) in oral liquid preparations, except when included in the Second Schedule.

TRIPELENNAMINE—

(a) in oral solid preparations containing 30 dosage units or less, except when included in the Second Schedule;

(b) in oral liquid preparations, except when included in the Second Schedule.

TRIPROLIDINE—

(a) in oral solid preparations containing 30 dosage units or less, except when included in the Second Schedule;

(b) in oral liquid preparations, except when included in the Second Schedule.

Fourth Schedule

ACEDAPSONE.

ACETANILIDE and alkyl acetanilides, for human therapeutic use.

ACETAZOLAMIDE.

ACETOHEXAMIDE.

ACETYLCHOLINE and other choline esters.

ACETYLCYSTEINE.

ACETYLDIHYDROCODEINE when compounded with one or more other medicaments—

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

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

except when included in the Second Schedule.

ACETYLMETHYLDIMETHYLOXIMIDOPHENYLHYDRAZINE.

ADIPHENINE.

ADRENALINE, natural or synthetic, except in preparations containing 1 per cent or less of adrenaline.

ALCURONIUM.

ALPHADOLONE.

ALPHA-RECEPTOR BLOCKING AGENTS including phentolamine and phenoxybenzamine.

ALPHAXALONE.

ALPRAZOLAM.

AMANTADINE.

AMBENONIUM.

AMBUCETAMIDE.

AMBUTONIUM

AMETHOCAINE.

AMIKACIN.

AMILORIDE.

AMINOMETRADINE.

AMINOPHENAZONE and derivatives.

AMINOPTERIN.

AMINOREX.

AMIODARONE.

AMIPHENAZOLE.

AMISOMETRADINE.

AMITRIPTYLINE and other compounds not elsewhere specified in these schedules structurally derived therefrom by substitution in the side chain.

AMODIAQUINE.
AMOXYCILLIN.
AMPHOMYCIN.
AMPHOTERICIN.
AMPICILLIN.
AMYGDALIN (Laetrile)
AMYLOCAINE.
ANABOLIC steroidal agents.
ANAESTHETICS LOCAL, being synthetic cocaine substitutes, except when included in the Second Schedule.
ANGIOTENSINAMIDE.
ANTAZOLINE, except when included in the Second or Third Schedule.
ANTIBIOTICS not elsewhere specified, except—
(a) AVOPARCIN when intended for use as an animal feed additive;
(b) NISIN.
ANTIFOLIC ACID substances not elsewhere specified in these Schedules.
ANTIHISTAMINES not elsewhere specified in these Schedules except when included in the Second or Third Schedule.
ANTIMALARIAL SUBSTANCES not elsewhere specified.
ANTIMONY, organic compounds of, for therapeutic use.
ANTITUBERCULAR SUBSTANCES not elsewhere in these Schedules including isoniazid and its derivatives, para aminosalicylic acid and thiacetazone.
APOMORPHINE.
APROTININ.
ASPIRIN when combined with caffeine, paracetamol or salicylamide or any derivative of these substances.
ATENOLOL.
ATROPINE METHONITRATE.
AZAPERONE.
AZAPETINE.
AZATADINE.
AZLOCILLIN.

BACITRACIN except—
(a) when included in the Sixth Schedule;
(b) in animal feedstuffs for growth promotion in concentration of 50 mg/kg or less of the total active antibiotic principle;
(c) in milk replacers for calves and starter rations for pigs in concentrations of 100 mg/kg or less of the total active antibiotic principle.
BACLOFEN.
BAMIPINE, except when included in the Second or Third Schedule.
BARBITURIC ACID and its derivatives.
BECLAMIDE.
BEMEGRIDE.
BENACTYZINE and other substances structurally derived from diphenylmethane with ataractic properties when used for therapeutic purposes.
BENORYLATE.
BENSERAZIDE.
BENZAMINE, except when included in the Second Schedule.
BENZHEXOL.
BENZILONIUM.
BENZOCAINE, except when included in the Second Schedule.
BENZOYL PEROXIDE in preparations for external human therapeutic use, except when included in the Second or Third Schedule.

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) when in the the Second or Eighth Schedules;

(b) ephedrine, pseudoephedrine and phenylephrine in preparations exempted from the Second Schedule.

BENZTROPINE.

BENZYDAMINE.

BENZYLPENICILLIN (including procaine pencillin), except when included in the Sixth Schedule.

BETAHISTINE.

BETA-RECEPTOR BLOCKING AGENTS including alprenolol, propranolol and practolol.

BETANIDINE.

BIPERIDEN.

BISMUTH compounds of, for human therapeutic or cosmetic use, except bismuth citrate when incorporated in hair colourant preparations in concentrations of 0.5 per cent w/w or less and except when included in the Fifth Schedule.

BLEOMYCIN.

BORON COMPOUNDS for human therapeutic or cosmetic use except—

(a) in dusting powders or cosmetics containing 1 per cent or less of boron;

(b) in unit dose preparations for periodontal disease containing 100 mg or less of boron.

BRETYLIUM.

BROMIDES, inorganic, for therapeutic use.

BROMOCRIPTINE.

BROMODIPHENHYDRAMINE, except when included in the Second or Third Schedule.

BROMOFORM for therapeutic use.

BROMPHENIRAMINE except when included in the Second or Third Schedule.

BROMVALETONE.

BUCLIZINE, except when included in the Second or Third Schedule.

BUFEXAMAC, except when included in the Second Schedule.

BUMETANIDE.

BUPIVACAINE.

BUPRENORPHINE.

BUSULPHAN.

BUTACAINE.

BUTYLAMINOBENZOATE, except when included in the Second Schedule.

BUTYLCHLORAL HYDRATE.

CALCITONIN.

CALCITRIOL.

CALCIUM CARBIMIDE for therapeutic use.

CAMPHORATED OIL.

CAMPHOTAMIDE.

CANDICIDIN.

CANTHARIDIN, except when included in the Second Schedule.

CAPREOMYCIN.

CAPTODIAME.

CAPTOPRIL.

CAPURIDE.

CARAMIPHEN.

CARBACHOL.
CARBAMAZEPINE.
CARBARYL for human therapeutic use, except when included in the Second Schedule.
CARBAZOCHROME.
CARBENICILLIN.
CARBENOXOLONE, except when included in the Second Schedule.
CARBIDOPA.
CARBIMAZOLE.
CARBINOXAMINE, except when included in the Second or Third Schedule.
CARBOCROMEN.
CARBROMAL.
CARDIAC GLYCOSIDES not elsewhere specified in these Schedules.
CARINDACILLIN.
CEFACLOR.
CEFOTAXIME.
CEFOXITIN.
CEPHACETRILE.
CEPHALEXIN.
CEPHALORIDINE.
CEPHALOTHIN.
CEPHAMANDOLE.
CEPHAPIRIN.
CEPHAZOLIN.
CEPHRADINE.
CHENODEOXYCHOLIC ACID.
CHLORAL FORMAMIDE.
CHLORAL HYDRATE, except—

(a) when included in the Third Schedule;
(b) in preparations for topical use containing 2 per cent or less of chloral hydrate.

CHLORAMPHENICOL—

(a) for human therapeutic use;
(b) in eye preparations for animal use;
(c) in other preparations for animal use, except as prohibited in the Seventh Schedule.

CHLORAZANIL.
CHLORBUTOL in preparations for human oral use, except in preparations containing 0.5 per cent or less of chlorbutol as a preservative.
CHLORCYCLIZINE.
CHLORDIAZEPOXIDE and other substances structurally derived from benzodiazepine with ataractic properties when used for therapeutic purposes.
CHLORMERODRIN.
CHLORMETHIAZOLE.
CHLORMEZANONE.
CHLOROFORM when specifically prepared and packed as a therapeutic agent for the induction of inhalation anaesthesia.
1-(4-CHLOROPHENOXY)-1-IMIDAZOL-1-YL-3,3-DIMETHYL-2-BUTANONE for human use.
CHLOROPYRILENE except when included in the Second or Third Schedule.
CHLOROQUINE.
CHLOROTHIAZIDE and other substances structurally derived from benzothiadiazine for therapeutic use.
CHLORPHENIRAMINE, except when included in the Second or Third Schedule.

CHLORPHENOXAMINE, except when included in the Second or Third Schedule.
CHLORPHENTERMINE.
CHLORPROMAZINE and other substances structually derived from phenothiazine with ataractic properties when used for therapeutic purposes.
CHLORPROPAMIDE.
CHLORPROTHIXENE.
CHLORTETRACYCLINE, except when included in the Sixth Schedule.
CHLORTHALIDONE.
CHLORZOXAZONE.
CHOLESTYRAMINE for human therapeutic use.
CHYMOPAPAIN by injection for human therapeutic use.
CIMETIDINE.
CINCHOCAINE.
CINNARIZINE, except when included in the Second or Third Schedule.
CINOXACIN.
CISPLATIN.
CLANOBUTIN by injection for the treatment of animals.
CLEMASTINE, except when included in the Second or Third Schedule.
CLEMIZOLE, except when included in the Second or Third Schedule.
CLIDINIUM.
CLINDAMYCIN.
CLOBAZAM.
CLOBETASONE-17-BUTYRATE.
CLOFENAMIDE.
CLOFIBRATE.
CLOMIPHENE and other products specifically prepared to stimulate ovulation.
CLOMIPRAMINE.
CLOMOCYCLINE.
CLONAZEPAM.
CLONIDINE.
CLOPAMIDE.
CLOPROSTENOL for treatment of animals.
CLORAZEPATE.
CLOREXOLONE.
CLOTRIMAZOLE.
CLOXACILLIN.
CLOZAPINE.
CODEINE when compounded with one or more other medicaments—

(a) in divided preparations containing not more than 30 mg of codeine per dosage unit; or
(b) in undivided preparations with a concentration of not more than one per cent of codeine,

except when included in the Second Schedule.
COLASPASE.
COLCHICINE.
COLESTIPOL for human therapeutic use.
COLISTIN.
CORTISONE and steroid suprarenal cortical hormones, either natural or synthetic.
COUMARIN derivatives and phenylindanedione derivatives for therapeutic use.
CURARE, TUBOCURARINE, d-TUBOCURARINE, d-TUBOCURARINEDIMETHYLETHER, and all synthetic quaternary ammonium compounds, and other compounds having curarising properties.

CYCLANDELATE.
CYCLIRAMINE, except when included in the Second or Third Schedule.
CYCLIZINE
CYCLOFENIL.
CYCLOPENTOLATE.
CYCLOPROPANE when specifically prepared and packed as a therapeutic agent for the induction of inhalation anaesthesia.
CYCLOSERINE.
CYCRIMINE.
CYPROHEPTADINE, except when included in the Second or Third Schedule.

DACARBAZINE.
DANAZOL.
DANTROLENE.
DAPSONE and all derivatives of 4,4-diaminodiphenylsulphone.
DEANOL.
DEBRISOQUINE
DEMECARIUM BROMIDE.
DEMECLOCYCLINE.
DEPTROPINE, except when included in the Second or Third Schedule.
DESIPRAMINE.
DESMOPRESSIN (D.D.A.V.P.).
DEXBROMPHENIRAMINE, except when included in the Second or Third Schedule.
DEXCHLORPHENIRAMINE, except when included in the Second or Third Schedule.
DEXTROMETHORPHAN, except when included in the Second Schedule.
DEXTROPROPOXYPHENE.
DEXTRORPHAN, except in preparations containing 1 per cent or less of dextrorphan.
DIBENZEPIN.
DICHLORALPHENAZONE.
DICHLORPHENAMIDE.
DICLOFENAC.
DICYCLOMINE, except in preparations containing 0.1 per cent or less of dicyclomine.
DIETHAZINE.
DIETHYLCARDAMAZINE for human therapeutic use.
DIETHLPROPION.
DIFENOXIN in preparations containing, per dosage unit, 0.5 mg 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 100 mg 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 the Second Schedule.
DIHYDROSTREPTOMYCIN, except when included in the Sixth Schedule.
DIISOPROPYLAMINE DICHLOROACETATE.
DIMENTHYDRINATE, except when included in the Second or Third Schedule.

