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No. 62] PERTH: TUESDAY, 29th JUNE [1965

AT a meeting of the Executive Council held in the Executive Council Chamber, at Perth, this 23rd day of June, 1965, the following Order in Council was authorised to be issued:—

Poisons Act, 1964.

ORDER IN COUNCIL.

WHEREAS by section 21 of the Poisons Act, 1964, it is provided that the Governor may from time to time, by Order in Council, amend any of the Schedules referred to in section 20 of that Act by—

(a) the addition thereto or the deletion therefrom of any substance;

(b) the transference of any substance from any Schedule to any other Schedule; or

(c) the alteration of any item in any Schedule;

and whereas it is deemed expedient to amend those Schedules by alteration of items therein to the extent that those Schedules are amended to read as hereinafter described: Now, therefore, His Excellency the Governor, acting by and with the advice of the Executive Council, doth hereby amend the Schedules to the said Poisons Act, 1964, by alteration of the items contained therein to the extent that those Schedules are amended to read in the manner set out in the Appendix to this Order in Council, to take effect on and from the day upon which notice of this Order in Council is published in the *Government Gazette*.

W. S. LONNIE,
Clerk of the Council.

APPENDIX.

FIRST SCHEDULE.

A substance specified in this Schedule includes any compound and all preparations and admixtures containing any proportion thereof and these are therefore subject to all the restrictions of this Schedule unless specifically exempted or specifically included in any other Schedule.

ACONITE (ROOT OF ACONITUM NAPELLUS).

ALKALOIDS, the following, their salts, their derivatives and their salts—

APOMORPHINE and substances containing more than 0.2 per cent. of apomorphine.

ATROPINE and substances containing more than 0.25 per cent. of atropine.

BRUCINE and substances containing more than 0.2 per cent. of brucine.

COLCHICINE and substances containing more than 0.5 per cent. of colchicine.

CONIINE and substances containing more than 0.1 per cent. of coniine.

COTARNINE.

HOMATROPINE and substances containing more than 0.25 per cent. of homatropine.

HYOSCINE and substances containing more than 0.25 per cent. of hyoscine.

HYOSCYAMINE and substances containing more than 0.25 per cent. of hyoscyamine.

SPARTEINE.

ANTIMONY and substances containing more than the equivalent of 1 per cent. of antimony trioxide, except chlorides in polishes.

ARSENIC and substances containing more than the equivalent of 0.5 per cent. of arsenic trioxide, except when prepared and packed to comply with the requirements of Schedule 6.

BELLADONNA and substances containing more than 0.25 per cent. of the alkaloids of belladonna calculated as hyoscyamine.

BROMINE as such.

CROTON OIL.

ELATERIUM.

HYDROCYANIC ACID and substances containing more than 0.15 per cent. of hydrocyanic acid, except when included in Schedule 7.

HYOSCYAMUS and substances containing more than 0.25 per cent. of alkaloids calculated as hyoscyamine.

LOBELIA and substances containing more than 0.5 per cent. of alkaloids except for smoking or burning.

MERCURIC CHLORIDE and substances containing more than 0.5 per cent. of mercuric chloride, except in batteries or when included in Schedule 6.

MERCURIC IODIDE and substances containing more than 2 per cent. of mercuric iodide, except when included in Schedule 6.

MERCURIC NITRATE and substances containing more than the equivalent of 3 per cent. of mercury (Hg), in such form.

MERCURIC-POTASSIUM IODIDE and substances containing more than the equivalent of 2 per cent. of mercuric-iodide, in such form.

MERCURY, organic compounds, and substances containing more than the equivalent of 0.5 per cent. of mercury (Hg), in organic combination, except for therapeutic use, or when included in the Sixth Schedule.

NUX VOMICA.

PHOSPHORUS YELLOW and substances containing more than 0.5 per cent. of free yellow phosphorus.

SAVIN, oil of.

STRAMONIUM and substances containing more than 0.25 per cent. of the alkaloids of stramonium except for smoking or burning.

STRYCHNINE and substances containing more than 0.2 per cent. of strychnine.

TANSY, oil of.

VERATUM, its alkaloids, their salts.

SECOND SCHEDULE.

A substance specified in this Schedule includes any compound, and all preparations and admixtures containing any proportion thereof and these are subject to all the restrictions of this Schedule unless specifically exempted or specifically included in any other Schedule.

ACID ACETIC GLACIAL as such.

AMPHETAMINE, its derivatives, in appliances for inhalation in which the substance is absorbed upon inert solid material.

ANTIMONY in substances containing the equivalent of 1 per cent. or less of antimony trioxide, except chlorides in polishes.

APOMORPHINE in substances containing 0.2 per cent. or less of apomorphine.

ARSENIC in substances containing the equivalent of 0.5 per cent. or less of arsenic trioxide, except when prepared and packed to comply with the requirements of Schedule 6.

ATROPINE in substances containing 0.25 per cent. or less of atropine.

BARBITURIC ACID, its derivatives, and their salts in substances containing 0.2 per cent. or less of barbituric acid, its derivatives and their salts.

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

BROMIDE metallic, including ammonium, in medicinal preparations or admixtures containing more than 300 mg. of metallic bromide or ammonium bromide in each adult dose.

BRUCINE in substances containing 0.2 per cent. or less of brucine.

CANTHARIDES (CANTHARIDIN) in substances containing 0.01 per cent. or less of cantharidin.

CHLOROFORM and substances containing more than 10 per cent. of chloroform.

COCAINE, its salts, its derivatives, their salts, in substances containing 0.1 per cent. or less of cocaine.

COCAINE, synthetic substitutes for—capable of use for local anaesthesia having a solubility in water of more than 1 per cent., in ointments containing 0.5 per cent. or less of such substances.

COCAINE, synthetic substitutes for—capable of use for surface anaesthesia having a solubility in water of 1 per cent. or less, in preparations and admixtures containing 2.5 per cent. or less of such substances and—

(1) Lozenges, pastilles, tablets, capsules, containing 30 mg. or less of such substance in each;

(2) Suppositories or bougies containing 200 mg. or less of such substances in each; and

(3) Preparations for external use containing 10 per cent. or less of such substances.

CODEINE in substances containing 1 per cent. or less of codeine

COLCHICINE in sustances containing 0.5 per cent. or less of colchicine.

CREOSOTE in substances containing more than 3 per cent. by weight of creosote.

CRESOL in substances containing more than 3 per cent. by weight of cresol.

DEXTROMETHORPHAN in substances containing 1 per cent. or less of dextromethorphan.

DEXTROPROPOXYPHENE in substances containing 1 per cent. or less of dextropropoxyphene.

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

DIAMINES, and alkylated benzene diamine derivatives, except when included in Schedule 5.

ETHER ANAESTHETIC and substances containing more than 10 per cent. of ether anaesthetic.

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

p-ETHOXY PHENYL UREA.

ETHYL MORPHINE in substances containing 1 per cent. or less of ethyl morphine.

FLUORIDES, metallic, including ammonium fluoride, when intended for ingestion, except in dentifrices containing 0.5 per cent. or less of fluoride.

GELSEMIUM.

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

HYDROCYANIC ACID in substances containing 0.15 per cent. or less of hydrocyanic acid.

HYOSCINE and its derivatives in substances containing 0.25 per cent. or less of hyoscine and its derivatives.

HYOSCYAMINE and its derivatives in substances containing 0.25 per cent. or less of hyoscyamine and its derivatives.

HYOSCYAMUS in substances containing 0.25 per cent. or less of alkaloids calculated as hyoscyamine.

IODINE and solutions containing more than 2.5 per cent. of iodine.

JABORANDI, alkaloids of, and their salts in substances containing more than 0.025 per cent. of the alkaloids of jaborandi.

LEAD SALTS and compounds of lead for medicinal use, except in machine spread plasters.

LOBELIA in substances containing 0.5 per cent. or less of the alkaloids of lobelia except for smoking or burning.

MERCURIC AMMONIUM CHLORIDE (Ammoniated Mercury).

MERCURIC CHLORIDE in substances containing 0.5 per cent. or less of mercuric chloride, except in batteries or when prepared and packed to comply with the requirements of Schedule 6.

MERCURIC IODIDE in substances containing 2 per cent. or less of mercuric iodide.

MERCURIC NITRATE in substances 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 substances containing the equivalent of 2 per cent. or less of mercuric iodide, in such form.

MERCURY (METALLIC) as such.

MERCURY, organic compounds of, in substances containing the equivalent of 0.5 per cent. or less of mercury (Hg) in organic combination, except when included in Schedule 6 or as a preservative in substances containing 0.01 per cent. or less of mercury.

MORPHINE (except derivatives and their salts unless specifically included in this Schedule) in substances containing 0.2 per cent. or less of morphine calculated as anhydrous morphine.

NITRIC ACID and preparations containing more than 10 per cent. of nitric acid.

NITROPHENOLS, ortho, meta, para.

NUX VOMICA in substances containing 0.2 per cent. or less of strychnine.

OPIUM (except its alkaloids, their derivatives, their salts unless specifically included in this Schedule) in substances containing 0.2 per cent. or less of morphine calculated as anhydrous morphine.

OXALIC ACID and metallic oxalates, except in laundry blue and polishes.

PHENOL (CARBOLIC ACID) and its homologues boiling below 220°C. and liquid substances containing more than 3 per cent. by weight of phenol, or its homologues boiling below 220°C.

PHOLCODINE in substances containing 1 per cent. or less of pholcodine.

PILOCARPINE and its salts in substances containing more than 0.025 per cent. of the alkaloid.

POTASSIUM CHLORATE and substances containing more than 10 per cent. of potassium chlorate.

SELENIUM, its salts and compounds, except in substances other than for human therapeutic use containing 2.5 per cent. or less of selenium.

SILVER NITRATE.

STAPHISAGRIA and substances containing more than 0.2 per cent.

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

STRYCHNINE in substances containing 0.2 per cent. or less of strychnine.

SULPHURIC ACID and substances and preparations containing more than 35 per cent. by weight of sulphuric acid (H_2SO_4).

THIRD SCHEDULE.

A substance specified in this Schedule includes any compound, and all preparations and admixtures containing any proportion thereof and these are therefore subject to all the restrictions of this Schedule unless specifically exempted or specifically included in any other Schedule.

ADRENALINE, natural or synthetic, its salts, in concentrations of more than 0.01 per cent. but not exceeding 1 per cent. of the base.

AMYL NITRITE.

ANTIHISTAMINES, all tertiary nitrogenous organic bases which possess pharmacological properties characteristic of antihistamine compounds in preparations labelled and packed for the treatment of motion sickness in packs of 10 doses or less.

BROMVALETONE.

CARBROMAL.

EPHEDRA, alkaloids of, both natural and synthetic and their salts, except in substances for external use containing less than 1 per cent. of the alkaloids.

ERYTHRITYL TETRANITRATE and other nitric esters of polyhydric alcohols.

GLYCERYL TRINITRATE.

IMIDAZOLE DERIVATIVES with vaso-pressor activity.

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

MERCUROUS CHLORIDE (CALOMEL) in substances for internal use, except when contained in teething powders or preparations for infants.

METHOXYPHENAMINE.

NOR-ADRENALINE, its salts, its N-alkyl derivatives, their salts, in concentrations of more than 0.01 per cent. but not exceeding 1 per cent. of the base.

OCTYL NITRITE.

PAPAVERINE.

PHENAZONE.

SANTONIN.

SODIUM NITRITE for therapeutic use.

FOURTH SCHEDULE.

ACETANILIDE and alkyl acetanilides.

ACETYL METHYL DIMETHYL OXIMIDO PHENYL HYDRAZINE.

ACETAZOLAMIDE.

ADRENALIN, natural or synthetic, its salts, in concentrations of more than 1 per cent. of the base.

ALLYLISOPROPYLACETYLUREA.

AMIDOPYRINE, its salts, its derivatives and their salts.

AMPHETAMINE, its salts, its derivatives and their salts, except when the base is supplied for inhalation absorbed upon an inert solid material.

ANALEPTICS such as Bemegride, Leptazol, Picrotoxin and Niketharnide.

ANTIBIOTICS, Penicillin, Streptomycin, Chloramphenicol, Tetracycline, their derivatives and any other antibiotic substances derived from natural sources.

ANTICHOLINE ESTERASES such as Dyflos, Neostigmine and its salts and other organo-phosphorus compounds with anticholine esterase activity when used for therapeutic purposes.

ANTICONVULSANT SUBSTANCES such as hydantoin derivatives, oxazolidine dione derivatives, and Primidone.

ANTIFOLIC ACID SUBSTANCES such as Aminopterin, Teropterin and Orthopterin.

ANTIHISTAMINES, all tertiary nitrogenous organic bases which possess pharmacological properties characteristic of antihistamine compounds except in preparations labelled and packed for the treatment of motion sickness in packs of 10 doses or less.

ANTIMALARIAL SUBSTANCES such as Amodiaquin, Chloroquine, Mepacrine, Pamaquin, Primaquine, Pyrimethamine, Proguanil and Sonotoquine, their salts (except Quinine and its salts).

ANTIMONY organic compounds for parenteral use.

ANTIPARKINSONIAN SUBSTANCES such as Benzhexol, Caramiphen, Diethazine, Ethopropazine, Procyclidine and their salts.

ANTITHYROID SUBSTANCES such as Carbimazole, Methimazole and Thiouracil and its derivatives except Thiourea.

ANTITUBERCULAR SUBSTANCES such as Isoniazide and its derivatives, para Aminosalicylic acid and its salts and Thiacetazone.

ARSENIC organic compounds for parenteral use.

ATARACTIC SUBSTANCES such as Benactyzine, Azacyclonol, Hydroxyzine and Meprobamate and their salts, their derivatives and their salts.

BARBITURIC ACID, its derivatives and their salts in substances containing more than 0.2 per cent. of barbituric acid or its derivatives and their salts.

BROMOFORM.

BUTYL CHLORAL HYDRATE.

CALCIUM CARBIMIDE.

CANTHARIDES, its alkaloids, their salts and substances containing more than 0.01 per cent. of Cantharidin.

CARBACHOL.

CHINIOFON and other derivatives of 7-iodo-8-hydroxy quinoline and their salts for internal use by humans.

CHLORAL FORMAMIDE.

CHLORAL HYDRATE.

CHLORAZANIL.

CHLORMERODRIN.

CHLORPHENTERMINE.

CHLORPROMAZINE and its salts and other derivatives of phenothiazine and their salts used as ataractics.

CHLORZOXAZONE.

COCAINE, synthetic substitutes for—capable of use for local anaesthesia having a solubility in water of more than 1 per cent. and all preparations except ointments containing 0.5 per cent. or less of such substances.

COCAINE, synthetic substitutes for—capable of use for surface anaesthesia having a solubility in water of 1 per cent. or less and preparations containing more than 2.5 per cent. of such substances except—

(a) Lozenges, pastilles, tablets, capsules containing 30 mg. or less in each;

(b) Suppositories, bougies containing 200 mg. or less in each; and

(c) Preparations for external use containing 10 per cent. or less.

CODEINE, its salts and other ethers of morphine such as ethyl morphine and pholcodine and their salts and substances containing more than 1 per cent of the organic base.

CORTICOTROPHIN and other pituitary hormones for parenteral use in humans.

CORTISONE and steroid suprarenal cortical hormones, either natural or synthetic, or their derivatives.

COUMARIN derivatives and phenylindanedione derivatives used as anticoagulants in the treatment of humans.

CURARE, TUBOCURARINE, d-TUBOCURARINE, d-TUBOCURARINE-DIMETHYL-ETHER, and all synthetic quaternary ammonium compounds having curarising and ganglionic paralyzing effects such as polymethylene bistrimethyl ammonium compounds, Gallamine, Laudexium methyl sulphate, Suxamethonium, Pentolinium, Mecamylamine, Pempidine and Trimetaphan.

CYTOTOXIC SUBSTANCES with blood destroying and/or anti-cancer properties such as Buzulphan, Mustine and Tretamin.

DAPSONE and all derivatives of 4,4′-diaminodiphenylsulphone.

DEXTRAN SULPHATE.

DEXTROMETHORPHAN and its salts except preparations containing 1 per cent. or less of dextromethorphan.

DEXTROPROPOXYPHENE and its salts except preparations containing 1 per cent. or less of dextropropoxyphene.

DEXTRORPHAN and its salts except preparations containing 1 per cent. or less of dextrorphan.

DIBUTAMIDE.

DICYCLOMINE and its salts.

DIETHYLPROPION.

DIGITALIS, its glycosides and their derivatives.

DINITROCRESOLS for therapeutic use.

DINITRONAPHTHOLS for therapeutic use.

DINITROPHENOLS for therapeutic use.

DINITROTHYMOLS for therapeutic use.

DISULFIRAM (except when used for industrial purposes).

EMETINE and its salts, except in Tincture of Ipecacuanha.

ERGOT, its alkaloids, their salts, derivatives of such alkaloids, and their salts.

ETHOHEPTAZINE and substances containing more than 1 per cent. of ethoheptazine.

ETHOXYZOLAMIDE.

ETHYL MORPHINE and substances containing more than 1 per cent. of ethyl morphine.

GLUTETHIMIDE.

HEPARIN.

HYDRALLAZINE.

HYPOTENSIVE SUBSTANCES (such as Apresoline, Trimetaphan, Dihydrallazine, Reserpine, Hexamethonium and Pentamethonium).

IMIPRAMINE.

ION EXCHANGE RESINS, anionic and cationic—for internal use in human beings.

ISOAMINILE.

KHELLIN.

LYSERGIC ACID DIETHYLAMIDE and its derivatives.

MEFENAMIC ACID.

MELANIN STIMULATORS such as Ammoidin, Methoxsalen, 8-Methoxypsoralen, 8-MOP, Meladinine, Meloxine and Xanthotoxin.

MEPHENESIN and its derivatives.

METHANTHELINE, its salts, its derivatives, their salts.

METHAQUALONE.

MERCUROUS CHLORIDE (CALOMEL) when contained in teething powders or preparations for infants.

MERCURY salts and compounds—for parenteral use.

METHYLPENTYNOL and other substituted alkynes for internal use by humans.

METHYL PHENIDATE.

MONO-AMINE OXIDASE INHIBITORS, Ipromiazid, Isocarboxazid, Nialamide, Phenelzine, Pheniprazine and other substances for which monoamine oxidase inhibition is claimed.

MORPHINE ANTAGONISTS such as Nalorphine, Tacrine and Amiphenazole.

MYLICON.

NICOTINYL ALCOHOL.

NOR-ADRENALINE and its salts in concentrations of more than 1 per cent. of the base.

PARALDEHYDE.

PHENACEMIDE.

PHENMETRAZINE.

PHENYLBUTAZONE and its derivatives.

PHENYL-TERTIARY BUTYLAMINE.

PHOLCODINE in substances containing more than 1 per cent. of pholcodine.

PHYSOSTIGMINE.

POTASSIUM PERCHLORATE for therapeutic use.

PROCAINAMIDE.

PROLINTANE.

QUINIDINE.

RAUWOLFIA, its alkaloids, their salts, derivatives of such alkaloids, their salts.

SEX HORMONES, natural or synthetic, their derivatives and their substitutes.

STROPHANTHUS and its glycosides and their derivatives.

SULPHANILAMIDE, its salts, its derivatives, their salts.

SULPHONAL and alkyl sulphonals.

THYROID and its extract, and its active principles.

TOLAZOLINE.

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, antitoxins, and antigens for human parenteral use.

VACCINES—Veterinary live virus.

YOHIMBA, its alkaloids, their salts.

FIFTH SCHEDULE.

Hazardous Substances.

AMMONIA and substances containing 5 per cent. or less by weight of free ammonia (NH_3) except in medicinal substances for internal use, or when used in appliances for inhalation in which the substance is absorbed upon an inert solid material.

CALCIUM, POTASSIUM AND SODIUM HYPOCHLORITES.

COAL TAR SOLVENTS.

DICOPHANE in substances containing 10 per cent. or less of dicophane, except in the case of fertilisers containing less than 2 per cent.

EPOXY RESINS AND POLYESTER RESINS.

FORMALDEHYDE in substances containing 5 per cent. or less of formaldehyde.

GAMMA BENZENE HEXACHLORIDE in substances containing less than 10 per cent. of the gamma isomer, except in the case of fertilisers containing less than 2 per cent.

HYDROCARBONS, LIQUID, distilling under 300° C. when tested according to method D86-61 of the American Society for Testing and Materials and preparations containing more than 25 per cent. of such hydrocarbons when packed in containers of four gallons or less.

HYDROCHLORIC ACID in substances containing 10 per cent. or less by weight of hydrochloric acid (HCl).

KEROSENE and preparations containing more than 25 per cent. of kerosene when packed in containers of four gallons or less.

METALDEHYDE in substances containing 5 per cent. or less of metaldehyde.

4:7 METHANOINDENE and all substitution and/or addition products, such as Chlordane and Heptachlor in preparations containing 2 per cent. or less of the substance.

METHYLATED SPIRIT and substances containing more than 25 per cent. of methylated spirit when packed in containers of four gallons or less.

MINERAL TURPENTINE and preparations containing more than 25 per cent. of mineral turpentine when packed in containers of four gallons or less.

NICOTINE and its salts in preparations containing 1 per cent. or less of the base, except in tobacco in any form.

NITRIC ACID in substances containing 10 per cent. or less by weight of nitric acid.

OIL OF TURPENTINE and preparations containing more than 25 per cent. of oil of turpentine when packed in containers of four gallons or less.

PETROL and preparations containing more than 25 per cent. of petrol when packed in containers of four gallons or less.

POTASSIUM BROMATE in substances containing 0.5 per cent. or less of potassium bromate.

POTASSIUM HYDROXIDE in substances containing 5 per cent. or less of potassium hydroxide.

PYRETHRIN in substances containing 10 per cent. or less of pyrethrin.

SODIUM BROMATE in substances containing 0.5 per cent. or less of sodium bromate.

SODIUM HYDROXIDE in substances containing 5 per cent. or less of sodium hydroxide.

WHITE SPIRIT and preparations containing 25 per cent. or more of white spirit when packed in containers of four gallons or less.

ZINC CHLORIDE in substances containing 5 per cent. or less of zinc chloride.

ZINC SULPHATE in substances containing 5 per cent. or less of zinc sulphate.

ZINC SULPHOCARBOLATE in substances containing 5 per cent. or less of zinc sulphocarbolate.

SIXTH SCHEDULE.

ACETONYL BENZYL—4—HYDROXYCOUMARIN and all substances containing it.

AMMONIA and substances containing more than 5 per cent. by weight of free ammonia (NH_3) except in medicinal substances for internal use, or when used in appliances for inhalation in which the substance is absorbed upon an inert solid material.

ANILINE except substances containing 1 per cent. or less of aniline.

ARNICA and all liquid preparations of arnica.

ARSENIC and preparations containing arsenic when used for agricultural, pastoral or horticultural purposes.

BARIUM salts of (except barium sulphate) and in all substances containing barium salts (except barium sulphate).

BENZENE.

CAMPHORATED OIL.

CARBON BISULPHIDE.

CARBON TETRACHLORIDE except when used for the treatment of humans or in fire extinguishers or in refill containers for such extinguishers.

CHLOROALLYLDIETHYLTHIOCARBAMATE (CDEC)

2-CHLORO-N-N-DIALLYLACETAMIDE (CDAA)

CHROMATES and dichromates of alkali metals.

CHROMIC ACID.

COPPER salts and compounds (inorganic) in substances containing the equivalent of 1 per cent. or more of copper (Cu).

DICHLOROETHYLENE.

DICHLOROETHYL ETHER.

DICHLOROPROPANE.

DICHLOROPROPENE.

DICOPHANE and all substances containing more than 10 per cent. of dicophane.

DIMETHANONAPHTHALENE and all substitution and/or addition products, such as Aldrin and Dieldrin.

DINOCAP (KARATHANE).

DIETHYLENE DIOXIDE
DINITROCRESOLS
DINITROPHENOLS
} and their homologues in substances unless contained in Schedule 7.

DISULFIRAM except for therapeutic use.

ETHER SOLVENT.

ETHYL BROMIDE.

ETHYLENE DIBROMIDE.

ETHYLENE OXIDE.

FERRIC DIMETHYLDITHIOCARBAMATE (FERBAM).

FORMALDEHYDE and substances containing more than 5 per cent. of formaldehyde.

GAMMA BENZENE HEXACHLORIDE and substances containing 10 per cent. or more of the gamma isomer.

HYDROCHLORIC ACID and substances containing more than 10 per cent. by weight of hydrochloric acid (HCl).

HYDROFLUORIC ACID, HYDROSILICOFLUORIC ACID, their salts and other fluorine compounds and all preparations except for therapeutic use and not specifically included in this or any other schedule and except in dentifrices containing less than 0.5 per cent. of fluoride.

IODINE in liquid substances containing 2.5 per cent. or less of iodine.

MERCURY, organic compounds of, when used for agricultural, pastoral or horticultural purposes.

MERCURIC CHLORIDE, when used for agricultural, pastoral or horticultural purposes.

METALDEHYDE and substances containing more than 5 per cent. metaldehyde.

4:7 METHANOINDENE and all substitution and/or addition products, such as Chlordane and Heptachlor and all preparations containing more than 2 per cent. of such substances.

METHYL ALCOHOL except in methylated spirit.

METHYL CHLORIDE.

METHYLENE CHLORIDE.

NICOTINE and its salts in preparations containing more than 1 per cent. of the base, except in tobacco in any form.

NITROBENZENE except in solid or semi-solid polishes, or soaps containing 1 per cent. or less, or other substances containing 0.1 per cent. or less of nitrobenzene.

ORGANO-PHOSPHORUS COMPOUNDS organic fluorophosphates, organic pyrophosphates, organic thiophosphates and any other organo-phosphorus compound used as an insecticide except when included in Schedule 7.

OXALIC ACID and METALLIC OXALATES in polishes.

PENTACHLOROPHENOL except in paints containing 0.25 per cent. or less of pentachlorophenol.

PERMANGANATES.

PICRIC ACID.

PHENOL (CARBOLIC ACID) and its homologues boiling below 220°C. in substances except for therapeutic use containing 3 per cent. or less by weight of phenol or its homologues boiling below 220°C. except paints containing 0.25 per cent or less by weight.

PHENYLENE DIAMINES when used in hair-tinting composition.

PHOSPHIDES, METALLIC.

PHOSPHORUS YELLOW.

POTASSIUM BROMATE and substances containing more than 0.5 per cent. of potassium bromate.

POTASSIUM HYDROXIDE and substances containing more than 5 per cent. of potassium hydroxide (KOH) except in accumulators and batteries.

PYRETHRINS in substances containing more than 10 per cent. of pyrethrins.

SELENIUM in substances other than for human therapeutic use containing 2.5 per cent. or less of selenium.