DIMETHINDENE, except when included in the Second or Third Schedule.
DIMETHISOQUIN, except when included in the Second Schedule.
DIMETHOTHIAZINE, except when included in the Second or Third Schedule.
DIMETHOXANATE.
DIMETHYL SULPHOXIDE for therapeutic use, except when included in the Sixth Schedule.
DINITROCRESOLS for therapeutic use.
DINITRONAPHTHOLS for therapeutic use.
DINITROPHENOLS for therapeutic use.
DINITROTHYMOLS for therapeutic use.
DINOPROST for treatment of animals.
DIPERODON.
DIPHEMANIL METHYLSULPHATE, except in preparations for topical use.
DIPHENYDRAMINE, except when included in the Second or Third Schedule.
DIPHENIDOL.
DIPHENOXYLATE in preparations containing per dosage unit 2.5 mg or less of diphenoxylate calculated as base, and a quantity of atropine sulphate equivalent to at least 1 per cent of the dose of diphenoxylate.
DIPHENYLPYRALINE, except when included in the Second or Third Schedule.
DIPIVEFRIN.
DIPYRIDAMOLE.
DISOPHENOL.
DISOPYRAMIDE.
DISULFIRAM, except when used for industrial purposes.
DITHIAZANINE, except in preparations containing 2 per cent or less of dithiazanine for the treatment of animals.
DITOPHAL.
DOBUTAMINE.
DOMPERIDONE.
DOPAMINE.
DOTHIEPIN.
DOXAPAM.
DOXEPIN.
DOXORUBICIN.
DOXYCYCLINE.
DOXYLAMINE, except when included in the Second or Third Schedule.
DROPERIDOL.
DROSTANOLONE.

ECONAZOLE, except when included in the Third or Sixth Schedule.
EDETIC ACID for human therapeutic use in preparations for injection or infusion.
EMBRAMINE, except when included in the Second or Third Schedule.
EMETINE, except in preparations containing 0.2 per cent or less of emetine.
ENFLURANE when specifically prepared and packed as a therapeutic agent for the induction of inhalation anaesthesia.
EPICILLIN.
ERGOT.
ERYTHROMYCIN except—

(a) when included in the Sixth Schedule;
(b) in animal feedstuffs for growth promotion in concentrations of 50 mg/kg or less of the total active antibiotic principle;
(c) in milk replacers for calves or starter rations for pigs in concentrations of 100 mg/kg or less of the total active antibiotic principle.

ETHACRYNIC ACID.
ETHAMBUTOL.
ETHAMIVAN.
ETHCHLORVYNOL.
ETHER when specifically prepared and packed as a therapeutic agent for the induction of inhalation anaesthesia.
ETHINAMATE.
ETHOGLUCID.
ETHOHEPTAZINE, except in preparations containing 1 per cent or less of ethoheptazine.
ETHOPROPAZINE.
ETHOXZOLAMIDE.
ETHYL CHLORIDE when specifically prepared and packed as a therapeutic agent for the induction of inhalation anaesthesia.
ETHYLENE when specifically prepared and packed as a therapeutic agent for the induction of inhalation anaesthesia.
ETHYLMORPHINE when compounded with one or more other medicaments—
(a) in divided preparations containing not more than 100 mg 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 the Second Schedule.
ETHYLOESTRENOL.
ETIDOCAINE.
ETIDRONATE.
ETRETINATE.
ETOPOSIDE.

FENCAMFAMIN.
FENFLURAMINE.
FENOPROFEN.
FENOTEROL, except when included in the Third Schedule.
FENPRIPRAMIDE.
FENPIPRANE.
FENPROSTALENE for the treatment of animals.
FLAVOPHOSPHOLIPOL except:
(a) when included in the Sixth Schedule;
(b) in animal feedstuffs for growth promotion in concentrations of 50 mg/kg or less of the total active antibiotic principle.
FLUCLOXACILLIN.
FLUCYTOSINE.
FLUFENAMIC ACID.
FLUNISOLIDE.
FLUNITRAZEPAM.
FLUNIXIN MEGLUMINE, for animal use.
FLUOROURACIL and other substances structurally derived from uracil with cytotoxic properties when used for therapeutic purposes.
FLUOXYMESTERONE.
FLUPROSTANOL for treatment of animals.
FLURAZEPAM.
FLUROXENE when specifically prepared and packed as a therapeutic agent for the induction of inhalation anaesthesia.
FLUSPIRILENE.

FRAMYCETIN.
FRUSEMIDE.
FUSIDIC ACID.

GALANTHAMINE.
GALLAMINE.
GENTAMYCIN.
GLIBENCLAMIDE
GLIBORNURIDE
GLICLAZIDE.
GLUCAGON.
GLUTETHIMIDE.
GLYCERYL TRINITRATE in preparations for injection.
GLYCOPYRROLATE.
GLYMIDINE.
GRAMICIDIN.
GRISEOFULVIN.
GUANACLINE.
GUANETHIDINE.

HALCINONIDE.
HALOPERIDOL and other substances structurally derived from butyrophenone with ataractic properties when used for therapeutic purposes.
HALOPYRAMINE, except when included in the Second or Third Schedule.
HALOTHANE when specifically prepared and packed as a therapeutic agent for the induction of inhalation anaesthesia.
HEPARIN.
HETACILLIN.
HEXACHLOROPHANE in preparations for use on infants and in all other preparations, except when included in the Second or Sixth Schedule.
HEXAMETHONIUM.
HEXOCYCLIUM.
HISTAPYRRODINE, except when included in the Second or Third Schedule.
HUMAN CHORIONIC GONADOTROPHIN except when included in the Third Schedule.
HYDRALAZINE.
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 included in the Sixth Schedule;
(b) in preparations in concentrations of 50mg/kg or less of hygromycin.
HYOSCINE BUTYLBROMIDE.
HYPOTHALMIC RELEASING FACTORS when used for diagnostic purposes.

IBUFENAC.
IBUPROFEN.
IDOXURIDINE, except when included in the Third Schedule.

IMIPRAMINE.
INDAPAMIDE.
INDOMETHACIN.
INOSITOL NICOTINATE for internal use.
ION-EXCHANGE RESINS, anionic and cationic, for internal use in humans.
IPRATROPIUM.
IRON compounds, injectable preparations for human therapeutic use.
ISOAMINILE.
ISOCONAZOLE.
ISOETHARINE.
ISOMETHEPTENE.
ISOPRENALINE.
ISOPROPAMIDE, except when included in the Second Schedule.
ISOXUPRINE.

KANAMYCIN.
KETAMINE.
KETOPROFEN.
KHELLIN.
KITASAMYCIN, except—

(a) when included in the Sixth Schedule;
(b) in animal feedstuffs for growth promotion in concentrations of 100mg/kg or less of the total active antibiotic principle.

LABETALOL.
LAUDEXIUM METHYLSULPHATE.
LEFETAMINE.
LEPTAZOL.
LEVAMISOLE—

(a) for human therapeutic use;
(b) in preparations for the prevention or treatment of heartworm in dogs.

LEVODOPA.
LIDOFLAZINE.
LIGNOCAINE, except when included in the Second Schedule.
LINCOMYCIN.
LITHIUM salts for therapeutic use, except in preparations containing 0.01 per cent or less of lithium.
LOPERAMIDE.
LORAZEPAM.
LOXAPINE.
LYMECYCLINE.

MAFENIDE.
MAPROTILINE.
MAZINDOL.
MEBEVERINE.
MEBHYDROLIN.
MECAMYLAMINE.
MECLOFENOXATE.
MECLOZINE.
MEDAZEPAM.
MEFENAMIC ACID, except when included in the Third Schedule.

MEFLOQUINE.
MEFRUSIDE.
MEPACRINE.
MEPENZOLATE.
MEPHENESIN and its derivatives except guaiphenesin when included in the Second Schedule.
MEPHENTERMINE.
MEPIVACAINE.
MEPROBAMATE.
MEPYRAMINE, except when included in the Second or Third Schedule.
MERCAPTOPURINE and other substances structurally derived therefrom with cytotoxic properties when used for therapeutic purposes.
MERCUROUS CHLORIDE for therapeutic use.
MERCURY organic compounds of, for therapeutic use, except preparations for topical use containing 0.5 per cent or less of mercury.
METARAMINOL.
METFORMIN.
METHACYCLINE.
METHANDIENONE.
METHANDRIOL.
METHANTHELINIUM.
METHAZOLAMIDE.
METHDILAZINE, except when included in the Second or Third Schedule.
METHENOLONE.
METHICILLIN.
METHIMAZOLE.
METHIXENE.
METHOCARBAMOL.
METHOTREXATE.
METHOXSALEN.
METHOXYFLURANE when specifically prepared and packed as a therapeutic agent for the induction of inhalation anaesthesia.
METHYLANDROSTANOLONE.
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 the Third Schedule.
MINOCYCLINE.
MINOXIDIL.
MITHRAMYCIN.
MITOBRONITOL.
MITOMYCIN.

MONENSIN, except—

(a) in animal feedstuffs containing 33 mg/kg or less of monensin;

(b) when included in the Sixth Schedule.

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

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

MOPERONE.

MORPHINE ANTAGONISTS including nalorphine, naloxone and levallorphan.

MOXALACTAM.

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

NALBUPHINE.

NALIDIXIC ACID.

NANDROLONE.

NAPROXEN, except when included in the Third Schedule.

NARASIN except—

(a) when included in the Sixth Schedule;

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

NATAMYCIN, except where used as a permitted food additive.

NEOMYCIN, except when included in the Sixth Schedule.

NEOSTIGMINE.

NETILMICIN.

NICOCODINE when compounded with one or more other medicaments—

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

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

except when included in the Second Schedule.

NICODICODINE when compounded with one or more other medicaments—

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

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

except when included in the Second Schedule.

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

NICOTINIC ACID, where the recommended daily dose exceeds 250 mg.

NICOTINYL ALCOHOL for internal use.

NIFEDIPINE.

NIFENAZONE.

NIKETHAMIDE.

NIRIDAZOLE.

NITRAZEPAM.

NITROFURAN and its derivatives for human therapeutic use, except when included in the Third Schedule.

NITROUS OXIDE when specifically prepared and packed as a therapeutic agent for the induction of inhalation anaesthesia.

NOMIFENSINE.

NORADRENALINE (excluding its derivatives).

NORCODEINE when compounded with one or more other medicaments—

(a) in divided preparations containing not more than 100 mg 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 the Second Schedule.

NORETHANDROLONE.

NORTRIPTYLINE.

NOVOBIOCIN, except when included in the Sixth Schedule.

OCTAMYLAMINE.

OCTATROPINE.

OLEANDOMYCIN except—

(a) when included in the Sixth Schedule;

(b) in animal feedstuffs for growth promotion in concentrations of 50 mg/kg or less of the total active antibiotic principle.

OPIPRAMOL.

ORCIPRENALINE.

ORGANOPHOSPHORUS COMPOUNDS with anticholinesterase activity for human therapeutic use, except when included in the Second Schedule.

ORNIDAZOLE.

ORNIPRESSIN.

ORPHENADRINE.

ORTHOCAINE.

ORTHOPTERIN.

OXACILLIN.

OXANDROLONE.

OXAZEPAM.

OXPRENOLOL.

OXYBUPROCAINE.

OXYMESTERONE.

OXYMETHOLONE.

OXYPHENBUTAZONE.

OXYPHENCYCLIMINE.

OXYPHENONIUM.

OXYTETRACYCLINE, except when included in the Sixth Schedule.

PAMAQUINE.

PANCURONIUM.

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

PARALDEHYDE.

PARAMETHADIONE.

PAROMOMYCIN.

PEMOLINE.

PEMPIDINE.

D-PENICILLAMINE.

PENTAMETHONIUM.

PENTHIENATE.

PENTOLINIUM.

PERHEXILENE.

PHENACEMIDE and other substances structurally derived from acetylurea with anticonvulsant properties when used for therapeutic purposes.