SODIUM BROMATE and substances containing more than 0.5 per cent. of sodium bromate.

SODIUM CHLORATE in substances containing more than 10 per cent. of sodium chlorate when used for agricultural, pastoral or horticultural purposes or for the extermination of pests and vermin.

SODIUM HYDROXIDE and substances containing more than 5 per cent. of sodium hydroxide (NaOH).

SULPHURIC ACID in substances containing 35 per cent. or less by weight of sulphuric acid (H_2SO_4) except in accumulators, batteries and fire extinguishers.

TETRACHLOROETHYLENE except when used for the treatment of humans and for veterinary purposes.

TETRAMETHYL-THIURAM-DISULPHIDE (THIRAM).

THALLIUM and its salts in preparations containing 0.5 per cent. or less of thallium in containers of not more than 4 ozs.

THIOUREA and its derivatives and in substances containing Thiourea except for therapeutic use.

TOLUENE.

TOLUENE DI-ISOCYANATE.

TOXAPHENE.

TRICHLOROETHYLENE except when specially prepared for medicinal purposes.

TRICHLOROPHENOL.

XYLENE.

ZINC CHLORIDE and substances containing more than 5 per cent. of zinc chloride.

ZINC DIMETHYLDITHIOCARBAMATE (ZIRAM).

ZINC ETHYLENE-BIS-DITHIOCARBAMATE (ZINEB).

ZINC SULPHATE and substances containing more than 5 per cent. of zinc sulphate.

ZINC SULPHOCARBOLATE and substances containing more than 5 per cent. of zinc sulphocarbolate.

SEVENTH SCHEDULE.

Special Poisons

Substances or preparations of exceptional danger which require special precautions and restrictions in manufacture, use and sale.

CARBON TETRACHLORIDE except when used for the treatment of humans or in fire extinguishers or in refill containers for fire extinguishers.

CHLORINE.

CHLOROPICRIN.

DINITROCRESOLS, DINITROPHENOLS and their homologues in substances containing more than 50 per cent. of such substances except for therapeutic use.

FLUOROACETIC ACID, its salts and derivatives and all preparations and admixtures of them.

HYDROCYANIC ACID and substances containing more than 0.15 per cent. of hydrocyanic acid; cyanides and substances containing more than the equivalent of 0.15 per cent. of hydrocyanic acid, except when included in Schedule 1.

METHYL BROMIDE.

ORGANO-PHOSPHORUS COMPOUNDS.

Dimefox (Hanane)
T.E.P.P.

Substances containing more than 20 per cent. of:—

Demeton
Endothion
EPN
Ethion
FAC 20
Guthion
Methyl parathion
Parathion
Phosdrin
Phosphamidon
Schradan
Thimet
Trithion

Substances containing more than 50 per cent. of:—

Delnav (AC 528)
Demeton methyl
Demeton-S-methyl
Dition.

TETRACHLOROETHANE.

THALLIUM and its salts, derivatives, compounds and all preparations and admixtures thereof except as in Schedule 6.

EIGHTH SCHEDULE.

(Drugs of Addiction)

A substance specified in this Schedule includes any active principle, alkaloid, derivative, natural or synthetic, salt, compound and all preparations and admixtures containing any proportion thereof and these are therefore subject to all the restrictions of this Schedule unless specifically exempt.

ACETYLDIHYDROCODEINE (Acetylcodone).

ACETYLMETHADOL (3-acetoxy-6-dimethylamino-4,4-diphenylheptane).

ALLYPRODINE (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).

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).

CANNABIS and CANNABIS RESIN and EXTRACTS and TINCTURES of CANNABIS.

CLONITAZENE (2-para-chlorobenzyl-1-diethylaminoethyl-5-nitrobenzimidazole).

COCA LEAF.

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 containing more than 0.1 per cent. of cocaine.

CODEINE-N-OXIDE.

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

DESOMORPHINE (dihydrodesoxymorphine).

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

DIACETYLNALORPHINE (diacetyl-N-allylnormorphine).

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

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

DIHYDROCODEINE (Paracodine).

DIHYDROMORPHINE.

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

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

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

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

DIPHENOXYLATE 1-(3-cyano-3,3-diphenylpropyl)-4-phenylpiperidine-4-carboxylic acid ethyl ester) excluding preparations containing not more than 2.5 mgm of diphenoxylate and not less than 25 micrograms of atropine (sulphate) per dosage unit.

DIPIPANONE (4, 4-diphenyl-6-piperidine-3-heptanone).

ECGONINE, its ESTERS and DERIVATIVES which are convertible to ECGONINE and COCAINE.

EHYLMETHYLTHIAMBUTENE (3-ethylmethylamino-1, 1-di-(2'-thienyl)-1-butene).

ETONITAZENE (1-diethylaminoethyl-2-para-ethoxybenzyl-5-nitrobenzimidazole).

ETOXERIDINE (1-(2-(2-hydroxyethoxy)ethyl)-4-phenylpiperidine-4-carboxylic acid ethyl ester).

FURETHIDINE (1-(2-tetrahydrofurfuryloxyethyl)-4-carboxylic acid ethyl ester).

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

HEROIN (diacetylmorphine).

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).

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).

METHYLDESORPHINE (6-methyl-delta 6-deoxymorphine).

METHYLDIHYDROMORPHINE (6-methyldihydromorphine).

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 and any solution or dilution in an inert substance whether liquid or solid in any proportion of morphine and all preparations and admixtures (not being such a solution or dilution as aforesaid) containing more than 0.2 per cent. of morphine calculated as anhydrous morphine.

MORPHINE DERIVATIVES (except Codeine, Ethyl morphine and Pholcodine) 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).

NICOMORPHINE (3, 6-dinicotinylmorphine).

NORACYMETHADOL ((±)-alpha-3-acetoxy-6-methylamino-4,4-diphenylheptane).

NORLEVORPHANOL ((-)-3-hydroxymorphinan).

NORMETHADONE (6-dimethylamino-4,4-diphenyl-3-hexanone).

NORMORPHINE (desmethylmorphine).

OPIUM in any form except the alkaloid Papaverine—and in any solution or dilution in an inert substance whether liquid or solid in any preparation of opium and all preparations and admixtures (not being such a solution or dilution as aforesaid) containing more than 0.2 per cent. of morphine calculated as anhydrous morphine.

OXYCODONE (14-hydroxydihydrocodeinone).

OXYMORPHONE (14-hydroxydihydromorphinone).

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).

PHENOMORPHAN (3-hydroxy-N-phenethylmorphinan).

PHENOPERIDINE (1-(3-hydroxy-3-phenylpropyl)-4-phenylpiperidine-4-carboxylic acid ethyl ester).

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

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

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

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

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

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

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

THEBACON (acetyl dihydrocodeinone).

THEBAINE.

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

POISONS ACT, 1964.

Department of Public Health,
Perth, 23rd June, 1965.

HIS Excellency the Governor in Executive Council, acting in pursuance of section 64 of the Poisons Act, 1964, and section 11 of the Interpretation Act, 1918-1962, has been pleased to make the regulations set forth in the Schedule hereunder, to have and take effect on and after the 1st day of July, 1965.

D. J. R. SNOW,
Acting Commissioner of Public Health.

SCHEDULE.

REGULATIONS.

Citation.

1. These regulations may be cited as the Poisons Act Regulations, 1965.

Interpretation.

2. In these regulations unless the context requires otherwise—

"Approved Name" means—

(a) the common name given to any substance by the British Standards Institution or the Australian Standards Association; or

(b) the English name by which any poison or substance is described in the British Pharmacopoeia, the Australian Pharmaceutical Formulary, or the British Pharmaceutical Codex; or

(c) if the substance is not described in the British Pharmacopeia, the Australian Pharmaceutical Formulary, or the British Pharmaceutical Codex, the approved name as published by the General Medical Council of Great Britain; or

(d) if the substance is not described in the British Pharmacopoeia, the Australian Pharmaceutical Formulary, or the British Pharmaceutical Codex, and has not been given an approved name by the General Medical Council of Great Britain, the name given to the substance in any standard book approved by the Commissioner for the purpose of these regulations; or

(e) the systematic chemical name using the English system of nomenclature;

"Child" means a person under the age of twelve years;

"Direction" means regular and frequent supervision but does not necessarily imply continuous personal supervision;

"Dispense" in relation to a medicine or a poison means supplying the medicine or poison on and in accordance with a prescription duly given by a medical practitioner, a dentist or a veterinary surgeon;

"Experienced Person" means a person who for at least five years has been employed in the manufacture, handling or selling of poisons;

"Fourth Schedule drug" means any substance included in the Fourth Schedule to the Act;

"Internal Use" means a substance which is given parenterally, or orally, or a substance which is administered by way of a body orifice for the purpose of absorption and the production of a systemic effect;

"Manufacture" includes the processes of refining, manipulating and mixing any poison or hazardous substance (including such substance in the raw state);

"New Drug" has the same meaning as that term has in and for the purposes of section 37 of the Act;

"Permit" means a permit granted pursuant to the Act;

"Personal Supervision" means close and continuous control requiring the actual presence of the person exercising the supervision;

"Poisons Cupboard" means a substantially made cupboard provided with an effective locking device, and having the word "Poison" conspicuously painted on the outside of the cupboard;

"Qualified Person" means—

(a) a medical practitioner, pharmaceutical chemist, dentist, veterinary surgeon;

(b) a person who is the holder of a degree, approved by the Commissioner, conferred by a University of the British Commonwealth;

(c) a person who is eligible to be—

(i) a Fellow or Associate of the Royal Australian Chemical Institute; or

(ii) a Fellow, Associate or Licentiate of the Royal Institute of Chemistry; or

(d) any other person approved of by the Commissioner;

"Quarter" means any one of the three-monthly periods of any year ending on the 31st March, 30th June, 30th September or 31st December;

"Sale" includes exposing or offering for sale or having in possession for sale, whether by wholesale or retail, and also delivery with or without consideration, in any shop or store or premises appurtenant thereto by the keeper thereof or by his servant or agent; and the verb "to sell" has a corresponding meaning;

"Supply" includes "distribute" and "sell" but the administration to a patient of any substance specified in any of the Schedules to the Act by a medical practitioner or dentist, or by a nurse when acting under the direction of a medical practitioner, or the administration of any substance specified in any of the Schedules to the Act to an animal under the direct personal supervision or under the direction of a veterinary surgeon, shall not be deemed to be supplying within the meaning of these regulations;

"the Act" means the Poisons Act, 1964.

Licences and Permits.

3. A licence to procure, manufacture and supply by wholesale dealing poisons (other than drugs of addiction) shall authorise the licensee to procure, manufacture and supply (according to the endorsement thereon) by wholesale dealing such substances as are specified in the licence, and shall be in the Form No. 1 in Appendix A to these regulations.

4. (1) A licence to procure, manufacture and supply by wholsale dealing drugs of addiction shall authorise the licensee to procure, manufacture, and supply by wholesale dealing drugs of addiction on or from the premises described in the licence, and shall be in the Form No. 2 in Appendix A to these regulations.

(2) In addition to any other conditions required by these regulations the licence shall be subject to the following conditions:—

(a) The manufacture shall be carried out—

(i) by a qualified person whose name appears on the licence; or

(ii) by an experienced person acting under the personal supervision of a qualified person whose name appears on the licence;

(b) the supply shall be carried out—

(i) by a qualified person whose name appears on the licence; or

(ii) by an experienced person whose name appears on the licence;

but should the person whose name appears on the licence cease employment or be unable to exercise the necessary supervision, the Commissioner may authorise, in writing, some other person having the required qualification to act in his stead.

Pharmaceutical Chemist's Licence to Sell Poisons.

5. A pharmaceutical chemist shall not sell or supply any poison except at or from a pharmacy registered under the Pharmacy Act, 1964, and described in the licence issued under these regulations as provided in Form No. 3 in Appendix A to these regulations.

Retailer's Licence to Sell Poisons Specified in the Sixth Schedule to the Act.

6. This licence shall authorise the licensee to procure, and to sell by retail, the poisons specified in the Sixth Schedule to the Act, at the premises described in the licence, and shall be in the Form No. 4 in Appendix A to these regulations.

Retailer's Licence to Sell Poisons Specified in the First, Second or Sixth Schedules to the Act.

7. This licence shall authorise the licensee to procure, and to sell by retail, poisons specified in either the First, Second or Sixth Schedules to the Act at the premises described in the licence, and shall be in the Form No. 5 in Appendix A to these regulations.

Retailer's Licence to Sell Poisons Specified in the Seventh Schedule to the Act.

8. This license shall authorise the licensee to procure, and to sell by retail, poisons specified in the Seventh Schedule to the Act at the premises described in the licence, and shall be in the Form No. 6 in Appendix A to these regulations.

Poisons Permit (Industrial).

9. This permit shall authorise the holder to purchase from a manufacturer or wholesale dealer such poisons as are specified in the permit, which shall be in the Form No. 7 in Appendix A to these regulations.

Poisons Permit (Educational, Advisory or Research).

10. This permit shall authorise the holder to purchase from a manufacturer or wholesale dealer such poisons as are specified in the permit, which shall be in the Form No. 8 in Appendix A to these regulations.

Licence to Hawk, Peddle or Distribute Poisons.

11. This licence issued under section 48 of the Act shall authorise the licensee to sell or distribute in the areas specified in the licence such poisons as are included in the licence, subject to the conditions, limitations and restrictions specified therein, and such licence shall be in the Form No. 9 in Appendix A to these regulations.

Application for Licences or Permits.

12. (1) A person desirous of obtaining a licence or permit pursuant to these regulations shall lodge with the Commissioner an application in or to the effect of such of the Forms Nos. 1A to 11A in Appendix A to these regulations as is appropriate in the particular case, together with the appropriate fee prescribed in Appendix G to these regulations.

(2) Where the applicant for a licence or permit applies on behalf of a corporate body or firm the application shall contain the name of a natural person who, in respect of the premises named in the application, shall be responsible for carrying out the provisions of the Act and these regulations.

Licences and Permits—General Conditions.

13. Every licence or permit issued pursuant to these regulations shall be subject to these regulations and the conditions, limitations and restrictions set out in the licence or permit.

14. Every licence or permit issued pursuant to these regulations shall be valid until the thirtieth of June next following the day of issue, unless sooner cancelled, suspended or revoked, and may thereafter be renewed annually at the discretion of the Commissioner on payment of the prescribed fee (if any).

15. A licence or permit shall not be issued to any person under the age of 21 years, unless he is a qualified person approved by the Commissioner.

16. A sale of any poison shall not be made by any person other than the licensee or a person, not less than 18 years of age, acting on his behalf.

17. A licence or permit is not transferable from one person to another.

Provided that—

(a) a licence or permit held in the name of a person on behalf of a firm or corporate body may, on endorsement by the Commissioner, be transferred into the name of another person on behalf of the firm or corporate body;

(b) the holder of a licence or permit who ceases to carry on or conduct the business or practice to which the licence or permit relates shall within 14 days surrender such licence or permit to the Commissioner.

18. The holder of a licence shall keep such licence displayed in a conspicuous place within the premises specified in the licence.

Containers.

19. (1) Except as provided by these regulations a container in which poison is stored, sold, supplied or transported shall comply with the following conditions:—

(a) The container shall be either a bottle, jar, drum, bag, carton or other type of container approved by the Commissioner, be made of material which is impervious to the poison and of suitable texture and sufficient strength to withstand the ordinary risks of storage, handling or transport without leakage, and be adequately sealed.

(b) The container shall not be a collapsible tube, except where the poison stored or supplied therein is prepared and packed for therapeutic use.

(c) A paper bag shall not be used as the sole container of poison, unless of a type approved by the Commissioner.

(d) (i) A bottle or jar made of glass, having a capacity of 40 fl. ozs. or less, shall have either the words "Poison" or "Not to be taken" embossed or indelibly branded thereon, together with prominent ribs, grooves, points or other distinctive designs, which shall extend over at least one third of the surface, so as to render it distinguishable by touch as a vessel reserved to contain poison.

(ii) A bottle or jar made of glass, having a capacity of more than 40 fl. ozs., shall have either the words "Poison" or "Not to be taken" embossed or indelibly branded thereon in a conspicuous place and in letters not less in depth than 1/32 of the depth of the container.

(iii) A bottle or jar made of plastic shall comply with the Australian Standard Specification for Plastics Containers for Poisonous Substances as published by the Standards Association of Australia.

(e) A container made of metal shall have the word "Poison" embossed or indelibly branded on at least one side and on the top of the container, or on at least one side and on any removable lid, and in each case the word "Poison" shall be in letters not less in depth than 1/32 of the depth of the container, but if the word "Poison" is not embossed it must be left exposed and not covered by a label.

(2) This regulation does not apply in respect of a vessel containing a medicine packed for human internal use.

Labels.

20. Except as provided by these regulations, a person shall not sell any poison or hazardous substance unless the container immediately containing it bears thereon or has securely affixed to it a label bearing the following particulars appropriate to the Schedule to the Act in which the poison or hazardous substance is specified, and any other required particulars:—

First Schedule to the Act.

(a) POISON.
(b) Keep out of reach of children.
(c) The approved name of the poison or poisons and the proportion or percentage of that poison or those poisons in the contents.
(d) Name and address of manufacturer, wholesaler or retailer.
(e) First Aid measures if the substance is included in Appendix C to these regulations.

Second Schedule to the Act.

When not prepared and packed for internal use.

(a) POISON.
(b) Keep out of reach of children.
(c) The approved name of the poison or poisons and the proportion or percentage of that poison or those poisons in the contents.
(d) Name and address of manufacturer, wholsesaler or retailer.
(e) First Aid measures if the substance is included in Appendix C to these regulations.

When prepared and packed for internal use.

(a) CAUTION.
(b) Keep out of reach of children.
(c) The approved name of the poison or poisons and the proportion or percentage of that poison or those poisons in the contents.
(d) Use strictly in accordance with directions.
(e) Name and address of manufacturer, wholesaler or retailer.

Third Schedule to the Act.

(a) CAUTION.
(b) Keep out of reach of children.
(c) The approved name of the poison or poisons and the proportion or percentage of that poison or those poisons in the contents.
(d) Use strictly as directed.
(e) Directions for use.
(f) First Aid measures if the substance is included in Appendix C to these regulations.

Fourth Schedule to the Act.

(a) CAUTION.
(b) Keep out of reach of children.

(c) The approved name of the poison or poisons and the proportion or percentage of that poison or those poisons in the contents.

(d) Use only on prescription.

(e) When supplied by wholesale—manufacturer's or wholesaler's name and address.

When supplied by retail—name and address of retail vendor.

Fifth Schedule to the Act.

(a) CAUTION.

(b) Keep out of reach of children.

(c) The approved name of the hazardous substance or hazardous substances and the proportion or percentage of that hazardous substance or those hazardous substances in the contents.

(d) If swallowed seek medical advice.

(e) First Aid measures if the substance is included in Appendix C to these regulations.

(f) If the substance is included in Appendix D to these regulations, the label shall contain the wording specified therein.

(g) Name and address of manufacturer, wholesaler or retailer.

Sixth Schedule to the Act.

(a) POISON.

(b) Keep out of reach of children.

(c) The approved name of the poison or poisons and the proportion or percentage of that poison or those poisons in the contents.

(d) First Aid measures if the substance is included in Appendix C to these regulations.

(e) If the substance is included in Appendix D to these regulations, the label shall contain the wording specified therein.

(f) Name and address of manufacturer, wholesaler or retailer.

Seventh Schedule to the Act.

Substances in this Schedule shall be labelled according to the requirements set out in Appendix E to these regulations.

Eighth Schedule to the Act.

(a) POISON.

(b) Keep out of reach of children.

(c) The approved name of the poison or poisons and the proportion or percentage of that poison or those poisons in the contents.

(d) Supply without prescription or possession without authority is illegal.

(e) Manufacturer's or supplier's name and address.

21. Notwithstanding the provisions of regulation 20 of these regulations, a medicine containing any poison dispensed or supplied by a pharmaceutical chemist—

(i) for human internal use shall comply with this regulation if it is labelled according to the instructions given on the prescription, together with the identifying number of the prescription and the name and address of the pharmacy at which it is supplied;

(ii) for external therapeutic use shall comply with this regulation if it is labelled with the words "not to be taken" together with the directions, prescription identification number and the name and address of the pharmacy at which it is supplied.

22. Wherever the word "Poison" or "Caution" is required to be shown on a label it shall appear in red on a white background and be surrounded by a red frame. Such word shall form the first line of the principal label and no other word or words shall appear on the same line. The word "Poison" or "Caution" as the case may be shall be in bold face sans serif capital letters of a size not less than half the size of the largest lettering on the label and in any case not less than six points face measurement.

23. A label shall not be attached or affixed to any bottle containing any poison or hazardous substance in such a manner that the embossed points, ridges, flutes, stars. name of the article or the prescribed words blown thereon are covered or obliterated: Provided that the label may cover the front panel of the bottle and extend around the adjacent sides, if the embossed matter on the back panel of the bottle and the name of the article and the prescribed words blown thereon are not covered or obliterated.

24. Any preparation containing any poison or hazardous substance which is supplied for veterinary use, whether in pursuance of a prescription or otherwise, shall be labelled "For veterinary use only" or "For animal treatment only".

Containers and Labels—General.

25. The Commissioner may approve, in writing, a container or label which does not comply with these regulations if, having regard to the nature of the poison and the purpose for which it is to be used, it is unlikely that the interests of safety will be adversely affected by the use of such container or label.

26. The Commissioner may, in the interest of safety, suspend or prohibit the use of any form of container or label for the packing or labelling of any poison.

27. Wherever it is required that the words—

"Keep out of reach of children" or

"First Aid Measures" or

the approved name of the poison or poisons

shall appear on a label, such words or particulars shall be shown—

(a) in bold face sans serif capitals of not less than six points face measurement; and

(b) in such colour or colours as to afford a distinct contrast to the background colour.

Calculation or Percentages.

28. Where required, percentages may be expressed in units other than those in section 51 of the Act.

Storage.

29. Any person having a poison specified in Appendix F to these regulations in or on any premises for the purpose of sale, or to be used in his profession, business, trade or industry shall store that poison in a poisons cupboard securely locked, which cupboard shall be securely fastened to a portion of the premises and not be used for any purpose other than the storage of poison.

Provided that when such poison is stored in bulk quantities it may be stored in a securely locked room, approved by the Commissioner and reserved for the storage of poison.

30. Any person having a hazardous substance or a poison, other than those specified in Appendix F to these regulations, in or on any premises for the purpose of sale or use in his profession, business, trade or industry shall keep that poison or hazardous substance in such a manner as to preclude contamination of any food, drink or condiment by the poison or hazardous substance; and to preclude access to the poison or hazardous substance by children.

Disposal of Poisons.

31. A person shall not dispose of any poison in any place or manner likely to constitute a risk to the public.

Notification of Loss or Theft of Poison.

32. Every person who loses any poison or from whom any poison is stolen shall immediately notify a member of the Police Force of such loss or theft.

Poisons Not to be Sold to Persons Under 16 years.

33. A person who sells or supplies any poison to any person who is apparently under sixteen years of age commits an offence against these regulations, but this regulation does not apply to sales of pharmaceutical preparations or medicines by persons licensed to sell poisons by retail.

Arsenic and Strychnine to be Coloured.

34. (1) It is an offence against these regulations—

(a) to sell any arsenic or any preparation or compound of arsenic which in its natural state is colourless or white in colour unless such arsenic, preparation or compound is mixed with some black substance in a proportion sufficient to render the mixture a gray colour; or

(b) to sell any strychnine or any preparation or compound of strychnine which in its natural state is colourless or white in colour unless such strychnine, preparation or compound is mixed with some red substance in a proportion sufficient to render the mixture a pink colour.

(2) Notwithstanding the provisions of subregulation (1) of this regulation whenever according to the representation of the purchaser the arsenic or strychnine or compound or preparation of arsenic or strychnine is not required for pastoral or agricultural use or for the destruction of vermin, but is required for a purpose for which the colouring matter would render it unfit, such poison may be sold without such admixture.

New Drugs.

35. (1) Before any new drug is first offered for sale to the public the manufacturer, importer or distributor, as the case may require, shall make application to the Commissioner to classify the new drug by determining the Schedule (if any) to the Act in which it is to be included.

(2) An application made under this regulation shall be in Form No. 10 of Appendix A to these regulations.

Supply of Fourth Schedule Drugs.

36. (1) Subject to the Act and these regulations a Fourth Schedule drug shall not be sold or supplied to any person unless—

(a) he is authorised under regulation 40 of these regulations to procure the drug; or

(b) he is the holder of a prescription written by a medical practitioner, dentist or veterinary surgeon, prescribing the drug according to the requirements of these regulations.

(2) A medical practitioner, pharmaceutical chemist, or veterinary surgeon or an assistant under the direct personal supervision of a medical practitioner, pharmaceutical chemist, or veterinary surgeon shall be the only person who shall dispense a Fourth Schedule drug.

(3) The following conditions shall be observed by persons dispensing such prescriptions:—

(a) The prescription shall not be dispensed more than the maximum number of times indicated thereon, and on each occasion upon which it is dispensed the prescription shall be stamped or marked to show clearly the date upon which it is dispensed and the name and address of the pharmacy at which it is dispensed.

(b) The person who dispenses a prescription which does not clearly indicate the maximum number of times it is to be dispensed, or which has reached the last occasion upon which it may be dispensed according to the maximum indicated thereon, shall write in ink, stamp or mark in legible letters across such prescription the word "cancelled".

(c) Before the drug is handed to the purchaser the prescription, whether given in writing or otherwise, shall be copied in full into a Prescription Book, and the entry shall bear an identifying letter or number, show the date upon which the drug is dispensed, and

be signed or initialled by the person who actually dispensed the drug. For the purpose of this paragraph any card system or other system of recording approved by the Commissioner shall be deemed to be the Prescription Book.

In the case of a repeated prescription, an entry in the Prescription Book of the fact of the repeat signed or initialled and dated as required by this paragraph shall be sufficient compliance with this regulation. The label on the bottle or package containing the drug shall be marked with the identifying letter or number of the prescription as appearing in the Prescription Book.

The Prescription Book shall be kept at the place at which the Fourth Schedule drug was dispensed and shall be produced on demand to any person authorised in that behalf under the Act or regulations.

(d) A prescription marked "cancelled" or that is more than six months old shall not be dispensed.

(e) A prescription which is illegible or defaced or which appears to be for the purpose of enabling some unauthorised person to obtain a Fourth Schedule drug, or which does not appear to be genuine, shall not be dispensed.

(f) A pharmaceutical chemist to whom a prescription referred to in paragraph (e) of this subregulation is presented shall retain the prescription and forthwith inform the Commissioner of the relevant circumstances and the reasons for his refusal to dispense the prescription.

Prescriptions.