PHENACETIN.
PHENAZONE for internal use.
PHENAZOPYRIDINE.
PHENETHICILLIN, except when included in the Sixth Schedule.
PHENFORMIN.
PHENGLUTARIMIDE.
PHENINDAMINE, except when included in the Second or Third Schedule.
PHENIRAMINE, except when included in the Second or Third Schedule.
PHENOXYBENZAMINE.
PHENOXYMETHYLPENICILLIN, except when included in the Sixth Schedule.
PHENSUXIMIDE and other substances structurally derived from succinamide with anticonvulsant properties when used for therapeutic purposes.
PHENTERMINE.
PHENTHIMENTONIUM.
PHENYAPIN.
PHENYLBUTAZONE.
PHENYLPROPANOLAMINE, except when included in the Second Schedule.
PHENYLTOLOXAMINE, except when included in the Second or Third Schedule.
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 100 mg 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 the Second Schedule.
PHYSOSTIGMINE.
PICROTOXIN.
PILOCARPINE, except in preparations containing 0.025 per cent or less of pilocarpine.
PIMOZIDE.
PINDOLOL.
PIPENZOLATE.
PIPERACILLIN.
PIPERIDOLATE.
PIPOBROMAN.
PIPRADROL.
PIROXICAM.
PITUITARY, its extracts, its active principles or their synthetic substitutes, except when included in the Seventh Schedule.
PIZOTIFEN.
POLYMETHYLENE BISTRIMETHYL AMMONIUM COMPOUNDS.
POLYMYXIN.
POTASSIUM PERCHLORATE for therapeutic use.
PRAMOXINE, except when included in the Second Schedule.
PRAZEPAM.
PREGNENOLONE ACETATE, except in preparations for topical use.
PRENYLAMINE.
PRILOCAINE.
PRIMAQUINE.
PRIMIDONE.
PROBENECID.
PROCAINAMIDE.

PROCAINE.
PROCARBAZINE.
PROCHLORPERAZINE.
PROCYCLIDINE, except when included in the Second Schedule.
PROGUANIL.
PROLINTANE.
PROMETHAZINE, except when included in the Second or Third Schedule.
PROPANIDID.
PROPANTHELINE, except in preparations for topical use.
PROPYLHEXEDRINE, except when included in the Second Schedule.
PROQUAZONE.
PROSTAGLANDINS, except where separately specified in this Schedule.
PROSTIANOL, for treatment of animals.
PROTHIONAMIDE.
PROTRIPTYLINE.
PROXYMETACAINE.
PYRIDOSTIGMINE.
PYRIMETHAMINE.
PYRROBUTAMINE, except when included in the Second or Third Schedule.

QUINETHAZONE.
QUINIDINE.

RANITIDINE.
RAUWOLFIA SERPENTINA.
RIFAMPICIN.
RITODRINE.
ROLITERACYCLINE.
ROSOXACIN.

SALBUTAMOL, except when included in the Third Schedule.
SALICYLAMIDE when combined with aspirin, caffeine or paracetamol or any derivative of these substances.
SALINOMYCIN, except—
(a) in animal feedstuffs containing 60 mg/kg or less of the total active principal;
(b) when included in the Sixth Schedule.
SELENIUM except—
(a) when included in the Fifth or Sixth Schedule;
(b) when included in animal feedstuffs containing 0.1 g/tonne or less of selenium in total feed;
(c) in compressed pellets for control of selenium responsive conditions in sheep.
SEX HORMONES, natural or synthetic, their substitutes in all preparations, including cosmetics; except—
(a) their derivatives and their substitutes without sex hormonal activity;
(b) when specifically named in this or any other schedule.
SILVER SULPHADIAZINE.
SISOMYCIN.
SODIUM CROMOGLYCATE, except when included in the Third Schedule.
SODIUM FLUORIDE, in preparations for human ingestion, except when included in the Second Schedule.

SODIUM NITROPRUSSIDE for human therapeutic use.
SODIUM VALPROATE.
SONTOQUINE.
SPARTEINE.
SPECTINOMYCIN.
SPIRAMYCIN, except—
(a) when included in the Sixth Schedule;
(b) in animal feedstuffs for growth promotion in pigs or poultry in concentrations of 50 mg/kg or less of the total active antibiotic principle.
SPIRONOLACTONE.
STANOLONE.
STANOZOLOL.
STREPTOMYCIN, except when included in the Sixth Schedule.
STROPHANTUS 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 included in the Sixth Schedule;
(b) sulphaquinoxaline when incorporated in baits for the destruction of vermin and in animal feedstuffs containing 200 mg/kg or less of sulphaquinoxaline;
(c) Oryzalin;
(d) when specifically named in this or any other schedule.
SULPHATROXAZOLE, for animal use.
SULPHINPYRAZONE.
SULPHOMYXIN.
SULPHONAL and alkyl sulphonals.
SULTHIAME.
SUXAMETHONIUM.

TACRINE.
TAMOXIFEN.
TEMAZEPAM.
TENIPOSIDE.
TERBUTALINE, except when included in the Third Schedule.
TEROPTERIN.
TETRABENAZINE.
TETRACYCLINE, except when included in the Sixth Schedule.
THENALIDINE, except when included in the Second or Third Schedule.
THENYLDIAMINE, except when included in the Second or Third Schedule.
THEOPHYLLINE and derivatives therefrom, except when included in the Third Schedule.
THIACETARSAMIDE, in preparations for the prevention or treatment of heart worm in dogs.
THIAMBUTOSINE.
THIAZOSULPHONE.
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 its extract, and its active principles.
TIAMULIN except—
(a) when included in the Sixth Schedule;
(b) in prepared animal feedstuffs.
TICARCILLIN.
TIEMONIUM.
TIGLOIDINE.
TIMOLOL.
TINIDAZOLE.
TIPEPIDINE.
TOBRAMYCIN.
TOCAINIDE.
TOLAZAMIDE.
TOLAZOLINE for internal use.
TOLBUTAMIDE.
TOLPROPAMINE, except when included in the Second or Third Schedule.
TRANEXAMIC ACID.
TRETAMINE.
TRIAMTERENE.
TRIAZIQUONE.
TRIAZOLAM.
TRICHLOROETHYLENE when specifically prepared and packed as a therapeutic agent for the induction of inhalation anaesthesia.
TRICLOFOS.
TRICYCLAMOL.
TRIDIHEXETHYL.
TRIFLUPERIDOL.
TRIMEPRAZINE, except when included in the Second or Third Schedule.
TRIMETAPHAN.
TRIMETHOBENZAMIDE, except when included in the Second or Third Schedule.
TRIMETHOPRIM.
TRIMIPRAMINE and other compounds structurally derived therefrom by substitution in the side chain.
TRIMUSTINE.
TRIOXYSALEN.
TRIPELENNAMINE, except when included in the Second or Third Schedule.
TRIPROLIDINE, except when included in the Second or Third Schedule.
TROXIDONE and other substances structurally derived from oxazolidinone with anticonvulsant properties when used for therapeutic purposes.
TYLOSIN except-
(a) when included in the Sixth Schedule;
(b) in animal feedstuffs for growth promotion in concentrations of 50 mg/kg or less of the total active antibiotic principle;
(c) in milk replacers for calves or starter rations for pigs in concentrations of 100 mg/kg or less of the total active antibiotic principle.

URETHANE (excluding its derivatives), for therapeutic use.
URETHANES AND UREIDES having or purporting to have soporific hypnotic or narcotic properties not specifically included in this or any other schedule.

VACCINES, sera, toxoids, and antigens for human parenteral use.
VACCINES, veterinary live virus.

VALNOCTAMIDE.
VERAPAMIL.
VERATRUM for therapeutic use.
VIDARABINE,
VINCA ALKALOIDS, including semi-synthetic derivatives.
VINYL ETHER when specifically prepared and packed as a therapeutic agent for the induction of inhalation anaesthesia.
VIPRYNIUM
VIRGINIAMYCIN except—
(a) when included in the Sixth Schedule;
(b) in animal feedstuffs for growth promotion in concentrations of 50 mg/kg or less of the total active antibiotic principle.
VISNADINE.
VITAMIN A in preparations containing more than 10 000 IU per recommended daily dosage for human use.
VITAMIN D in preparations containing more than 25 micrograms per recommended daily dosage for human use.

XANTHINE OXIDASE INHIBITORS including allopurinol.
XANTHINOL NICOTINATE.
XYLAZINE.

YOHIMBINE.

ZOMEPIRAC.

Excluding however, the substances hereinbefore mentioned when contained in the products listed as exemptions, set out after the Eighth Schedule in this Appendix.

Fifth Schedule.

ACETIC ACID (excluding its salts and derivatives) in preparations containing more than 30 per cent acetic acid except when—
(a) included in the Second or Sixth Schedule;
(b) for therapeutic use.
ACETIC ANHYDRIDE (excluding its salts and derivatives) in preparations containing more than 30 per cent acetic anhydride, except when included in the Sixth Schedule or for therapeutic use.
ACETONE when packed in containers of 20 litres or less, except—
(a) in preparations containing 25 per cent or less of acetone;
(b) when packed in containers of 60 ml or less.
AKLOMIDE.
ALACHLOR.

ALKALINE SALTS, being the carbonate, orthosilicate, metasilicate or tribasic phosphate salts of sodium or potassium, and in any combination, except—

(a) in preparations containing 10 per cent or less of combined substances;

(b) in solid preparations the pH of which in 1 per cent (w/w) aqueous solution is 11.5 or less;

(c) in liquid preparations having a pH of 11.5 or less.

ALLOXYDIM.

AMETRYNE.

AMITROLE.

AMMONIA (excluding its salts and derivatives other than ammonium hydroxide) in preparations containing 5 per cent or less of free ammonia, except—

(a) in medicinal preparations for internal use;

(b) in appliances for inhalation in which the substance is absorbed upon an inert solid material;

(c) in preparations containing 0.5 per cent or less of free ammonia.

AMMONIUM THIOCYANATE.

ARSENIC organic compounds of, in preparations containing 3 per cent or less of arsenic, when prepared for use as herbicides or defoliants.

BARIUM SILICOFLUORIDE when coated on paper in an amount not exceeding 8 mg per sq. cm.

BENDIOCARB in preparations containing 2 per cent or less of bendiocarb.

BENTAZONE.

BENZOYL PEROXIDE, except when included in the Second, Third or Fourth Schedule or when used as an approved food additive.

S-BENZYL N,N-DI-(SEC-BUTYL)-THIOLOCARBAMATE.

BHC (excluding the gamma-isomer) in preparations containing 10 per cent or less of BHC.

BIOALLETHRIN, including sinbioallethrin, except in preparations containing 10 per cent or less of bioallethrin.

BIORESMETHRIN, except in preparations containing 10 per cent or less of bioresmethrin.

BISMUTH COMPOUNDS when used in a dusting powder for local application and containing less than 3 per cent of bismuth.

BORIC ACID and BORAX except—

(a) in preparations containing 1 per cent or less of boron;

(b) in hand cleaning preparations;

(c) when included in the Fourth Schedule.

BUTHIDAZOLE.

BUTOXYCARBOXIM in solid preparations containing 10 per cent or less.

2-iso-BUTYLAMINO-4-ETHYLAMINO-6-METHOXY-1,3,5-TRIAZINE.

CADMIUM SULPHIDE in preparations containing 2.5 per cent or less of cadmium sulphide for human therapeutic use.

CAMPHOR, except—

(a) in preparations containing 10 per cent or less of camphor;

(b) when included in the Fourth Schedule.

CARBARYL in preparations containing 10 per cent or less carbaryl, except when included in the Second or Fourth Schedule; or when impregnated in plastic resin strip material containing 20 per cent or less of carbaryl.

CHLORDECONE in preparations containing 5 per cent or less of chlordecone.

CHLORFENAC.

CHLORFENSON.

CHLORINATING COMPOUNDS AND BLEACHES containing more than 4 per cent of available chlorine, except—

(a) when included in the Seventh Schedule;

(b) when included elsewhere in this Schedule.