37. A prescription for the supply of a Fourth Schedule drug shall comply with the following conditions:—

(a) It shall be written in ink in the handwriting of the prescriber.

(b) It shall bear the name, address and signature of the prescriber.

(c) It shall bear the name and address of the patient or, in the case of a prescription for veterinary use, the name and address of the person having the care of the animal for which it is intended.

(d) It shall bear the date on which it is written.

(e) It shall clearly indicate the quantity to be supplied, the maximum number of times it may be repeated and (where applicable) the interval at which it may be repeated.

(f) A prescription written by a dentist shall be for dental purposes only and shall be marked as such and a prescription written by a veterinary surgeon shall be for veterinary use only and shall be marked "For veterinary use only" or "For animal treatment only".

(g) If a prescription contains an unusual dose the prescriber shall indicate that such dose is intended, by underlining that part of the prescription and initialling the same in the margin.

(h) A prescription shall not bear the impression of a rubber stamp or other such contrivance in lieu of the written signature of the medical practitioner, dentist or veterinary surgeon by whom it has been issued.

(i) A prescription shall not be written in cipher.

Dispensing Fourth Schedule Drugs in Emergency Cases.

38. Where a medical practitioner, dentist or veterinary surgeon in a case of emergency orally or by telephone or telegram directs the dispensing of a Fourth Schedule drug, he shall forthwith write a prescription complying with the conditions prescribed in regulation 37 of these regulations, mark such prescription to show that it has been given as a confirmation of instructions given by him orally or by telephone or telegram, and despatch such prescription within 24 hours to the person to whom the instructions were given.

Fourth Schedule Drugs for Veterinary Use.

39. (1) Notwithstanding the provisions of regulation 36 of these regulations, a pharmaceutical chemist is authorised to supply for veterinary use a Fourth Schedule drug listed in Appendix H to these regulations without a

prescription if in the circumstances the purchaser satisfies such pharmaceutical chemist that it is not reasonably practicable for him to obtain such a prescription provided that he records in a book kept for the purpose details showing the name and address of the purchaser and the form and quantity of the drug supplied.

(2) Any preparation containing a Fourth Schedule drug for veterinary use which is registered under the Veterinary Medicines Act and labelled in accordance with the requirements of that Act and the regulations made thereunder may be sold, without prescription, by a pharmaceutical chemist or by a person holding a permit as provided in Form No. 11 in Appendix A to these regulations.

Special Authority to Purchase Fourth Schedule Drugs.

40. Until in any particular case such authority is withdrawn—

(a) a medical practitioner;

(b) a pharmaceutical chemist;

(c) a dentist;

(d) a veterinary surgeon;

(e) an analyst appointed under the Health Act, 1911;

(f) a matron of a hospital registered under the Hospitals Act, 1927;

(g) any other person authorised in writing by the Commissioner

is authorised to procure any Fourth Schedule drug to the extent that it is required for the purpose of his profession or employment, but such authority does not entitle any person to have in his possession any Fourth Schedule drug in quantity greater than is permitted by the Commissioner.

Delivery of a Fourth Schedule Drug on Order.

41. (1) A Fourth Scheduule drug shall not be delivered to any person other than a person authorised by these regulations to purchase the drug or except on the authority of a written order signed by such authorised person, and the person supplying the drug shall satisfy himself that the authority is genuine.

(2) This regulation shall not be deemed to apply to medicines dispensed in pursuance of the foregoing regulations relating to the dispensing of Fourth Schedule drugs.

DRUGS OF ADDICTION.

Authority for Prescribed Persons to Procure and have Drugs of Addiction.

42. (1) Until in any particular case such authority is withdrawn—

(a) a medical practitioner;

(b) a pharmaceutical chemist employed in dispensing medicines at any public hospital or at a pharmacy for which a licence is held under regulation 5 of these regulations;

(c) a dentist;

(d) a veterinary surgeon;

(e) an analyst registered under the Health Act, 1911;

(f) a nurse employed in a public hospital (so far as the possession or use of such drug of addiction is required in connection with its administration to a patient under the instruction of a medical practitioner); and

(g) a person in possession of a permit granted by the Commissioner under these regulations,

is hereby authorised to procure and be in possession of any drug of addiction for the purpose of his profession or employment, subject to the conditions, limitations and restrictions imposed by the Commissioner.

(2) A person to whom a prescription for a drug of addiction has been given is hereby authorised to procure and have possession of the drug of addiction to the extent specified in the prescription.

(3) The authority under this regulation to procure and be in the possession of any drug of addiction does not entitle the holder to procure or have in his possession any drug of addiction in any quantity greater than is permitted by the Commissioner.

Authority for Pharmacists to Retail, Compound and Dispense Drugs of Addiction.

43. (1) Until in any particular case such authority is withdrawn, every pharmaceutical chemist holding a Pharmaceutical Chemist's licence to sell poisons under these regulations is hereby authorised, subject to the conditions, limitations and restrictions imposed by the Commissioner, to procure and to manufacture at his registered premises in the ordinary course of his retail business any preparation, admixture, or extract of any drug of addiction, and to carry on at his registered premises the business of dispensing or compounding any drug of addiction, and also of retailing and supplying a drug of addiction, but only to persons licensed or authorised under these regulations to be in possession of or to procure the drug of addiction.

(2) The authority under this regulation does not in any way entitle the holder to procure, manufacture, sell, distribute, supply, or have in his possession any drug of addiction in any quantity greater than is permitted by the Commissioner.

Register of Drugs of Addiction.

44. (1) Any person authorised to manufacture, distribute, sell or possess any drug of addiction (other than a person having possession by the authority of a prescription from a medical practitioner or veterinary surgeon to the extent shown in the prescription) shall keep or cause to be kept a Register in the form or to the effect of Appendix B to these regulations, and shall enter or cause to be entered in such Register accurate records of the drugs of addiction manufactured, procured, used, supplied or kept by him or on his behalf.

(2) The entries in such Register shall be written in ink on the day of the transaction, and such Register shall be kept on the premises at which the drugs of addiction are kept, manufactured, or disposed of; and where the holders of a licence or other authorised person has drugs of addiction on other premises, he shall keep, or cause to be kept, such Register on those premises also.

(3) All such Registers shall be at all times available for inspection by persons authorised by or under the Act or the regulations to inspect such Registers.

(4) Alterations, obliterations or cancellations shall not be made in any Register, but any mistake made in any entry may be corrected by a marginal or foot note, initialed and dated.

(5) Every person required by these regulations to keep a Register of Drugs of Addiction shall enter in the Register—

(a) the name and quantity of every drug of addiction received, manufactured, sold, used or otherwise disposed of;

(b) the date of each transaction;

(c) the name and address of the person or firm concerned in each transaction;

(d) the balance remaining on hand after each transaction;

(e) in the case of a pharmaceutical chemist the identifying number of the prescription;

(f) in the case of a manufacturer or distributor an identifying number of the order or other authority on which the drug of addiction is supplied.

(6) A separate page of the Register shall be used for each drug of addiction recorded, so that the balance on hand at any time in respect of each drug of addiction will be clearly apparent.

Inventory of Drugs of Addiction.

45. (1) An inventory of drugs of addiction held in stock shall be made—

(a) at intervals of not more than one month by every person required to keep a Register of Drugs of Addiction; and

(b) by a person who is about to relinquish control of drugs of addiction; and

(c) forthwith on assuming control by any person who assumes control of drugs of addiction.

(2) If such inventory of drugs of addiction in stock does not agree with the balance recorded in the Register, the person required to keep the Register shall immediately notify the Commissioner in writing, of the discrepancy.

46. (1) Where a person authorised to have drugs of addiction in his possession for the purpose of his profession or employment does not manufacture, retail, dispense or compound drugs of addiction, or where such dispensing or compounding is done by a medical practitioner, dentist, or veterinary surgeon for the purpose of treatment under his instructions, or his direct personal supervision, it shall be a sufficient compliance with regulation 44 of these regulations if such person keeps a record of—

(a) the drugs of addiction obtained by him and the quantities of each;

(b) the person or firm from whom he obtained such drugs of addiction;

(c) the drugs of addiction disposed of or used by him, the quantities of each, and the date of such disposal or use;

(d) the manner in which such drugs of addiction were disposed of or used; and

(e) the drugs of addiction remaining in his possession and the quantities of each.

(2) The records required to be kept pursuant to this regulation shall be in a book, either written in ink, or gummed or pasted on the pages when the invoices or other documents are used for the purpose, and shall, together with the drugs of addiction then in the possession of the authorised person, be produced for inspection on demand by a person appointed or authorised by or under the Act to inspect those records.

Records to be Retained for Two Years and Available on Demand.

47. (1) All records, registers, prescription books, invoices and other documents relating to drugs of addiction and transactions in regard thereto shall be kept by the person licensed or authorised to have drugs of addiction in his possession for not less than two years from the latest date on which such record, register, prescription book, invoice or other document was made or acted upon.

(2) The records, registers, prescription books, invoices or other documents and stocks of drugs of addiction on hand shall be made available for inspection on demand by a person authorised by or under the Act or regulations or by a member of the Police Force.

(3) In the event of a register being lost or destroyed the person to whom such register belongs shall upon becoming aware of the loss or destruction make and forward to the Commissioner a statutory declaration concerning that loss or destruction and shall immediately take stock of all drugs of addiction in his possession and enter particulars of those stocks in a new register in accordance with the requirements of these regulations.

(4) A person authorised or licensed to procure and be in possession of a drug of addiction, on ceasing to be so authorised or licensed shall, if requested by the Commissioner, surrender any records, registers, prescription books, invoices or other documents and stocks of drugs of addiction that are in his possession to the Commissioner.

Quarterly Returns from Manufacturers and Wholesalers.

48. Every person who holds a licence to manufacture, distribute or sell drugs of addiction by wholesale shall within seven days after the end of each quarter forward to the Commissioner a return in respect of that quarter showing the following details relating to each drug of addiction manufactured, distributed, sold or held in stock, namely—

(a) the stock on hand on the first day of the quarter;

(b) the quantity manufactured or obtained during the quarter;

(c) the name and address of every person to whom supplies were delivered and details of the drugs of addiction supplied during the quarter; and

(d) the balance of stock held at the close of business on the last day of the quarter.

Drugs of Addiction for Use on Ships and Aircraft.

49. (a) Ships.—

(1) The master of a ship is authorised to procure and be in possession of any drug of addiction which is necessary to complete the equipment of the ship in order to comply with the requirements of the Navigation Act.

(2) The holder of a licence or other authorised person may supply such drug on receipt of a written order signed by the master of the ship and endorsed by the manager, or a person authorised in writing by him, of the ship's agents in Western Australia certifying that the drug of addiction is necessary to complete the equipment of the ship in order to comply with the requirements of the Navigation Act.

(b) Aircraft.—

(1) The person in charge of an aircraft is authorised to be in possession of a drug of addiction for the purpose of medical treatment on such aircraft but only in such quantity as does not exceed the quantity required by the Department of Civil Aviation to be carried on the aircraft.

(2) The holder of a licence or other authorised person may supply such drug of addiction on receipt of a written order signed by the manager, or a person authorised in writing by him, of the airline company or firm responsible for the operation of the aircraft in Western Australia.

(c) Any person who supplies a drug of addiction pursuant to this regulation shall within 24 hours of such supply report the details to the Commissioner or the officer in charge of the nearest Police Station.

Drugs of Addiction at Hospitals.

50. (a) Where a Pharmaceutical Chemist is Employed.—The pharmaceutical chemist in charge of the pharmacy department of a hospital shall be responsible for ordering, storing and issuing all drugs of addiction in such hospital and for keeping records of drugs of addiction as required by these regulations.

(b) Where a Pharmaceutical Chemist is not Employed.—The matron of a hospital or other person authorised by the Commissioner shall be responsible for ordering, issuing and storing all drugs of addiction in such hospital and for keeping records of drugs of addiction as required by these regulations.

Prescriptions.

51. A prescription for the supply of a drug of addiction shall comply with the following conditions:—

(a) It shall be written in ink in the handwriting of the prescriber.

(b) It shall bear the name, address and signature of the medical practitioner or veterinary surgeon by whom it is written.

(c) It shall bear the name and full address of the patient or, in the case of a prescription for veterinary use, the name and full address of the person having the care of the animal for which it is intended.

(d) It shall bear the date on which it is written.

(e) It shall clearly indicate the quantity to be supplied, the maximum number of times it may be repeated and (where applicable) the intervals at which it may be repeated. Provided that a prescription written by a veterinary surgeon may be dispensed once only.

(f) A prescription written by a veterinary surgeon shall be for veterinary use only and shall be marked "For veterinary use only" or "For animal treatment only."

(g) If a prescription contains an unusual dose the prescriber shall indicate that such is intended, by underlining that part of the prescription and initialling the same in the margin.

(h) A prescription shall not bear the impression of a rubber stamp or other such contrivance in lieu of the written signature of the medical practitioner or veterinary surgeon by whom it has been issued;

(i) a prescription shall not be written in cipher.

Dispensing Drugs of Addiction.

52. (1) Subject to the Act and these regulations a drug of addiction shall be dispensed or supplied only in accordance with a prescription complying with the requirements of the regulations.

(2) A medical practitioner, pharmaceutical chemist, or veterinary surgeon or an assistant under the direct personal supervision of a medical practitioner, pharmaceutical chemist, or veterinary surgeon shall be the only person who shall dispense a drug of addiction.