CHLORNIDINE.

CHLOROCRESOL, except in preparations containing 3 percent or less chlorocresol.

2-CHLORO-N-[(4-METHOXY-6-METHYL-1,3,5-TRIAZIN-2-YL)AMINOCARBONYL] BENZENE SULPHONAMIDE.

1-(4-CHLOROPHENOXY)-1-IMIDAZOL-1-YL-3,3-DIMETHYL-2-BUTANONE in concentrations of more than 2 per cent, except when included in the Fourth or Sixth Schedule.

CHLOROPROPYLATE.

CHLOROTHALONIL.

CLANOBUTIN for animal use, except when included in the Fourth Schedule.

COPPER SULPHATE, except for internal human therapeutic use.

4-CPA.

CUPRIMYXIN for the treatment of animals.

CYANATRYN.

CYANOACRYLIC ACID ESTERS.

CYANURIC ACID (excluding its salts and derivatives).

CYCLOHEXANONE PEROXIDE.

CYPERMETHRIN in preparations containing 10 per cent or less of cypermethrin.

2,4-D.

2-4-DB.

DDT in preparations containing 10 per cent or less of DDT, except for human therapeutic use.

2,4-DES.

N,N-DIALLYLDICHLOROACETAMIDE, except in preparations containing ten percent or less of N,N-diallyldichloroacetamide.

DICAMBA.

DICHLONE.

p-DICHLOROBENZENE.

DICHLOROISOCYANURATES and in preparations containing more than 4 per cent available chlorin

1-[2(2,4-DICHLOROPHENYL)-4-PROPYL-1,3-DIOXALAN-2-YL-METHYL]-IH-1,2,4-TRIAZOLE, in concentrations of 20 per cent or less.

1-[2-(2,4-DICHLOROPHENYL)-2-(2-PROPENYLOXY)ETHYL]-1H-IMIDAZOLE.

3,6-DICHLOROPICOLINIC ACID.

DICHLORVOS—

(a) when impregnated in plastic resin strip material containing 20 per cent or less dichlorvos;

(b) sustained release resin pellets for veterinary use containing 20 per cent or less dichlorvos;

(c) in aerosol packs containing 10 grams or less of dichlorvos.

DICLORAN.

DICOFOL.

DIMETHIRIMOL.

DIMETHYLFORMAMIDE in preparations containing 10 per cent or less of dimethylformamide.

DINITRAMINE.

DIPHENAMID.
DODINE.

EPOXY RESINS LIQUID and all amines and organic anhydrides used as curing agents for epoxy resins.
EPTC.
ETHEPHON (excluding its salts and derivatives).
ETHER PREPARATIONS for use in internal combustion engines.
ETHOFUMESATE.
ETHOXYQUIN, except in preparations containing 10 per cent or less of ethoxyquin.
ETHYLENE GLYCOL when packed and labelled as a boiling point or freezing point modifier and containing 10 mg/kg of denatonium benzoate as a bittering agent.
EUCALYPTUS OIL, except in preparations containing 25 per cent or less of eucalyptus oil.

FENARIMOL.
FENBUTATIN-OXIDE.
FENOPROP.
FENSON.
FENTHION in preparations containing 20 per cent or less of fenthion when packed in single-use containers having a capacity of 1.0 ml or less.
FLAMPROP-METHYL.
FLUCHLORALIN.
FORMIC ACID (excluding its salts and derivatives).
FOSPIRATE when impregnated in plastic resin strip material containing 20 per cent or less of fospirate.
FURALAXYL.

GLUTARALDEHYDE, in preparations containing 5 per cent or less of glutaraldehyde, except when included in the Second Schedule.
GLYPHOSATE.

HEXAZINONE.
HYDROCARBONS, LIQUID, including kerosene, mineral turpentine, white petroleum spirit, toluene, xylene and light mineral and paraffin oils (but excluding their derivatives) distilling under 300°C, except—

(a) toluene and xylene when included in the Sixth Schedule;
(b) in containers having a capacity of more than 20 litres;
(c) in solid or semi-solid cleaning and polishing preparations;
(d) in preparations containing 25 per cent or less of a total of such liquid hydrocarbons;
(e) in preparations packed in pressurised aerosol containers;
(f) in adhesives packed in containers each containing 50 grams or less of adhesive.

HYDROCHLORIC ACID (excluding its salts and derivatives) in preparations containing 10 per cent or less of hydrochloric acid (HCL) except—

(a) in preparations containing 0.5 per cent or less of hydrochloric acid (HCL);
(b) for thereapeutic use.

HYDROFLUORIC ACID and HYDROSILICOFLUORIC ACID in preparations containing 0.5 per cent or less of hydroflouric acid or hydrosilicoflouric acid except in substances containing 15 mg/kg or less of fluoride ion.

HYDROGEN PEROXIDE (excluding its salts and derivatives), except in preparations containing 6 per cent (20 vol) or less of hydrogen peroxide.

IODOFENPHOS.

ISOPROPYL-N-(3-N-ETHYL-N-PHENYLCARBAMOYLOXY) PHENYL-CARBAMATE.

KEROSENE, see HYDROCARBONS LIQUID.

LEAD COMPOUNDS, in preparations for use as hair cosmetics.

LEVAMISOLE in preparations containing 15 per cent or less of levamisole for the treatment of animals, except when included in the Fourth Schedule.

LINDANE in preparations containing 10 per cent or less of lindane, except when included in the Second Schedule.

MALDISON in preparations containing 10 per cent or less maldison except—

(a) when included in the Second Schedule;

(b) in other preparations containing 2 per cent or less maldison.

MANCOZEB.

MANEB.

MCPA.

MCPB.

MECOPROP.

MEPIQUAT.

METALDEHYDE in preparations containing 2 per cent or less of metaldehyde.

METHABENZTHIAZURON.

METHAZOLE.

METHIOCARB in pelleted preparations containing 2 per cent or less of methiocarb.

METHOXYCHLOR.

METHYLATED SPIRIT INDUSTRIAL, as defined by the Spirits Act 1906 of the Parliament of the Commonwealth or any Act in substitution for that Act, as amended from time to time, except in containers having a capacity of more than 5 litres, and except in preparations containing 75 per cent or less methylated spirits industrial.

METHYLENE CHLORIDE except when used in aerosols.

METHYL ETHYL KETONE when packed in containers of 20 litres or less, except in preparations containing 25 per cent or less of ketones included in the Fifth Schedule.

METHYL ETHYL KETONE PEROXIDE.

METHYL-ISO-AMYL KETONE when packed in containers of 20 litres or less, except in preparations containing 25 per cent or less of ketones included in the Fifth Schedule.

METHYL-ISO-BUTYL KETONE when packed in containers of 20 litres or less, except in preparations containing 25 per cent or less of ketones included in the Fifth Schedule.

METHYL N-(METHOXYACETYL)-N-(2,6-XYLYL) ALANINATE.

METIRAM.

METOLACHLOR.

METRIBUZIN.

MEZINEB.

MINERAL TURPENTINE, see HYDROCARBONS LIQUID.

NAA.

NALED when impregnated in plastic resin strip material containing 20 per cent or less of naled.

NAPHTHALENE as such.

NAPHTHALENE ACETIC ACID, except in preparations containing 25 per cent or less of naphthalene acetic acid.

NITRIC ACID (excluding its salts and derivatives) in preparations containing 10 per cent or less of nitric acid as such, except preparations containing 0.5 per cent or less of nitric acid.

NORBORMIDE.

OFURACE.

ORGANO-TIN COMPOUNDS not elsewhere included in this schedule in preparations containing 1 per cent or less of such compounds.

OXYCARBOXIN.

OXYTHIOQUINOX.

PARA-DICHLOROBENZENE.

PEBULATE.

PENDIMETHALIN.

PERACETIC ACID in concentrations of 10 per cent or less.

PETROL when packed in containers of 20 litres or less, except preparations containing 25 per cent or less of petrol.

ortho-PHENYLPHENOL, except in preparations containing 3 per cent or less of the phenylphenol.

PHOSPHONIC ACID, except in preparations containing 10 per cent or less phosphonic acid.

PHOSPHORIC ACID (excluding its salts and derivatives) except—

(a) when packed in containers with a capacity of not less than 10 litres and labelled with the word "CORROSIVE", in bold face *sans serif* capital letters of a height of not less than 1 cm;

(b) in preparations containing 350 g/litre or less of phosphoric acid;

(c) in solid and semi—solid preparations;

(d) in professional dental kits.

PIRIMICARB in preparations containing 0.5 per cent or less pirimicarb.

POLY (HEXAMETHYLENE BIGUANIDE), except in preparations containing 5 per cent or less of poly hexamethylene biguanide.

POTASSIUM HYDROXIDE (excluding its salts and derivatives) in preparations containing 5 per cent or less of potassium hydroxide, except—

(a) in preparations containing 0.5 per cent or less of potassium hydroxide;

(b) in accumulators and batteries.

POTASSIUM SULPHIDE in preparations for metal treatment in containers each containing 50 g or less of potassium sulphide.

PROMETRYN.

PROPANIL.

PROPIONIC ACID (excluding its salts and derivatives) in preparations containing more than 30 per cent propionic acid, except—

(a) when included in the Sixth Schedule;

(b) for therapeutic use.

PROPOXUR—

(a) in dust preparations containing 3 per cent or less of propoxur;

(b) in granular sugar-based fly baits containing 1 per cent or less of propoxur providing that the preparation also contains a dark colouring agent and separate bittering agent.

(c) in aerosol packs containing 10 g or less of propoxur.

PRYNACHLOR.

PYRETHRINS, naturally occurring, being pyrethrolone, cinerolone or jasomline esters of chrysanthemic or pyrethric acids, except in preparations containing 10 per cent or less of such substances.

PYRINURON in preparations containing 10 per cent or less of pyrinuron.

PYRITHIONE ZINC in preparations containing 2 per cent or less of pyrithione zinc.

QUATERNARY AMMONIUM COMPOUNDS and in preparations containing more than 10 per cent quaternary ammonium compounds, except when included in any other schedule.

QUINTOZENE.

SALICYLANILIDE.

SECBUMETON.

SELENIUM SULPHIDE in preparations containing 2.5 per cent or less of selenium sulphide for topical therapeutic use.

SETHOXYDIM.

SODIUM CHLORATE.

SODIUM HYDROGEN SULPHATE.

SODIUM HYDROXIDE (excluding its salts and derivatives) in preparations containing 5 per cent or less of sodium hydroxide, except in preparations containing 0.5 per cent or less of sodium hydroxide.

SODIUM NITRITE except—

(a) in preparations containing 1 per cent or less of sodium nitrite;

(b) for therapeutic use.

SODIUM SULPHIDE in preparations for metal treatment in containers each containing 50 g or less of sodium sulphide.

STYRENE (excluding its derivatives) when packed in containers of 20 litres or less.

SULPHAMIC ACID, except in preparations containing 10 per cent or less of sulphamic acid.

2,3,6-TBA.

TCA—see TRICHLOROACETIC ACID.

TDE in preparations containing 10 per cent or less TDE.

TERBUMETON.

TERBUTRYN.

TETRACHLOROETHYLENE in preparations containing 5 per cent or less tetrachloroethylene except—

(a) when prepared for use for the treatment of humans and for the treatment of animals;

(b) when packed in containers of 50 ml or less; or

(c) when absorbed into an inert solid material.

TETRACHLORVINPHOS, except in animal feedstuffs containing 0.2 per cent or less of tetrachlorvinphos.

TETRAMETHRIN, except in aerosol packs.

THIOBENCARB.

TRIADIMEFON, in wettable powders containing 25 per cent or less of triadimefon.

TRIADIMENOL.

TRI-ALLATE.

TRICHLOROACETIC ACID, alkali salts of.

1,1,1-TRICHLOROETHANE when packed in containers of 20 litres or less except—

(a) in preparations containing 25 per cent or less of 1,1,1-trichloroethane;

(b) when used in aerosols other than for therapeutic use;

(c) when packed in containers of 50 mls or less.

TRICHLOROISOCYANURIC ACID when compressed in block form for use in swimming pools.

TRIETAZINE.

TURPENTINE OIL when packed in containers of 20 litres or less except in preparations containing 25 per cent or less of turpentine oil.

VERNOLATE.

WARFARIN, in rodent baits containing 0.1 per cent or less of warfarin.

ZINEB.

ZIRAM.

Excluding however, the substances hereinbefore mentioned when contained in the products listed as exemptions, set out after the Eighth Schedule in this Appendix.

Sixth Schedule.

ACEPHATE.

ACETIC ACID (excluding its salts and derivatives) in preparations containing more than 80 per cent acetic acid, except when included in the second Schedule.

ACETIC ANHYDRIDE (excluding its salts and derivatives) in preparations containing more than 80 per cent of acetic anhydride, except—

(a) when included in the Fifth Schedule; or

(b) for therapeutic use.

ACIFLUORFEN.

ALDRIN.

ALLIDOCHLOR.

ALPHA-CHLOROHYDRIN.

AMIDITHION.

2-AMINO-BUTANE.

AMINOCARB in preparations containing 25 per cent or less of aminocarb.

2-AMINO-5-DIETHYLAMINO TOLUENE.

2-AMINO-5-N-ETHYL-N-B(HYDROXY ETHYL) AMINO TOLUENE.

2-AMINO-5-N-ETHYL-N-B(METHANE SULPHONAMIDE ETHYL) AMINO TOLUENE.

2-AMINO-5-N-ETHYL-N-B(METHOXYETHYL AMINO TOLUENE) DI-p-TOLUENE.

AMITRAZ.

AMMONIA (excluding its salts and derivatives other than ammonium hydroxide) except—

(a) in preparations containing 5 per cent or less of free ammonia;

(b) in medicinal preparations for internal use;

(c) in appliances for inhalation in which the substance is absorbed in an inert solid material.

ANILINE (excluding its salts and derivatives), except in preparations containing 1 per cent or less of aniline.

ARECOLINE.

ARSENIC—

(a) in ant poisons containing 0.5 per cent or less arsenic trioxide;

(b) organic compounds of arsenic prepared for use as herbicides or defoliants, except when included in the Fifth Schedule;

(c) in animal feedstuff premixes containing 4 per cent or less of arsenic;

(d) in preparations for therapeutic use in animals except when included in the Fourth Schedule.

AZAMETHIPHOS.

AZOBENZENE.

AZOCYCLOTIN.

BACITRACIN in animal feedstuff premixes for growth promotion purposes containing concentrations greater than 50 mg/kg but not more than 20 000 mg/kg of the total antibiotic principle.

BARBAN.

BARIUM, salts of (except barium sulphate) except—

(a) Paint containing barium metaborate;

(b) when included in the Fifth Schedule.

BENDIOCARB—

(a) in wettable powders containing 80 per cent or less of bendiocarb and when packed in containers or primary packs containing not less than 100 g of bendiocarb;

(b) in wettable powders containing 20 per cent or less of bendiocarb and not less than 0.002 per cent of denatonium benzoate and when packed in containers or primary packs containing not less than 48 g of bendiocarb and when used as a fly control preparation;

(c) in insoluble granular preparations containing 5 per cent or less of bendiocarb;

(d) except when included in the Fifth Schedule.

BENOMYL.

BENQUINOX.

BENSULIDE.

5-BENZYLFUR-3-YLMETHYL (1'R,3'S.E)-2',2'-DIMETHYL-3'-(2-OXO-2,3,4,5-TETRAHYDROTHIENYLIDENEMETHYL)-CYCLOPROPANE CARBOXYLATE.

BENZYLPENICILLIN including procaine penicillin in preparations for intramammary infusion in animals, containing not more than 100 000 international units per dose of benzylpenicillin or procaine penicillin, when suitably coloured with Brilliant Blue FCF or other approved colour as a marker and when packed in applicator devices specially designed for the purpose.

BERYLLIUM.

BHC (excluding the gamma-isomer) except when included in the Fifth Schedule.

BINAPACRYL.

BITHIONOL for treatment of animals.

BRODIFACOUM in preparations containing 0.25 per cent or less.

BROMADIOLONE in preparations containing 0.1 per cent.

BROMOFORM, except for therapeutic use.

BROMOPHOS.

BROMOPHOS-ETHYL.

BROMOXYNIL.

BROTIANIDE.

BUNAMIDINE.
BUTACARB.
BUTOXYCARBOXIM except when included in the Fifth Schedule.
2-BUTOXY-2'-THIOCYANO-DIETHYL ETHER.
BUTYNORATE.

CADMIUM, compounds of, except when included in the Fifth Schedule.
CAMBENDAZOLE.
CARBADOX except in animal feedstuffs containing 50 mg/kg or less of the total active principle.
CARBARYL, except when included in the Second, Fourth or Fifth Schedule.
CARBENDAZIM.
CARBON DISULPHIDE.
alpha-CHLORALOSE, when prepared for use as a pesticide.
CHLORDANE.
CHLORDECONE and substances containing more than 5 per cent of chlordecone.
CHLORFENETHOL.
CHLORMEQUAT.
N-[5-CHLORO-4-[(4-CHLOROPHENYL)-CYANOMETHYL]-2-METHYL-PHENYL] -2-HYDROXY-3,5-DIIODOBENZAMIDE.
CHLOROMETHIURON.
CHLOROPHACINONE.
1-(4-CHLOROPHENOXY)-1-IMIDAZOL-1-YL-3,3-DIMETHYL-2-BUTANONE in concentrations or more than 40 per cent, except when included in the Fourth Schedule.
CHLOROPICRIN in preparations containing 5 per cent or less of chloropicrin.
CHLORPYRIFOS.
CHLORPYRIFOS-METHYL.
CHLORTETRACYCLINE in preparations—

(a) for topical application to animals for ocular use only;
(b) in preparations for intramammary infusion in animals, containing not more than 100 000 international units per dose of chlortetracycline when suitably coloured with Brilliant Blue FCF or other approved colour as a marker and when packed in applicator devices specially designed for the purpose.

CHLORTHIAMID.
5[2-CHLOR-4(TRIFLUOROMETHYL)-PHENOXY]-Z-NITROBENZOATE-4.
CHROMATES AND DICHROMATES.
CHROMIUM TRIOXIDE (excluding its salts and derivatives).
COUMAPHOS in preparations containing 5 per cent or less of coumaphos.
COUMARIN DERIVATIVES and phenylindanedione derivatives not elsewhere included in the Schedules.
COUMATETRALYL.
CROTOXYPHOS.
CRUFOMATE.
CYANAZINE.
CYCLOSULFYNE.
CYHEXATIN.
CYPERMETHRIN, except when included in the Fifth Schedule.
CYTHIOATE.

DAZOMET.
DDT and DDT in preparations containing more than 10 per cent of DDT, except for human therapeutic use.

DELTAMETHRIN in aqueous formulation containing 1 per cent or less of deltamethrin, when no other organic solvent, other than a glycol, is present.

DEMETON-O-METHYL AND DEMETON-S-METHYL in preparations containing 50 per cent or less of one or both demeton-O-methyl and demeton-S-methyl.

DI-ALLATE.

DIAZINON.

DICHLOFENTHION.

DICHLOFLUANID.

DICHLOROETHYLENE.

DICHLOROETHYL ETHER.

1-[2-(2,4-DICHLOROPHENYL)-4-ETHYL-1,3-DIOXOLAN-2-YL-METHYL]-1H-1,2,4-TRIAZOLE.

1-[2-(2,4-DICHLOROPHENYL)-4-PROPYL-1,3-DIOXALAN-2-YL-METHYL]-IH-1,2,4-TRIAZOLE, except when included in the Fifth Schedule.

N-(3,4-DICHLOROPHENYL)-N'-[2-(2"SULFOXY-4'-CHLORPHENOXY)-5 CHLORPHENYL] UREA (SODIUM SALT).

1,2-DICHLOROPROPANE.

1,3-DICHLOROPROPENE.

DICHLORVOS in preparations containing 50 per cent or less dichlorvos, except when included in the Fifth Schedule.

DICLOFOP-METHYL.

DIELDRIN.

DIETHYLENE DIOXIDE.

N,N-DIETHYL-p-PHENYLENE DIAMINE.

DIFFENZOQUAT.

2,3-DIHYDRO-5,6-DIMETHYL-1,4-DITHIIN-1,1,4,4-TETRAOXIDE.

DIHYDROSTREPTOMYCIN in preparations for intramammary infusion in animals containing not more than 100 000 international units per dose of dihydrostreptomycin when suitably coloured with Brilliant Blue FCF or other approved colour as a marker and when packed in applicator devices specially designed for the purpose.

DIMETHANONAPHTHALENE and all substitution or addition, or substitution and addition products of Dimethanonaphthalene not elsewhere specified in these Schedules.

DIMETHOATE.

1,3-DI (METHOXYCARBONYL)-1-PROPEN-2-YL-DIMETHYL PHOSPHATE in preparations containing 25 per cent or less of 1,3-di (methoxycarbonyl)-1-propen-2-yl-dimethyl phosphate.

DIMETHYL FORMAMIDE, except when included in the Fifth Schedule.

2-(2',4'-DEMETHYL-PHENYLIMINO)-3-METHYL-4-THIAZOLINE.

DIMETHYL SULPHOXIDE—

(a) when not packed and labelled for theapeutic use;

(b) for veterinary use when combined with no other therapeutic substance;

(c) for veterinary therapeutic use in preparations containing copper salicylate as the only other therapeutically active ingredient.

DIMETILAN in preparations containing 25 per cent or less of dimetilan.

DIMETRIDAZOLE.

DINITROCRESOLS, DINITROPHENOLS and their homologues in preparations containing 5 per cent or less of such compounds, except for therapeutic use.

DINOCAP.

DIOXACARB.

DIPHACINONE.

DIQUAT.

DISULFIRAM except for therapeutic use.

DISULFOTON in granular preparations containing 5 per cent or less of disulfoton.

DITHIANON.

DITHIAZANINE in preparations containing 2 per cent or less of dithiazanine for the treatment of animals.

DITHIOCARBAMATES when prepared for agricultural, pastoral or horticultural purposes, except when included in the Fifth Schedule.

3,3'-DI-(TRIFLUOROMETHYL)-4,4'-DICHLORO-N,N'-DIPHENYLUREA.

DIUREDOSAN.

ECONAZOLE for external animal use.

ENDOSULFAN.

ENDOTHAL.

EPICHLOROHYDRIN except in preparations containing 2 per cent or less of epichlorohydrin.

ERYTHROMYCIN in preparations—

(a) for intramammary infusion in animals, containing not more than 100 000 international units per dose of erythromycin, when suitably coloured with Brilliant Blue FCF or other approved colour as a marker and when packed in applicator devices specially designed for the purpose;

(b) in animal feedstuff premixes for growth promotion purposes containing concentrations greater than 50 mg/kg but not more than 20 000 mg/kg of the total antibiotic principle.

ETHER SOLVENT.

ETHIOFENCARB.

ETHOATE-METHYL.

ETHOPROPHOS in granular formulations containing 10 per cent or less of ethoprophos.

ETHYL BROMIDE.

ETHYLENE CHLOROHYDRIN.

ETHYLENE DICHORIDE.

ETHYLENE GLYCOL, when packed and labelled as an anti-freeze, except when included in the Fifth Schedule.

ETHYLENE OXIDE.

ETRIDIAZOLE.

FAMPHUR in preparations containing 20 per cent or less of famphur.

FENAMINOSULF in preparations containing 10 per cent or less of fenaminosulf when labelled and packed as dry seed dressings.

FENAMIPHOS in granular preparations containing 5 per cent or less of fenamiphos.

FENAZAFLOR.

FENCHLORPHOS.

FENITROTHION.

FENTHION, except when included in the Fifth Schedule.

FENVALERATE.