(3) The following conditions shall be observed by persons dispensing prescriptions referred to in this regulation:—

(a) The prescription shall not be dispensed more than the maximum number of times indicated thereon, and on each occasion upon which it is dispensed it shall be stamped or marked in ink, by writing or otherwise, to show clearly the date upon which it is dispensed, and the name and address of the pharmacy at which it is dispensed.

(b) The person who dispenses a prescription which does not clearly indicate the maximum number of times such prescription is to be dispensed, or which has reached the last occasion upon which it can be lawfully dispensed according to the maximum indicated thereon, shall write in ink, stamp, or mark in legible letters across such prescription the word "cancelled".

(c) The person who dispenses a prescription shall enter, or cause to be entered, in the book prescribed by regulation 44 of these regulations, a proper record of the transaction which record shall be made in such a way as to be easily understood.

(d) Before the drug of addiction is handed to the purchaser, the prescription, whether given in writing or otherwise, shall be copied in full into a Prescription Book. The entry shall bear an identifying letter or number, and the date upon which the drug of addiction is dispensed, and be signed or initialled by the person who actually dispensed the drug of addiction. For the purpose of these regulations any card system or other system of recording approved by the Commissioner shall be deemed to be the Prescription Book.

(e) In the case of a repeated prescription, an entry in the Prescription Book of the fact of the repeat, signed or initialled and dated as prescribed shall be sufficient compliance with this regulation.

(f) The label on the bottle or package containing the drug of addiction shall be marked with the identifying letter or number of the prescription as appearing in the Prescription Book.

(g) The Prescription Book shall be kept at the place at which the drug of addiction was dispensed and shall be produced on demand to any person authorised in that behalf under the Act or these regulations.

(4) A prescription marked "cancelled" or that is more than six months old shall not be dispensed.

(5) A prescription which is illegible or defaced or which appears to be for the purpose of enabling some unauthorised person to obtain a drug of addiction, or which does not appear to be genuine, shall not be dispensed.

(6) A pharmaceutical chemist to whom a prescription referred to in subregulation 5 of this regulation is presented shall retain the prescription and forthwith inform the Commissioner of the relevant circumstances and the reasons for his refusal to dispense the prescription.

Dispensing Drugs of Addiction in Cases of Emergency.

53. Where a medical practitioner or veterinary surgeon in a case of emergency orally or by telephone or telegram directs the dispensing of a drug of addiction, he shall forthwith write a prescription complying with the conditions prescribed in regulation 51 of these regulations, mark such prescription so as to show clearly that it is in confirmation of the directions given by him orally or by telephone or telegram, and despatch such prescription within 24 hours to the person by whom the drug of addiction was dispensed.

Delivery of Drugs of Addiction on Order.

54. (1) A drug of addiction shall not be delivered to any person except on the authority of a written order signed by the person licensed or otherwise authorised to procure or be in possession of the drug of addiction.

(2) A drug of addiction shall not be delivered to any person not licensed, or otherwise authorised to be in possession of the drug of addiction, who purports to be sent by or on behalf of a person so licensed or authorised, unless the firstmentioned person produced an authority in writing signed by the person so licensed or authorised to receive the drug of addiction on his behalf, and unless the person supplying the drug of addiction is satisfied that the authority is genuine.

(3) This regulation does not apply to medicines dispensed in pursuance of the foregoing regulations.

Common Carrier Protected.

55. A common carrier or his employee is hereby authorised to be in possession of any drug of addiction so far only as the possession is necessary for the transport of the drug of addiction in the ordinary course of business.

Safe Custody of Drugs of Addiction.

56. (1) Any person licensed or authorised to have a drug of addiction in his possession shall store such drug of addiction in a poisons cupboard, securely locked.

(2) This regulation does not apply to—

(a) a person who has been supplied with the drug of addiction pursuant to a prescription from a medical practitioner or veterinary surgeon; or

(b) a medical practitioner or veterinary surgeon when transporting the drug of addiction for the purpose of his profession or practice, if such medical practitioner or veterinary surgeon takes reasonable precautions to protect such drug of addiction against theft or loss.

Labelling.

57. (1) A person shall not supply any drug of addiction unless the package or bottle containing the drug of addiction is plainly labelled or marked to show the quantity of such drug of addiction contained therein.

(2) A person shall not supply any preparation or admixture containing any drug of addiction, unless the package or bottle containing that preparation or admixture is plainly labelled or marked to show the total quantity of such preparation or admixture in the package or bottle and the percentage or quantity of the drug of addiction contained therein, or, in the case of tablets or other articles, the number of such tablets or articles in the package or bottle, and the percentage or quantity of the drug of addiction contained in each tablet or article.

(3) This regulation does not apply to any drug of addiction, preparation, or admixture dispensed in accordance with these regulations.

Improper Prescribing or Use of Drugs of Addiction.

58. (1) A medical practitioner or veterinary surgeon shall not knowingly give a prescription for a drug of addiction merely for purposes of addiction.

(2) A medical practitioner or dentist shall not knowingly supply or administer a drug of addiction merely for purposes of addiction.

Names of Persons from whom Licence or Authority Withdrawn to be Published.

59. The names of all persons from whom a licence or authorisation has been withdrawn shall be published in the *Government Gazette*.

Appeals.

60. (1) Any person desirous of appealing under the provisions of section 29 of the Act shall lodge with the Clerk of Petty Sessions of the Court of Petty Sessions held nearest to the place of business of the appellant notice of appeal in the Form No. 12 in Appendix A to these regulations.

(2) A copy of the notice shall be served on the Commissioner within seven days after lodging the notice with the Clerk of Petty Sessions.

(3) On proof by affidavit to the satisfaction of the magistrate that the notice has been duly served, the magistrate shall cause to be sent to the parties written notice of a date and time for the hearing of the appeal, which date shall not be less than fourteen clear days from the service of the notice.

61. If either party to the appeal neglects to appear personally or by counsel or solicitor on the date and at the time fixed for hearing, the magistrate may—

(a) where the appellant fails to appear, dismiss the appeal;

(b) where the Commissioner fails to appear, hear the appeal or adjourn it to some other date.

62. On the hearing and determination of the appeal the magistrate may make such order as to costs to be paid by either party to the appeal as he may think just.

63. On the hearing of the appeal, it shall proceed according to the procedure and rules of evidence applicable in the Court of Petty Sessions.

APPENDIX A

Form No. 1.

Poisons Act, 1964.

LICENCE TO PROCURE, MANUFACTURE AND SUPPLY POISONS (OTHER THAN DRUGS OF ADDICTION) BY WHOLESALE DEALING.

This licence is granted to.....and authorises him to procure, manufacture and supply by wholesale dealing on behalf of.....the poisons specified in the *1st, 2nd, 3rd, 4th, 6th, 7th Schedules to the Poisons Act.

Subject to the following conditions:—

1. The poisons will be manufactured at premises situated at.....
.....

(a) under the personal supervision of.....
who holds the qualification.....(or)

(b) under the direction of.....who
holds the qualification.....and
under the personal supervision of.....
who is an experienced person within the meaning of the regulations
(or)

(c) .....
.....

2. The poisons will be supplied from premises situated at.....
.....

(a) under the personal supervision of.....
who holds the qualification.....(or)

(b) under the direction of.....who holds the qualification.....and under the personal supervision of.....who is an experienced person within the meaning of the regulations (or)

(c) .....
.....

3. (a) .....

(b) .....

Dated at Perth.....19.....

Valid until 30th June, 19 .....

.....
Commissioner of Public Health.

* Strike out whichever is not applicable.

Form No. 1A.

Poisons Act, 1964.

APPLICATION FOR LICENCE TO PROCURE, MANUFACTURE AND SUPPLY POISONS (OTHER THAN DRUGS OF ADDICTION) BY WHOLESALE DEALING.

To the Commissioner of Public Health,
Public Health Department,
57 Murray Street,
Perth.

I, Mr.
Mrs.....
Miss (Full Name)

hereby apply for a licence to procure, manufacture and supply by wholesale dealing on behalf of.....the poisons specified in the *1st, 2nd, 3rd, 4th, 6th, 7th Schedules to the Poisons Act.

In support of this application I declare that—

1. The poisons will be manufactured at premises situated at.....
.....

(a) under the personal supervision of.....
who holds the qualification.....(or)

(b) under the direction of.....who holds the qualification.....and under the personal supervision of.....
who is an experienced person within the meaning of the regulations (or)

(c) .....
.....

2. The poisons will be supplied from premises situated at.....
.....

(a) under the personal supervision of.....
who holds the qualification.....(or)

(b) under the direction of.....who holds the qualification.....and under the personal supervision of.....who is an experienced person within the meaning of the regulations (or)

(c) .....
.....

Date.....

Signature of Applicant.
.....

* Strike out whichever is not applicable.

Form No. 2.

Poisons Act, 1964.

LICENCE TO PROCURE, MANUFACTURE AND SUPPLY BY WHOLESALE DEALING DRUGS OF ADDICTION.

This licence is granted to.....and authorises him to procure, manufacture and supply by wholesale dealing on behalf of .....the following drugs of addiction.....
.....

Subject to the following conditions:—

1. The drugs of addiction will be manufactured at premises situated at .....
   (a) under the personal supervision of..... who holds the qualification.....(or)
   (b) under the direction of.....who holds the qualification.....and under the personal supervision of.....who is an experienced person within the meaning of the regulations.
2. The drugs of addiction will be supplied from premises situated at..... .....
   (a) under the personal supervision of..... who holds the qualification.....(or)
   (b) under the direction of.....who holds the qualification.....and under the personal supervision of.....who is an experienced person within the meaning of the regulations.
3. (a).....
   (b).....

Dated at Perth.....19.....

Valid until 30th June, 19.....

.....
Commissioner of Public Health.

Form No. 2A.

Poisons Act, 1964.

APPLICATION FOR LICENCE TO PROCURE, MANUFACTURE AND SUPPLY BY WHOLESALE DEALING DRUGS OF ADDICTION.

To the Commissioner of Public Health,
Public Health Department,
57 Murray Street, Perth.

I, Mr. / Mrs. / Miss (Full Name) .....hereby apply for a licence to procure, manufacture and supply by wholesale dealing the following drugs of addiction.....
.....

In support of this application I declare that:—

1. The drugs of addiction will be manufactured at premises situated at..... .....
   (a) under the personal supervision of..... who holds the qualification.....(or)
   (b) under the direction of.....who holds the qualification.....and under the personal supervision of.....who is an experienced person within the meaning of the regulations.
2. The drugs of addiction will be supplied from premises situated at..... .....
   (a) under the personal supervision of..... who holds the qualification.....(or)
   (b) under the direction of.....who holds the qualification.....and under the personal supervision of.....who is an experienced person within the meaning of the regulations.

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

Form No. 3.

Poisons Act, 1964.

PHARMACEUTICAL CHEMIST'S LICENCE TO SELL POISONS.

This licence is granted to.....
and authorises him to sell poisons at premises situated at.....
.....

Dated at Perth.....19.....
Valid until 30th June, 19.....

.....
Commissioner of Public Health.

Form No. 3A.

Poisons Act, 1964.

APPLICATION FOR PHARMACEUTICAL CHEMIST'S LICENCE TO SELL POISONS.

To the Commissioner of Public Health,
Public Health Department,
57 Murray Street,
Perth.

I, Mr. / Mrs. / Miss .....
(Full Name)

a pharmaceutical chemist registered to practice in Western Australia, hereby apply for a licence to sell poisons at premises situated at.....
.....which premises are registered as a Pharmacy under the Pharmacy Act, 1964, the Registration Certificate in respect of which is No.....valid until 30th June, 19.....

.....
Signature of Applicant.

Date.....

Form No. 4.

Poisons Act, 1964.

LICENCE TO SELL BY RETAIL POISONS SPECIFIED IN THE 6TH SCHEDULE.

This licence is granted to.....
and authorises him to procure, and to sell by retail, on behalf of.....
.....the poisons specified in the 6th Schedule to the Poisons Act, 1964, at premises situated at.....
.....

Dated at Perth.....19.....
Valid until 30th June, 19.....

.....
Commissioner of Public Health.

Form No. 4A.

Poisons Act, 1964.

APPLICATION FOR LICENCE TO SELL BY RETAIL POISONS SPECIFIED IN THE 6TH SCHEDULE.

To the Commissioner of Public Health,
Public Health Department,
57 Murray Street,
Perth.

I, Mr. / Mrs. / Miss ..... (Full Name)
hereby apply for a licence to sell, by retail, on behalf of.....
.....the poisons specified in the 6th Schedule to the Poisons Act, 1964.

I declare that—

(a) I have attained the age of 21 years.
(b) The poisons will be sold only at premises siutated at.....
.....
(c) The poisons will not be sold by an assistant under 18 years of age.
(d) The poisons will not be sold to anyone who is apparently under 16 years of age.

Date.....

.....
Signature of Applicant.

Form No. 5.

Poisons Act, 1964.

LICENCE TO SELL BY RETAIL POISONS SPECIFIED IN THE 1ST, 2ND OR 6TH SCHEDULES.

This licence is granted to.....
and authorises him to procure, and to sell by retail, on behalf of.....
.....the poisons specified in the 1st, 2nd or 6th Schedules to the Poisons Act, 1964, at premises situated at.....
.....

Dated at Perth.....19.....

Valid until 30th June, 19.....

.....
Commissioner of Public Health.

Form No. 5A.

Poisons Act, 1964.

APPLICATION FOR LICENCE TO SELL BY RETAIL POISONS SPECIFIED IN THE 1ST, 2ND OR 6TH SCHEDULES.

To the Commissioner of Public Health,
Public Health Department,
57 Murray Street,
Perth.

I, Mr. / Mrs. / Miss ..... (Full Name)
hereby apply for a licence to sell, by retail, on behalf of.....
.....the poisons specified in the 1st, 2nd or 6th Schedules to the Poisons Act, 1964, at premises situated at.....
.....

I declare that—

(a) I have attained the age of 21 years.
(b) These premises are distant at least five miles from the nearest place at which a pharmaceutical chemist conducts a pharmacy.
(c) The poisons will not be sold by an assistant under 18 years of age.
(d) The poisons will not be sold to anyone who is apparently under 16 years of age.

Date.....

.....
Signature of Applicant.

Form No. 6.

Poisons Act, 1964.

LICENCE TO SELL BY RETAIL POISONS SPECIFIED IN THE 7TH SCHEDULE.

This licence is granted to.....
and authorises him to procure, and to sell by retail, on behalf of.....
....., at premises situated at
.....the following poisons specified in the 7th Schedule:—

.....

.....

Subject to the following conditions:—

.....

.....

Dated at Perth.....19.....
Valid until 30th June, 19.....

.....
Commissioner of Public Health.

Form No. 6A.

Poisons Act, 1964.

APPLICATION FOR LICENCE TO SELL BY RETAIL POISONS SPECIFIED IN THE 7TH SCHEDULE.

To the Commissioner of Public Health,
Public Health Department,
57 Murray Street,
Perth.

I, Mr. / Mrs. / Miss.....
(Full Name)
hereby apply for a licence to sell, by retail, on behalf of.....
.....poisons specified in the 7th Schedule:—

.....

.....

.....

I declare that—

(a) I have attained the age of 21 years.
(b) The poisons will be sold only at premises situated at.....
.....
(c) The poisons will be sold only by myself or by an assistant who is not less than 18 years of age.
(d) The poisons will not be sold to anyone who is apparently under 16 years of age.
(e) .....

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

Form No. 7.

Poisons Act, 1964.

POISONS PERMIT (INDUSTRIAL).

This permit is granted to.....and authorises him to purchase on behalf of..... from a manufacturer or wholesale dealer—

(a) the poisons specified in the..... Schedules to the Poisons Act, 1964;

(b) the following poisons:—

.....

.....

.....

This permit is issued subject to the following conditions:—

(1) the poisons will be stored only at premises situated at .....;

(2) the poisons will not be resold;

(3) the poisons will be used only for the following purposes:—

.....

.....

.....

(4) .....

.....

.....

Dated at Perth....., 19.....

Valid until 30th June, 19.....

.....

Commissioner of Public Health.

Form No. 7A.

Poisons Act, 1964.

APPLICATION FOR POISONS PERMIT (INDUSTRIAL).

To the Commissioner of Public Health,
Public Health Department,
57 Murray Street,
Perth.

I, Mr. / Mrs / Miss .....
(Full Name)

hereby apply on behalf of.....for a permit to purchase from a manufacturer or wholesale dealer—

(a) the poisons specified in the *1st, 2nd, 3rd, 4th, 6th, 7th, 8th Schedules to the Poisons Act, 1964; or

(b) the following poisons:—

.....

.....

.....

In support of this application I declare that—

(1) the poisons will be stored only at premises situated at.....

.....

(2) the poisons will not be resold;

(3) the poisons will be used only for the following purposes:—

.....

.....

.....

(4) .....

.....

.....

Date.....

.....

Signature of Applicant.

*Strike out whichever does not apply.

Form No. 8.

Poisons Act, 1964.

POISONS PERMIT (EDUCATIONAL, ADVISORY OR RESEARCH).

This permit is granted to.....and authorises him to purchase on behalf of..... from a manufacturer or wholesale dealer—

(a) the poisons specified in the..... Schedules to the Poisons Act, 1964.

(b) the following poisons:—

.....

.....

.....

This permit is issued subject to the following conditions:—

(1) the poisons will be stored only at premises situated at.....

.....

(2) the poisons will not be resold;

(3) the poisons will be used only for the following purposes:—

.....

.....

.....

(4) .....

.....

.....

Dated at Perth....., 19.....

Valid until 30th June, 19.....

.....

Commissioner of Public Health.

Form No. 8A.

Poisons Act, 1964.

APPLICATION FOR POISONS PERMIT.
(EDUCATIONAL, ADVISORY OR RESEARCH.)

To the Commissioner of Public Health,
Public Health Department,
57 Murray Street,
Perth.

I, Mr. / Mrs. / Miss ..... (Full Name)

hereby apply on behalf of.....for a permit to purchase from a manufacturer or wholesale dealer—

(a) the poisons specified in the *1st, 2nd, 3rd, 4th, 6th, 7th, 8th Schedules to the Poisons Act, 1964; or

(b) the following poisons:—

.....

.....

.....

In support of this application I declare that—

(1) the poisons will be stored only at premises situated at.....

.....

(2) the poisons will not be resold;

(3) the poisons will be used only for the following purposes:—

.....

.....

.....

(4) .....

.....

.....

Date.....

.....

Signature of Applicant.

*Strike out whichever does not apply.

Form No. 9.

Poisons Act, 1964.

LICENSE TO HAWK, PEDDLE OR DISTRIBUTE POISONS.

This licence is granted to .....
and authorises him to sell by hawking or peddling, or to distribute as a sample, the following poisons:—

.....
.....
.....

This licence is issued under section 48 of the Poisons Act, 1964, and is subject to the following conditions, limitations and restrictions:—

.....
.....
.....

Dated at Perth.....19.....

Valid until.....19.....

.....
Commissioner of Public Health.

Form No. 9A.

Poisons Act, 1964.

APPLICATION FOR LICENCE TO HAWK, PEDDLE OR DISTRIBUTE POISONS.

To the Commissioner of Public Health,
Public Health Department,
57 Murray Street,
Perth.

I, Mr. / Mrs. / Miss ..... (Full Name)

of ..... (address) hereby apply for a licence to hawk, peddle, or distribute as a sample, the following poisons:—

.....
.....
.....

In support of this application I declare that—

(a) I have attained the age of 21 years;

(b) the poisons will not be sold by an assistant under 18 years of age;

(c) the poisons will not be sold to anyone who is apparently under 16 years of age;

(d) the poisons will be sold or distributed only in the areas specified in the licence;

(e) .....
.....

Date.....19.....

.....
Signature of Applicant.

Form No. 10.

Poisons Act, 1964.

APPLICATION FOR CLASSIFICATION OF A NEW DRUG.

To the Commissioner of Public Health,
Public Health Department,
57 Murray Street,
Perth.

I (or we) ..... of ....., herewith make application for classification of the new drug.....
.....

I (or we) request that this drug be—

(a) included in Schedule.....;

(b) exempted from inclusion in any Schedule;

(c) preparations containing not more than ..... per cent. of the drug be .....

In support of this application I (we) submit the following information:—

1. The (a) approved name of the drug.....
(b) generic name of the drug .....

2. The trade name (or names) .....

3. The proprietary name (or names) .....

4. The chemical name .....

5. The chemical nature .....

6. The chemical structure and formula .....

7. Its description in precise chemical terms, together with its physical details .....

8. The nature and limits of any impurities present.....

9. Particulars of the tests and standards applied to control its potency, purity and safety during manufacture and storage .....

10. Full details of investigations made with respect to the safety and efficacy of the drug, including tests carried out by universities and/or research institutions, and clinical trials.

Note.—Full reports are required of adequate tests which will show whether or not the substance will be safe. The reports shall include detailed data derived from appropriate animal and other biological experiments in which the methods used and the results obtained are clearly set forth. Details of any reports which could bias an evaluation of the safety of the substance shall NOT be omitted. Special attention shall be given to—

(a) acute, sub-acute and chronic toxicity;

(b) uniformity of response within a species and among different species;

(c) occurrence of unusual or alarming reactions, such as carcinogenesis;

(d) known side effects;

(e) occurrence of sensitivity tolerance or idiosyncracy in response to the substance;

(f) Metabolism, rate, extent and mode of elimination of the substance;

(g) any tendency towards accumulation in the body;

(h) any special incompatability;

(i) method of assay.

11. A statement of the amounts of all ingredients, route of administration, proposed dosage, the claims to be made for such drug and a description of the pharmacteutical forms in which it is proposed to be sold.

13. Full details of proposed labelling and packaging.

14. Evidence of approval or rejection by any other statutory body or authority.

15. Complete bibliography of any publications relating to pharmacological and therapeutic actions, including clinical trials.

Form No. 11.

Poisons Act, 1964.

PERMIT TO SUPPLY FOR VETERINARY USE THE PREPARATIONS REFERRED TO IN REGULATION 39 (2).

This permit is granted to.....and authorises him to sell on behalf of..... at premises situated at..... the preparations referred to in regulation 39 (2).

Dated at Perth.....19.....

Valid until 30th June, 19.....

.....

Commissioner of Public Health.

Form No. 11A.

Poisons Act, 1964.

APPLICATION FOR PERMIT TO SUPPLY FOR VETERINARY USE THE PREPARATIONS REFERRED TO IN REGULATION 39 (2).

To the Commissioner of Public Health,

Valid until 30th June, 19.....

Public Health Department,
57 Murray Street,
Perth.

I, Mr. / Mrs / Miss.....
(Full Name)

hereby apply for a permit to sell on behalf of..... at premises situated at.....the preparations referred to in regulation 39 (2) of the Poisons Act Regulations, 1965.

Date.....19.....

.....

Signature of Applicant.

Form No. 12.

Poisons Act, 1964.

NOTICE OF APPEAL UNDER SECTION

IN the Court of Petty Sessions
at.....

BETWEEN

.....Appellant

and

.....Respondent

TAKE NOTICE that pursuant to the provisions of Section of the Poisons Act, 1964, I intend to appeal to the Magistrate of the abovenamed Court against your (a).....on the.....day of.....19..... (b) .....
.....
.....

Dated this.....day of.....19.....

.....

Appellant.

To the Commissioner of Public Health
And to.....

(a) State whether refusal, cancellation, order, etc., of the Commissioner.
(b) Set out particulars of the decision of the Commissioner.

APPENDIX B.

DRUG REGISTER OF DRUGS OF ADDICTION USED AND RECEIVED.

Date on which Transaction was Effected	Person, Body or Firm to Whom Sold or Supplied, or from Whom Received		Amount Used Sold or Supplied	Balance	Prescription Number	Prescriber	Dispenser
	Name and Address	Amount Received					

APPENDIX C.

POISONS AND HAZARDOUS SUBSTANCES REQUIRED TO BE LABELLED WITH FIRST AID MEASURES.

Except where otherwise stated, a poison or hazardous substance in this Appendix includes any derivative, compound, preparation or admixture included in relation to that item in the Schedule referred to.

First Schedule.—

Aconite, Antimony, Arsenic, Atropine, Belladonna, Brucine, Colchicine, Hydrocyanic acid, Hyoscine, Mercuric chloride, Mercuric iodide, Mercuric nitrate, Mercuric-potassium iodide, Phosphorus (yellow), Strychnine.

Second Schedule.—

Acid Acetic glacial, Chloroform, Creosote, Cresol, Diamines, Iodine, Lead salts, Nitric acid, Nitrophenols (ortho, meta and para), Oxalic acid, Phenol (Carbolic acid), Selenium, Sulphuric acid.

Third Schedule.—

Santonin, Sodium nitrite.

Fifth Schedule.—

Kerosene.

Sixth Schedule.—

Acetonyl benzyl-4-hydroxycoumarin (Warfarin), Ammonia, Aniline, Arsenic, Barium salts, Benzene, Carbon bisulphide, Carbon tetrachloride, Chloroallyldiethylthiocarbamate (CDEC), 2-Chloro-N-N-diallylacetamide (C.D.A.A.), Copper salts, Dicophane (D.D.T.), Dimethanonaphthalene, Dinocap (Karathane), Dinitrocresols, Dinitrophenols, Disulfiram, gamma Benzene hexachloride, Hydrochloric acid, Mercury (organic compounds), Mercuric chloride, Metaldehyde, 4:7 Methanoindene, Nicotine, Organo-phosphorus Compounds, Pentachlorophenol, Phenol (Carbolic acid), Phosphides (metallic), Phosphorus (yellow), Potassium bromate, Potassium hydroxide, Sodium bromate, Sodium hydroxide, Sulphuric acid in substances containing more than 10 per cent. by weight of sulphuric acid (except in accumulators, batteries or fire extinguishers), Tetramethyl-thiuram-disulphide (Thiram), Thallium, Toxaphene, Zinc chloride, Zinc dimethyldithiocarbamate (Ziram), Zinc ethylene-bis-(Dithiocarbamate) (Zineb).

Seventh Schedule.—

Carbon tetrachloride, Chlorine, Chloropicrin, Dinitrocresols, Dinitrophenols, Fluoroacetic acid, Hydrocyanic acid, Methyl bromide, Organo-phosphorus compounds, Tetrachloroethane, Thallium.

APPENDIX D.

POISONS AND HAZARDOUS SUBSTANCES REQUIRED TO BE LABELLED WITH A WARNING STATEMENT.

(a) "Avoid contact with the skin".

Formaldehyde, oxalic acid and metallic oxalates, phenol, zinc chloride, hydrochloric acid, hydrofluoric acid, sodium chlorate, sulphuric acid.

(b) "Avoid contact with the skin and avoid breathing its dust (or vapour)".