FERBAM.

FERROCYANIDES AND FERRICYANIDES, except in preparations containing 1 per cent or less of such substances.

FLAVOPHOSPHOLIPOL in animal feedstuff premixes for growth promotion purposes containing concentrations greater than 50 mg/kg but not more than 20 000 mg/kg of the total antibiotic principle.

FLUAZIFOP-BUTYL.

FORMALDEHYDE (excluding its derivatives), except in preparations containing 5 per cent or less of formaldehyde.

FORMOTHION.

FOSPIRATE, except when included in the Fifth Schedule.

FUMAGILLIN.

GLUTARALDEHYDE, except when included in the Second Schedule or in the Fifth Schedule.

GUAZATINE.

HCB.

HEXACHLOROPHANE in preparations for the treatment of animals.

HYDRAZINE.

HYDROCHLORIC ACID (excluding its salts and derivatives) except in preparations containing 10 per cent or less of hydrochloric acid (HCl).

HYDROFLUORIC ACID AND HYDROSILICOFLUORIC ACID AND OTHER FLUORINE COMPOUNDS except—

(a) when used for human therapeutic purposes;

(b) in dentifrices containing 1 000 mg/kg or less of fluoride ion;

(c) in preparations containing 3 per cent or less of sodium fluoride or sodium silicofluoride when used as preservatives;

(d) when included in the Second, Fourth, Fifth or Seventh Schedule;

(e) in substances containing 15 mg/kg or less of fluoride ion;

(f) ammonium fluosilicate in preparations containing 3.2 per cent or less of ammonium fluosilicate for pesticide purposes.

HYDROQUINONE except—

(a) when included in the Fourth Schedule;

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

8-HYDROXYQUINOLINE for topical use on animals.

HYGROMYCIN in animal feedstuff premixes for use as an anthelmintic containing concentrations greater than 50 mg/kg but not more than 20 000 mg/kg of hygromycin.

IMIDOCARB.

IODINE (excluding its salts and derivatives)—

(a) in iodophors, except in preparations containing 1.5 per cent or less of available iodine;

(b) in other liquid preparations containing 2.5 per cent or less of available iodine;

(c) in preparations for animal treatment only, except in iodophors containing 1.5 per cent or less of available iodine or in solid or semi-solid preparations in containing 2.5 per cent or less of available iodine.

IOXYNIL.

IRON compounds, in preparations for the treatment of animals, except in preparations containing 5 per cent or less of iron.

ISOCYANATES free organic, except in paints containing 0.1 per cent or less of free organic isocyanates.

KITASAMYCIN in animal feedstuffs premixes for growth promotion purposes containing concentrations greater than 100 mg/kg but not more than 20 000 mg/kg of the total antibiotic principle.

LASALOCID, except in animal feedstuffs containing 100 mg/kg or less of the total active antibiotic principle.

LAURYLISOQUINOLINIUM BROMIDE.

LEAD COMPOUNDS, except—

(a) in preparations for therapeutic use;

(b) in preparations for cosmetic use, except when included in the Fifth Schedule or containing 250 ppm or less of lead;

(c) in pencil cores, finger colours, showcard colours, pastels, crayons, poster paints or colours, or coloured chalk containing 0.01 per cent or less of lead.

LINDANE, except when included in the Second or Fifth Schedule.

MALDISON, except when included in the Second or Fifth Schedule.

MEBENDAZOLE for the treatment of animals.

MECLOFENAMIC ACID for the treatment of animals.

MENAZON.

MERCURIC IODIDE when prepared for use for agricultural, industrial, pastoral or horticultural purposes.

MERCURIC THIOCYANATE when prepared for use for photographic purposes.

MERCUROUS CHLORIDE, except preparations for internal use.

MERCURY, organic compounds of, when prepared for use for agricultural, pastoral or horticultural purposes, except when included in the Seventh Schedule.

METACRESOLSULPHONIC ACID AND FORMALDEHYDE CONDENSATION PRODUCT for animal use.

METALDEHYDE, except when included in the Fifth Schedule.

METAXAMINE.

METHACRIFOS.

METHAM.

METHIOCARB, except when included in the Fifth Schedule.

METHOMYL in fly-baits containing one per cent or less of methomyl and not less than 0.002 per cent of denatonium benzoate as a bittering agent.

METHYL ALCOHOL (excluding its salts and derivatives), except in methylated spirit.

N-METHYL CARBAMATES (as pesticides except when specifically included in any other Schedule).

METHYL CHLORIDE.

METHYLENE BISTHIOCYANATE, except in preparations containing one per cent or less of methylene bisthiocyanate.

METHYL ISOTHIOCYANATE.

1-(B METHYL SULPHONAMIDE ETHYL)-2-AMINO-3-N,N-DIETHYL-AMINO BENZENE.

MOLINATE.

MONENSIN in animal feedstuff premixes containing greater than 33 mg/kg but not more than 125 000 mg/kg of the total active principle.

NABAM—see DITHIOCARBAMATES.

NALED, except when included in the Fifth Schedule.

NAPHTHALOPHOS when specifically prepared and packed for use as a sheep drench.

NARASIN in animal feed premixes containing 120 g/kg or less of narasin.

NEOMYCIN in preparations for topical application to animals for ocular use only.

NICOTINE in preparations containing 3 per cent or less of nicotine when labelled and packed for the treatment of animals.

NIMIDANE in preparations containing 25 per cent or less of nimidane.

NITHIAMIDE, except in preparations containing 20 per cent or less of nithiamide.

NITRIC ACID (excluding its salts and derivatives), except in preparations containing 10 per cent or less of nitric acid as such.

NITROBENZENE except—

(a) in solid or semi-solid polishes;

(b) in soaps containing 1 per cent or less of nitrobenzene;

(c) in preparations containing 0.1 per cent or less of nitrobenzene.

NITROPHENOLS, ORTHO, META AND PARA.

NITROSCANATE.

NITROXYNIL.

NOVOBIOCIN in preparations for intramammary infusion in animals, containing not more than 100 000 international units per dose of novobiocin, when suitably coloured with Brilliant Blue FCF or other approved colour as a marker and when packed in applicator devices specially designed for the purpose.

2-n-OCTYL-4-ISOTHIAZOLIN-3-ONE.

OESTRADIOL-17-beta—

(a) in ear implants for growth promotion in bovine cattle;

(b) in combination with progesterone or testosterone in ear implants for growth promotion in bovine cattle.

OLAQUINDOX when intended for use as a growth promotant in pigs, except in animal feedstuffs containing 100 mg/kg or less of the total active principle.

OLEANDOMYCIN in animal feedstuff premixes for growth promotion purposes containing concentrations greater than 50 mg/kg but not more than 20 000 mg/kg of the total antibiotic principle.

OMETHOATE in preparations containing 50 per cent or less of omethoate.

ORGANO-TIN COMPOUNDS, being di-alkyl, tri-alkyl and triphenyl tin compounds where the alkyl group is methyl, ehtyl, propyl or butyl not elsewhere included in these schedules except—

(a) in plastics;

(b) in paints containing 3 per cent or less of such compounds calculated as a proportion of the non-volatile content of the paint;

(c) in other preparations containing 1 per cent or less of such compounds.

ORTHODICHLOROBENZENE.

OXADIAZON.

OXALIC ACID (excluding its salts and derivatives) and soluble oxalates.

OXANTEL EMBONATE for the treatment of animals.

OXFENDAZOLE.

OXYCLOZANIDE.

OXYTETRACYCLINE in preparations—

(a) for topical application to animals for ocular use only;

(b) for intramammary infusion in animals, containing not more than 100 000 international units per dose of oxytetracycline, when suitably coloured with Brilliant Blue FCF or other approved colour as a marker and when packed in applicator devices specially designed for the purpose.

PARAQUAT in granular preparations containing 3 per cent or less of paraquat.

PARBENDAZOLE.

PENTACHLOROPHENOL, except in preparations containing 0.5 per cent or less of pentachlorophenol.

PARACETIC ACID, except when included in the Fifth Schedule.

PERFLUIDONE.

PERMANGANATES.

PHENETHICILLIN in preparations for intramammary infusion in animals, containing not more than 100 000 international units per dose of phenethicillin, when suitably coloured with Brilliant Blue FCF or other approved colour as a marker and when packed in applicator devices specially designed for the purpose.

PHENKAPTON in preparations containing 50 per cent or less of phenkapton.

PHENOL and any homologue of phenol boiling below 220°C, and creosote, except preparations containing 3 per cent or less by weight of such substances or homologues for therapeutic use.

PHENOXYMETHYL PENICILLIN in preparations for intramammary infusion in animals, containing not more than 100 000 international units per dose of phenoxymethyl penicillin, when suitably coloured with Brilliant Blue FCF or other approved colour as a marker and when packed in applicator devices specially designed for the purpose.

PHENYLENE DIAMINES and alkylated phenylene diamines, not elsewhere specified in this Schedule—

(a) when used in hair dyes;

(b) in preparations packed and labelled for photographic purposes;

(c) in preparations packed and labelled for testing water except diethyl—or dimethyl-para-phenylene diamine in tablets containing 10 mg or less in opaque strip packaging labelled for water testing.

PHOSALONE.

PHOSMET.

PHOSPHIDES, METALLIC.

PHOSPHORUS YELLOW (excluding its salts and derivatives) in preparations containing 0.5 per cent or less of free phosphorus.

PHOXIM.

PICRIC ACID (excluding its derivatives), except in preparations containing 5 per cent or less of picric acid.

PINDONE.

PIPEROPHOS.

PIRIMICARB, except when included in the Fifth Schedule.

PIRIMIPHOS-ETHYL.

PIRIMIPHOS-METHYL.

POTASSIUM BROMATE, except in preparations containing 0.5 per cent or less of potassium bromate.

POTASSIUM CYANATE.

POTASSIUM HYDROXIDE, except in preparations containing 5 per cent or less of potassium hydroxide.

PROFENOFOS.

PROGESTERONE in a silicon rubber elastomer when used as a controlled release implant for synchronisation of oestrus in cattle.

PROMACYL.

PROMECARB in preparations containing 50 per cent or less of promecarb.

PROPACHLOR.

PROPETAMPHOS.

PROPIONIC ACID (excluding its salts and derivatives) in preparations containing more than 80 per cent propionic acid, except for therapeutic use.

PROPOXUR, except when included in the Second or Fifth Schedule.

PROTHIOPHOS.

PYRAZOPHOS.

PYRINURON except when included in the Fifth Schedule.

RAFOXANIDE.

SALINOMYCIN in animal feedstuff premixes containing greater than 60 mg/kg but not more than 60 000 mg/kg of the total active antibiotic principle.

SELENIUM, COMPOUNDS OF—

(a) in preparations containing 2.5 per cent or less of selenium—

(i) when packed and labelled for the blueing of gun barrels;

(ii) when packed and labelled for photographic purposes;

(b) in preparations containing 2.5 per cent or less of selenium when packed and labelled as vaccines, drenches or pastes for treatment of animals;

(c) in preparations containing 0.5 per cent or less of selenium when packed and labelled as other injections for treatment of animals;

(d) in premixes containing 2 per cent or less of selenium when packed and labelled for incorporation into animal feeds to provide 0.1 g/tonne or less of selenium.

SODIUM BROMATE, except in preparations containing 0.5 per cent or less of sodium bromate.

SODIUM HYDROXIDE (excluding its salts and derivatives), except—

(a) in preparations containing 0.5 per cent or less of sodium hydroxide;

(b) when included in the Fifth Schedule.

SPIRAMYCIN in animal feedstuff premixes for growth promotion purposes containing concentrations greater than 50 mg/kg but not more than 20 000 mg/kg of the total antibiotic principle.

STREPTOMYCIN in preparations for intramammary infusion in animals, containing not more than 100 000 international units per dose of streptomycin, when suitably coloured with a Brilliant Blue FCF or other approved colour as a marker and when packed in applicator devices specially designed for the purpose.

STRYCHNINE in grain baits containing 0.5 per cent or less of strychnine and registered as a pesticide.

SULPHANILAMIDE and its derivatives unless elsewhere specified in this Schedule when packed and labelled for treatment of ornamental caged birds or ornamental fish only.

SULPHAQUINOXALINE when packed and labelled for use as a coccidiostat in poultry except preparations containing 200 mg/kg or less of sulphaquinoxaline.

SULPHURIC ACID (excluding its salts and derivatives), except—

(a) in accumulators, batteries and fire extinguishers;

(b) in preparations containing 0.5 per cent or less of sulphuric acid (H^2SO^4).

SULPROPHOS.

2,4,5-T.

TCA—see TRICHLOROACETIC ACID.

TCMTB (2-[thiocyanomethylthio]benzothiazole).

TDE, except when included in the Fifth Schedule.

TEMEPHOS.

TERBUTHYLAZINE.

TERPENES, CHLORINATED.

TESTOSTERONE PROPIONATE, TESTOSTERONE DIPROPIONATE and TESTOSTERONE ENANTHATE in preparations labelled for treatment and prevention of pizzle (sheath) rot in wethers, and in preparation labelled for masculinisation of wethers for use as "teaser rams" to stimulate and detect reproductive activity in ewes.

TETRACHLOROETHYLENE, except—

(a) when prepared for use for the treatment of humans and for the treatment of animals;

(b) when packed in containers of 50 mls or less;

(c) when included in the Fifth Schedule.

TETRACYCLINE in preparations—

(a) for topical application to animals for ocular use only;

(b) for intramammary infusion in animals, containing not more than 100 000 international units per dose of tetracycline, when suitably coloured with a Brilliant Blue FCF or other approved colour as a marker and when packed in applicator devices specially designed for the purpose;

(c) when packed and labelled for treatment of ornamental caged birds or ornamental fish only.

TETRADIFON

TETRAMISOLE, including levamisole, in preparations for the treatment of animals except when included in the Fourth or Fifth Schedule.

THIAZAFLURON.

THIODICARB.

THIOMETON.

THIOUREA except for therapeutic use.

THIRAM.

TIAMULIN for animal use—

(a) in feedstuff premixes containing 25 per cent or less of tiamulin;

(b) in soluble concentrates containing 45 per cent or less of tiamulin.

ortho-TOLIDINE when packed and labelled in concentrations of 0.1 per cent or less of ortho-tolidine for the testing of water.

TOLUENE (excluding its derivatives), when packed in containers of 20 litres or less, except—

(a) in preparations containing 50 per cent or less of toluene or of both toluene and xylene;

(b) when packed in containers of 50 ml or less.

TRIADIMEFON, except when included in the Fifth Schedule.

S,S,S-TRIBUTYLPHOSPHOROTHIOATE.

TRICHLORFON.

TRICHLOROACETIC ACID, except when included in the Fifth Schedule.

TRICHLOROETHYLENE, except—

(a) when specifically prepared for medicinal purposes;

(b) when packed in containers of 50 ml or less.

TRICHLOROPHENOL.

TRICLOPYR.

TRIDEMORPH.

TRIETHYL PHOSPHATE.

TRIFLUOROMETHANE SULPHONIC ACID.

TYLOSIN and its salts in animal feedstuff premixes for growth promotion purposes containing concentrations greater than 50 mg/kg but not more 20 000 mg/kg of the total antibiotic principle.

VAMIDOTHION.

VIRGINIAMYCIN in animal feedstuff premixes for growth promotion purposes containing concentrations greater than 50 mg/kg but not more than 20 000 mg/kg of the total antibiotic principle.

WARFARIN, except—

(a) for therapeutic use;

(b) when included in the Fifth Schedule.

XYLENE (excluding its derivatives), when packed in containers of 20 litres or less except—

(a) in preparations containing 50 per cent or less of xylene or of both xylene and toluene;

(b) when packed in containers of 50 ml or less.

ZERANOL, in implants for use as a growth promotant in steer cattle.
ZINC CHLORIDE, except in preparations containing 5 per cent or less of zinc chloride.
ZINC p-PHENOLSULPHONATE, except in preparations containing 5 per cent or less of zinc p-phenolsulphonate.
ZINC SULPHATE, except for human therapeutic use and in preparations containing 5 per cent or less of zinc sulphate.

Excluding however, the substances hereinbefore mentioned when contained in the products listed as exemptions, set out after the Eighth Schedule in this Appendix.

Seventh Schedule.

ACROLEIN.
ALDICARB.
ALLYL ALCOHOL.
AMINOCARB, except when included in the Sixth Schedule.
4-AMINOPYRIDINE.
AMITON.
ANTU.
APRINOCID.
ARSENIC, except—
(a) for the specific purposes shown in the Fourth, Fifth or Sixth Schedule;
(b) in animal feedstuffs containing 75 mg/kg or less of arsenic.
AZINPHOS-ETHYL.
AZINPHOS-METHYL.

BENDIOCARB, except when included in the Fifth or Sixth Schedule.
BENZENE (excluding its derivatives) except—
(a) in preparations containing 1.5 per cent v/v or less of benzene;
(b) petrol containing 5 per cent v/v or less of benzene.
Note—see also "Carcinogenic Substances".
BETAHYDROXYETHYLHYDRAZINE.
BRODIFACOUM, except when included in the Sixth Schedule.
BROMADIOLONE, except when included in the Sixth Schedule.

CAMPHECHLOR.
CAPTAFOL.
CARBOFURAN.
CARBON TETRACHLORIDE.
CARBOPHENOTHION.

CARCINOGENIC SUBSTANCES—

2-Acetyl Aminofluorene
Acrylonitrile
Alphanaphthylamine
4-Aminobiphenyl
Benzene, (excluding derivatives) except—

(a) preparations containing 1.5 per cent v/v or less of benzene;

(b) petrol containing 5 per cent v/v or less of benzene.

Benzidine
Benzo(a)pyrene
Betanaphthylamine
Beta Propriolactone
Bis-Chloromethyl Ether
1,2-Dibromo-3-chloropropane
3,3'-Dichlorobenzidine
4-Dimethylamino Azobenzene
Methyl Chloromethyl Ether
4,4-Methylene Bis-(2-Chloroaniline)
4-Nitrobiphenyl
N-Nitrosodimethylamine
PCBs (polychlorinated biphenyls)
Toxaphene (Camphechlor)
Vinyl Chloride monomer.

CHLORAMPHENICOL for systemic use in food producing animals on a flock or herd basis.

CHLORDIMEFORM.

CHLORFENVINPHOS.

CHLORINE (excluding its salts and derivates).

5-CHLORO-3-METHYL-4-NITROPYRAZOLE.

CHLOROPICRIN, except when included in the Sixth Schedule.

CHLORTHIOPHOS, except when included in the Sixth Schedule.

COUMAPHOS, except when included in the Sixth Schedule.

CYANIDES—see hydrocyanic acid.

CYHALOTHRIN.

DELTAMETHRIN, except when included in the Sixth Schedule.

DEMETON.

DEMETON-0-METHYL and DEMETON-S-METHYL, except when included in the Sixth Schedule.

DIALIFOS.

1,2-DIBROMO-3-CHLOROPROPANE.

DICHLORVOS, except when included in the Fifth or Sixth Schedule.

DICROTOPHOS.

DIENOCHLOR.

DIMEFOX.

1,3-DI(METHOXYCARBONYL)-1-PROPEN-2-YL-DIMETHYL PHOSPHATE, except when included in the Sixth Schedule.

DIMETILAN, except when included in the Sixth Schedule.

DINITROCRESOLS, DINITROPHENOLS and their homologues and in preparations containing more than 5 per cent of such compounds, except for therapeutic use.

DIOXATHION.

DISULFOTON, except when included in the Sixth Schedule.

ENDOTHION.

ENDRIN.

ETHION.
ETHOPROPHOS, except when included in the Sixth Schedule.
ETHOXYETHYL MERCURY CHLORIDE.
ETHYLENE DIBROMIDE, except in and for the preparation of motor fuels.
ETHYL MERCURY CHLORIDE.

FAMPHUR, except when included in the Sixth Schedule.
FENAMINOSULF, except when included in the Sixth Schedule.
FENAMIPHOS, except when included in the Sixth Schedule.
FENSULFOTHION.
FENTHION-ETHYL.
FLUCYTHRINATE.
FLUNIXIN MEGLUMINE, except when included in the Fourth Schedule.
FLUORACETAMIDE.
FLUOROACETIC ACID.
FORMETANATE.

HALOFUGINONE, except in prepared stockfeeds containing 3 g/tonne or less of halofuginone.
HYDROCYANIC ACID, except—
(a) in preparations containing the equivalent of 0.15 per cent or less of hydrocyanic acid;
(b) for therapeutic use.

ISOCARBOPHOS
ISOFENPHOS.
IVERMECTIN.

LEPTOPHOS.

MAZIDOX.
MECARBAM.
MERCURIC CHLORIDE when prepared for use for agricultural, industrial, pastoral or horticultural purposes.
METHAMIDOPHOS.
METHAPYRILENE.
METHFUROXAM.
METHIDATHION.
METHOMYL, except when included in the Sixth Schedule.
METHYL BROMIDE.
MEVINPHOS.
MIPAFOX.
MIREX.
MONOCROTOPHOS.

NAPHTHALOPHOS, except when included in the Sixth Schedule.
NICOTINE except—
(a) when included in the Fourth or Sixth Schedule;
(b) in tobacco.
NIMIDANE, except when included in the Sixth Schedule.
NITROFEN.

OMETHOATE, except when included in the Sixth Schedule.
OXAMYL.
OXYFLUORFEN.

PARAQUAT, except when included in the Sixth Schedule.
PARATHION.
PARATHION-METHYL.
PHENKAPTON, except when included in the Sixth Schedule.
PHORATE.
PHOSFOLAN.
PHOSPHAMIDON.
POLYCHLORINATED BIPHENYLS see "Carcinogenic Substances".
PROMECARB, except when included in the Sixth Schedule.
PROTHOATE.

SCHRADAN.
STRYCHNINE, except when included in the First, Fourth or Sixth Schedule.
SULFALLATE.
SULFOTEP.
SULPHATROXAZOLE, except when included in the Fourth Schedule.

TEPP.
TERBUFOS.
TETRACHLOROETHANE.
THALIDOMIDE.
THALLIUM.
THIOFANOX.
THIONAZIN.
ortho-TOLIDINE, except when included in the Sixth Schedule and in solid-state diagnostic therapeutic reagents.
TRIAMIPHOS.
TRIAZBUTIL.
TRICHLOROISOCYANURIC ACID except—
(a) in preparations containing 4 per cent or less of available chlorine;
(b) when included in the Fifth Schedule.

VINYL CHLORIDE MONOMER see "Carcinogenic Substances".

Eighth Schedule.

ACETORPHINE (0^3-acetyl-7, 8 dihydro-7a (1 (R)-hydroxy-1-methylbutyl)-0^6-methyl-6, 14-endoetheno-morphine).
ACETYLDIHYDROCODEINE, except when included in the Second or Fourth Schedule.
ACETYLMETHADOL (3-acetoxy-6-dimethylamino-4, 4-diphenylheptane).
ALLYLPRODINE (3-allyl-1-methyl-4-phenyl-4-propionoxypiperidine).
ALPHACETYLMETHADOL (alpha-3-acetoxy-6-dimethylamino-4, 4-diphenylheptane).

ALPHAMEPRODINE (alpha-3-ethyl-1-methyl-4-phenyl-4-propionoxypiperidine).

ALPHAMETHADOL (alpha-6-dimethylamino-4, 4-diphenyl-3-heptanol).

ALPHAPRODINE (alpha-1, 3-dimethyl-4-phenyl-4-propionoxypiperidine).

2-AMINO-1-(2,5-DIMETHOXY-4-METHYLPHENYL) PROPANE (STP, DOM).

AMPHETAMINE.

ANILERIDINE (1-para-aminophenethyl-4-phenylpiperidine-4-carboxylic acid ethyl ester).

BENZETHIDINE (1-(2-Benzyloxyethyl)-4-phenylpiperidine-4-carboxylic acid ethyl ester).

BENZYLMORPHINE (3-benzylmorphine).

BETACETYLMETHADOL (beta-3-acetoxy-6-dimethylamino-4, 4-diphenylheptane).

BETAMEPRODINE (beta-3-ethyl-1-methyl-4-phenyl-4-propionoxypiperidine).

BETAMETHADOL (beta-6-dimethylamino-4, 4-diphenyl-3-heptanol).

BETAPRODINE (beta-1,3-dimethyl-4-phenyl-4-propionoxypiperidine).

BEZITRAMIDE (1-(3-cyano-3,3-diphenylpropyl)-4-(2-oxo-3-propionyl-1-benzimidazolinyl)piperidine).

BUFOTENINE.

CANNABIS AND CANNABIS RESIN and extracts of tinctures of cannabis.

CLONITAZENE (2-para-chlorobenzyl-1-diethylaminoethyl-5-nitro-benzimidazole).

COCAINE (methyl ester of benzoylecgonine), and any solution or dilution in an inert substance whether liquid or solid in any proportion and all preparations and admixtures.

COCA LEAF.

CODEINE (3-methylmorphine), except when included in the Second, Third or Fourth Schedule.

CODEINE-N-OXIDE.

CODOXIME (dihydrocodeinone-6-carboxymethyloxime).

CONCENTRATE OF POPPY STRAW (the material arising when poppy straw has entered into a process of concentration of its alkaloids).

DESOMORPHINE.

DEXAMPHETAMINE.

DEXTROMORAMIDE ((+)-4-(2-methyl-4-oxo-3, 3diphenyl-4-(1-pyrrolidinyl butyl) morpholine).

DIAMPROMIDE (N(-2-(methylphenethylamino) propyl) propionanilide).

DIETHYLTHIAMBUTENE (3-diethylamino-1,1-di-(2'-thienyl)-1-butene).

N,N-DIETHYLTRYPTAMINE.

DIFENOXIN (1-(3-cyano-3,3-diphenylpropyl)-4-phenylisonipecotic acid) except when included in the Fourth Schedule.

DIHYDROCODEINE, except when included in the Second or Fourth Schedule.

DIHYDROMORPHINE.

DIMENOXADOL (2-dimethylaminoethyl-1-ethoxy-1, 1-diphenylacetate).

DIMEPHEPTANOL (6-dimethylamino-4,4-diphenyl-3-heptanol).

3-(1,2-DIMETHYLHEPTYL)-1-HYDROXY-7,8,9,10-TETRAHYDRO-6,6,9-TRIMETHYL-6H-DIBENZO(b,d) PYRAN (DMPH).

DIMETHYLTHIAMBUTENE (3-dimethylamino-1,1-di-(2'-thienyl)-1-butene).

N,N-DIMETHYLTRYPTAMINE.

DIOXAPHETYL BUTYRATE (ethyl 4-morpholino-2,2-diphenylbutyrate).

DIPHENOXYLATE (1-(3-cyano-3,3-diphenylpropyl)-4-phenylpiperidine-4-carboxylic acid ethyl ester), except when included in the Fourth Schedule.

DIPIPANONE (4,4-diphenyl-6-piperidine-3-heptanone).
DROTEBANOL (3,4-dimethoxy-17-methylmorphinan-6 B, 14 diol).

ECGONINE, its esters and derivatives which are convertible to ecgonine and cocaine.
ETHYLMETHYLTHIAMBUTENE (3-ethylmethylamino-1, 1-di-(2'-thienyl) -1-butene).
ETHYLMORPHINE (3-ethylmorphine), except when included in the Second or Fourth Schedule.
N-ETHYL-1-PHENCYCLOHEXYLAMINE (PCE).
ETONITAZENE (1-diethylaminoethyl-2-para-ethoxybenzyl-5-nitrobenzimidazole).
ETORPHINE (7,8-dihydro-7a-(1(R)-hydroxy-1-methyl-butyl)-0^6, methyl-6,14-endoethenomorphine).
ETOXERIDINE (1-(2-(2-hydroxyethoxy) ethyl)-4-phenylpiperidine-4-carboxylic acid ethyl ester).

FENTANYL (1-phenethyl 4-N-propionyl-anilino piperidine).
FURETHIDINE (1-(2-tetrahydrofurfuryloxyethyl)-4-phenylpiperidine-4-carboxylic acid ethyl ester).

HEPTANE DERIVATIVES—having addiction properties, not specifically included in this Schedule.
HEROIN.
3-HEXYL-1-HYDROXY-7,8,9,10-TETRAHYDRO-6,6,9-TRIMETHYL-6H-DIBENZO (b,d) PYRAN (PARAHEXYL).
HYDROCODONE (dihydrocodeinone).
HYDROMORPHINOL (14-hydroxydihydromorphine).
HYDROMORPHONE (dihydromorphinone).
HYDROXYPETHIDINE (4-meta-hydroxyphenyl-1-methylpiperidine-4-carboxylic acid ethyl ester).

ISOMETHADONE (6-dimethylamino-5-methyl-4,4-diphenyl-3-hexanone).

KETOBEMIDONE (4-meta-hydroxyphenyl-1-methyl-4-propionylpiperidine).

LEVOMETHORPHAN ((-)-3-methoxy-N-methylmorphinan).
LEVOMORAMIDE ((-)-4-(2-methyl-4-oxo-3,3-diphenyl-4-(1-pyrrolidinyl) butyl) morpholine).
LEVOPHENACYLMORPHAN ((-)-3-hydroxy-N-phenacylmorphinan).
LEVORPHANOL ((-)-3-hydroxy-N-methylmorphinan).
LYSERGIC-ACID DIETHYLAMIDE (LSD).

MECLOQUALONE 3-(o-chlorophenyl)-2-methyl-4-(3H) quinazolinone.
MESCALINE, 2,5-DIMETHOXY-4-METHYLAMPHETAMINE, and other substances structurally derived from methoxyphenylethylamine having hallucinogenic properties.
METAZOCINE (2'-hydroxy-2,5,9,-trimethyl-6,7-benzomorphan).
METHADONE (6-dimethylamino-4,4-diphenyl-3-heptanone).
METHADONE INTERMEDIATE (4-cyano-2-dimethylamino-4,4-diphenylbutane).
METHAQUALONE.
METHYLAMPHETAMINE.
METHYLDESORPHINE (6-methyl-delta-6-desoxymorphine).

METHYLDIHYDROMORPHINE (6-methyldihydromorphine).
METHYLPHENIDATE.
1-METHYL-4-PHENYLPIPERIDINE-4-CARBOXYLIC ACID ESTERS.
METOPON (5-methyldihydromorphinone).
MORAMIDE INTERMEDIATE (2-methyl-3-morpholino-1, 1-diphenylpropane carboxylic acid).
MORPHERIDINE (1-(2-morpholinoethyl)-4-phenylpiperidine-4-carboxylic acid ethyl ester).
MORPHINE.
MORPHINE DERIVATIVES not specifically included in this Schedule.
MORPHINE METHOBROMIDE AND OTHER PENTAVALENT NITROGEN MORPHINE DERIVATIVES.
MORPHINE-N-OXIDE.
MORPHINE SUBSTITUTES not specifically included in this Schedule.
MYROPHINE (myristylbenzylmorphine).

NABILONE.
NICOCODINE (6-nicotinylcodeine), except when included in the Second or Fourth Schedule.
NICODICODINE (6-nicotinoyldihydrocodeine), except when included in the Second or Fourth Schedule.
NICOMORPHINE (3,6-dinicotinylmorphine).
NORACYMETHADOL ((±)alpha-3-acetoxy-6-methylamino-4,4-diphenylheptane).
NORCODEINE (N-demethylcodeine), except when included in the Second or Fourth Schedule.
NORLEVORPHANOL ((-)3-hydroxymorphinan).
NORMETHADONE (6-dimethylamino-4,4-diphenyl-3-hexanone).
NORMORPHINE (N-demethylated morphine).
NORPIPANONE (4,4-diphenyl-6-piperidine-3-hexanone).

OPIUM, in any form, except the alkaloids noscapine and papaverine.
OXYCODONE (14-hydroxydihydrocodeinone).
OXYMORPHONE (14-hydroxydihydromorphinone) except when included in the Fourth Schedule.

PENTAZOCINE.
PETHIDINE (1-methyl-4-phenylpiperidine-4-carboxylic acid ethyl ester).
PETHIDINE INTERMEDIATE A (4-cyano-1-methyl-4-phenylpiperidine).
PETHIDINE INTERMEDIATE B (4-phenylpiperidine-4-carboxylic acid ethyl ester).
PETHIDINE INTERMEDIATE C (1-methyl-4-phenylpiperidine-4-carboxylic acid).
PHENADOXONE (6-morpholino-4, 4-diphenyl-3-heptanone).
PHENAMPROMIDE (N(1-methyl-2-piperidinoethyl) propionanilide).
PHENAZOCINE (2-hydroxy-5,9-dimethyl-2-phenethyl-6,7-benzomorphan).
PHENCYCLIDINE.
1-(1-PHENCYCLOHEXYL) PYRROLIDINE (PHP or PCPY).
PHENMETRAZINE.
PHENOMORPHAN (3-hydroxy-N-phenethylmorphinan).
PHENOPERIDINE (1-(3-hydroxy-3-phenylpropyl)-4-phenylpiperidine-4-carboxylic acid ethyl ester).

PHOLCODINE (morpholinylethyl morphine), except when included in the Second or Fourth Schedule.

PIMINODINE (4-phenyl-1-(-3-phenylaminopropyl) piperidine-4-carboxylic acid ethyl ester).

PIPERIDINE DERIVATIVES having addiction properties, not specifically included in this Schedule.

PIRITRAMIDE (1-(3-cyano-3,3-diphenylpropyl)-4-(1-piperindino) piperidine-4-carboxylic acid amide).

PROHEPTAZINE (1-,3-dimethyl-4-phenyl-4-propionoxyazacycloheptane).

PROPERIDINE (1-methyl-4-phenylpiperidine-4-carboxylic acid isopropyl ester).

PROPIRAM.

PSILOCIN.

PSILOCYBIN.

RACEMETHORPHAN ((±)-methoxy-N-methylmorphinan).

RACEMORAMIDE((±)-4-(2-methyl-4-oxo-3,3-diphenyl-4-(1-pyrrolidinyl) butyl) morpholine).

RACEMORPHAN ((±)-3-hydroxy-N-methylmorphinan).

SUFENTANIL N-(4-(methoxymethyl)-1-(2-thienyl)-ethyl-4-piperidyl) propionanilide.

TETRAHYDROCANNABINOLS and 3- AND 4'-ALKYL homologues, including DMPH and PARAHEXYL, within one of those structural designations.

THEBACON (acetyl dihydrocodeinone).

THEBAINE.

1-(1-(2-THIENYL)CYCLOHEXYL) PIPERIDINE (TCP).

TILIDINE (±) ethyl-trans-2-(dimethylamine)-1-phenyl-3-cyclohexene-1-carboxyiate.

TRIMEPERIDINE (1,2,5-trimethyl-4-phenyl-4-propionoxypiperidine).

Exemptions.

The requirements of the Poisons Act 1964 and regulations made under that Act do not apply to the following products when containing a scheduled poison—

Timber and wallboard

Ceramics

Electrical components and electric lamps

Vitreous enamels

Explosives

Glazed pottery

Matches

Motor fuels, other than those containing methyl alcohol, unless specified in any of the Schedules.

Lubricants, unless specified in any of the Schedules

Paper

Photographic paper and film

Inorganic pigments unless specified in the Sixth Schedule

Paints, other than prepared for medicinal or cosmetic purposes.

Blankets moth proofed with dieldrin in the mill during finishing as directed by C.S.I.R.O.

Single-use tubes for the estimation of alcohol content of breath. ".

By His Excellency's Command,

R. G. COOPER,

Clerk of the Council.

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