Dichloroethylene, dicophane, dimethanonaphthalene, gamma benzene hexachloride, 4 : 7-methanoindene, phosphides (metallic), aniline, benzene, carbon bisulphide, carbon tetrachloride, chloropicrin, chromates, chromic acid, dichlorethyl ether, diethylene dioxide, dinitrocresols, dinitronaphthols, ether solvent, ethyl bromide, ethylene dibromide, ethylene oxide, methyl alcohol, methyl bromide, methyl chloride, methylene chloride, nicotine, nitrobenzene, organo-phosphorus compounds, pentachlorophenol, selenium, tetrachloroethylene, toluene, toxaphene, trichloroethylene, trichlorophenol, xylene.

(c) "Warning—this substance is inflammable".

Dichloroethylene, kerosene, methylated spirits, mineral turpentine, oil of turpentine, petrol, solvents and white spirits, benzene, carbon bisulphide, diethylene dioxide, ether solvent, ethylene oxide, methyl alcohol, toluene, xylene.

(d) "Warning—this substance may cause serious inflammation of the skin in certain persons and should be used only in accordance with expert advice".
Diamines.

(e) "Warning—this substance is caustic—avoid contact with the skin".
Potassium hydroxide, sodium hydroxide.

APPENDIX E.

LABELS FOR POISONS IN THE SEVENTH SCHEDULE.

Except where otherwise stated a substance in this Appendix includes any derivative, compound, preparation or admixture included in relation to that substance in the Seventh Schedule to the Act.

The label for any poison in the Seventh Schedule shall contain the following particulars:—

(a) Poison;
(b) Keep out of reach of children;
(c) Do not open until you have read safety directions;
(d) Approved name of the poison;
(e) The proportion or percentage of poison in the contents;
(f) Name and address of manufacturer, wholesaler or retailer;
(g) The following information appropriate to the particular item:—

CARBON TETRACHLORIDE.

Danger! Hazardous vapour and liquid.
May be fatal if inhaled or swallowed.

Precautions. Avoid breathing vapour by using adequate ventilation or a suitable respirator. Avoid contact with skin or clothing. Do not smoke or use naked lights. Store in a cool place.

FIRST AID MEASURES.

If inhaled: Remove patient from further exposure.
If splashing occurs: Remove contaminated clothing.
If splashing occurs on the skin: Wash off skin immediately.
If swallowed—only when patient conscious: Give milk drink (or water, if no milk). Give a cup of water with two teaspoonfuls of salt (if available). To assist in inducing vomiting tickle back of throat with finger. Give a tablespoonful of Epsom or Glauber Salts.
Call a doctor.

CHLORINE.

Irritant to the skin, extremely dangerous if inhaled.

Precautions: Avoid inhalation or contact with the skin. A suitable respirator should be available and used when required.

FIRST AID MEASURES.

Put on suitable respirator, then remove patient from further exposure.
If patient not breathing commence artificial respiration.
Call a doctor.

CHLOROPICRIN.

Extremely dangerous if inhaled, even in small doses (was used as a war gas). Very irritant to the skin.

Precautions. Avoid breathing fumes or contact with skin, eyes or clothing. A suitable respirator should be available and used as required.

FIRST AID MEASURES.—

Put on suitable respirator, then remove patient from further exposure.
If splashing occurs: Remove contaminated clothing.
If splashing occurs on the skin: Wash off skin immediately.
If splashing occurs in the eyes: Flush eyes with water for 5 minutes.
Call a doctor.

DINITROCRESOLS. DINITROPHENOLS.

Extremely dangerous and may be fatal if swallowed, inhaled or absorbed through the skin.

Precautions. Wear complete cover waterproof clothing including suitable respirator. Wash hands and face thoroughly before eating or smoking. Wash clothing and gloves daily.

FIRST AID MEASURES.—

If splashing occurs: Remove contaminated clothing.

If splashing occurs on the skin: Wash off skin immediately.

If splashing occurs in the eyes: Flush eyes with water for 5 minutes.

If swallowed—only when patient conscious: Give milk drink (or water if no milk). Give a cup of water with 2 teaspoonsful of salt (if available). To assist in inducing vomiting tickle back of throat with finger. Give a tablespoonful of Epsom or Glauber Salts.

Call a doctor.

FLUOROACETIC ACID.

Extremely poisonous if taken internally or inhaled.

Precautions. Avoid inhaling. Wash thoroughly before eating or smoking.

FIRST AID MEASURES.—

If swallowed—only when patient conscious: Give milk drink (or water if no milk). Give a cup of water with 2 teaspoonfuls of salt (if available). To assist in inducing vomiting tickle back of throat with finger. Give a tablespoonful of Epsom or Glauber Salts.

Call a doctor.

HYDROCYANIC ACID AND ALL CYANIDES.

Extremely poisonous when taken by mouth, inhaled or absorbed through the skin.

For solid and liquid preparations the label should also include:—

This preparation liberates poisonous gas on contact with water or acids.

Precautions. Store in a cool place. Avoid contact with skin. Wash hands before eating or smoking. When exposed to gas a suitable respirator must be worn.

FIRST AID MEASURES.—

If inhaled: Put on suitable respirator, then remove patient from further exposure.

If splashing occurs: Remove contaminated clothing.

If splashing occurs on the skin: Wash off skin immediately.

If swallowed—only when patient conscious: Give milk drink (or water if no milk). Give a cup of water with 2 teaspoonfuls of salt (if available). To assist in inducing vomiting tickle back of throat with finger. Give a tablespoonful of Epsom or Glauber Salts.

If patient unconscious: Do not give anything by mouth. Do not make patient vomit. If not breathing commence artificial respiration.

Call a doctor.

Use Amyl Nitrite Capsule in all cases. Break and hold lightly under nose for 15 seconds. If patient unconscious—repeat this up to 5 times at 15 second intervals.

METHYL BROMIDE.

Vapour extremely hazardous. Highly volatile and causes burns.

Precautions. Store in cool, well ventilated place. Do not breathe vapour. Avoid contact with skin, eyes or clothing. A suitable respirator should be available and used as required.

FIRST AID MEASURES.—

Put on suitable respirator, then remove patient from further exposure.

If splashing occurs. Remove contaminated clothing.

If splashing occurs on the skin: Wash off skin immediately.

If splashing occurs in the eyes: Flush eyes with water for 5 minutes.

For convulsions: Protect patient from injury. If breathing much obstructed, pull chin forward. Call a doctor.

Call a doctor.

Keep at rest.

ORGANO-PHOSPHORUS COMPOUNDS.

Extremely dangerous if swallowed, inhaled or absorbed through the skin.

Precautions. Wear complete cover protective waterproof clothing, including suitable respirator. Wash hands and face before eating or smoking. Wash clothing and gloves daily. Wash out and destroy empty containers.

FIRST AID MEASURES.—

If splashing occurs: Remove contaminated clothing.

If splashing occurs on the skin: Wash off skin immediately.

If splashing occurs in the eyes: Flush eyes with water for 5 minutes.

If swallowed—only when patient conscious:

Give milk drink (or water if no milk). Give a cup of water with 2 teaspoonfuls of salt (if available). To assist in inducing vomiting tickle back of throat with finger. Give a tablespoonful of Epsom or Glauber Salts.

If patient unconscious: Do not give anything by mouth. Do not make patient vomit. If not breathing commence artificial respiration.

For convulsions: Protect patient from injury. If breathing much obstructed, pull chin forward. Call a doctor.

Call a doctor.

TETRACHLORETHANE.

Danger! Vapour extremely hazardous.

Precautions. Avoid breathing vapour, or contact with skin, eyes or clothing.

FIRST AID MEASURES.—

If inhaled: Remove patient from further exposure.

If splashing occurs: Remove contaminated clothing.

If splashing occurs on the skin: Wash off skin immediately.

If swallowed—only when patient conscious: Give milk drink or water if no milk). Give a cup of water with 2 teaspoonfuls of salt (if available). To assist in inducing vomiting tickle back of throat with finger. Give a tablespoonful of Epsom or Glauber Salts.

Call a doctor.

THALIUM.

Extremely poisonous if taken by mouth or absorbed through the skin.

Precautions: Avoid contact with skin.

FIRST AID MEASURES.—

If swallowed—only when patient conscious: Give milk drink (or water if no milk). Give a cup of water with 2 teaspoonfuls of salt (if available). To assist in inducing vomiting tickle back of throat with finger. Give a tablespoonful of Epsom or Glauber Salts.

Call a doctor.

APPENDIX F.

POISONS WHICH ARE REQUIRED TO BE STORED IN A POISONS CUPBOARD.

Any substance included in the Eighth Schedule to the Act (Drugs of Addiction), Arsenic (white), Carbon bisulphide, Chloroform. Fluoroacetic acid, Hydrocyanic acid and cyanides, Nitrophenols, Organo-phosphorus compounds (dimefox, T.E.P.P., and substances containing more than 20 per cent. of thimet, phosdrin, parathion), Phosphorus (yellow), Selenium, Strychnine, Thallium.

APPENDIX G

FEES (ANNUAL).

		£	s.	d.
1.	Licence to Procure, Manufacture and Supply Poisons other than Drugs of Addiction (by Wholesale Dealing)	2	10	0
2.	Licence to Procure, Manufacture and Supply Drugs of Addiction (by wholesale Dealing)	2	10	0
3.	Pharmaceutical Chemist's Licence to Sell Poisons		10	0
4.	Retailer's Licence to Sell Poisons in Schedules 1, 2 and 6	1	0	0
5.	Retailer's Licence to Sell Poisons in Schedule 6		10	0
6.	Poisons Permit (Industrial)		10	0
7.	Licence to Hawk, Peddle, etc.	1	0	0
8.	Permit to Sell Veterinary Medicines containing 4th Schedule Drugs (reg. 39)		10	0

The fee for renewal of a licence shall be the same as for the original licence.

The fee for a licence issued after 31st December shall be one half the annual fee.

APPENDIX H.

FOURTH SCHEDULE DRUGS REFERRED TO IN REGULATION 39 (1).

Except where otherwise stated, a substance in this appendix includes any derivative, compound, preparation or admixture included in relation to that substance in the Fourth Schedule to the Act.—

Antihistamines, Ataractic substances, Chloral hydrate,
Chlorpromazine, Cortisone (for topical use only),
Penicillin, Streptomycin, Sulphanilamide.

PHARMACY ACT, 1964.

Department of Public Health,
Perth, 23rd June, 1965.

HIS Excellency the Governor in Executive Council, acting pursuant to the provisions of section 47 of the Pharmacy Act, 1964, and section 11 of the Interpretation Act, 1918-1962, with the recommendation of the Pharmaceutical Council of Western Australia constituted under the first mentioned Act, has been pleased to make the regulations set forth in the Schedule hereunder, to have and take effect on and after the 1st day of July, 1965.

D. J. R. SNOW,
Acting Commissioner of Public Health.

SCHEDULE.

REGULATIONS.

PART I.—PRELIMINARY.

1. These regulations may be cited as the Pharmacy Act Regulations, 1965.

2. The Pharmacy and Poisons Act Regulations, 1951 published in the *Government Gazette* on 12th October, 1951, and all subsequent amendments thereto are hereby revoked.

3. These regulations are divided into Parts as follows:—

Part I—Preliminary.—Regs. 1-4.
Part II—The Council of the Pharmaceutical Society.—Regs. 5-41.
Part III—Examinations and Practical Training.—Regs. 42-48.
Part IV—Registration of Pharmaceutical Chemists.—Regs. 49-53.
Part V—Annual Licenses to Practise.—Regs. 54-58.
Part VI—Registration of Pharmacies.—Regs. 59-65.
Part VII—Miscellaneous.—Regs. 66-67.
Appendix A—Forms.
Appendix B—Fees.
Appendix C—Schedule of basic dispensing equipment required in the pharmacy.

4. In these regulations unless the contrary intention appears—

"Deputy President" means the deputy president of the Council;
"dispensary" means that part of a pharmacy that is reserved for the dispensing of medicines;
"President" means the president of the Council; and
"the Act" means the Pharmacy Act, 1964.

PART II.—THE COUNCIL OF THE PHARMACEUTICAL SOCIETY.

5. The Council shall from time to time appoint a pharmaceutical chemist to be a returning officer, and to conduct elections in accordance with and as provided by these regulations.

6. The Council shall pay the returning officer for each and every election conducted by him the fee prescribed in Appendix B to these regulations, and a further sum sufficient to recoup him all authorised expenses incurred by him in conducting such election.

7. (1) The returning officer for the time being shall, in the month of March in each year, conduct an election to fill the vacancies resulting from effluxion of time of the period for which members holding office at the coming into operation of these regulations were elected or of those elected pursuant to subregulation (2) of this regulation.

(2) Of the seven members of the Council holding office at the coming into operation of the Act—

(a) the two members elected on 31st March, 1963 shall retire on 31st March, 1966;
(b) the three members elected on 31st March, 1964 shall retire on 31st March, 1967; and
(c) the two members elected on 31st March, 1965, shall retire on 31st March, 1968.

(3) Any vacancy resulting from the retirement of a member in pursuance of subregulation (2) of this regulation shall be filled by election and the person so elected shall hold office for three years.

(4) Any member who retires from the Council or whose term of office on the Council expires under this regulation shall, subject to the Act, be eligible for re-election to the Council.

(5) Where an extraordinary vacancy occurs such vacancy shall be filled by election of a pharmaceutical chemist in the manner prescribed for elections generally, and the member so elected shall hold office for the residue of the term during which the member in lieu of whom he is elected would have held office.

(6) At the first meeting of the Council after 31st March, 1966, and thereafter at the first meeting after the election of each Council, the members thereof shall elect two of their number to the respective offices of President and Deputy President, and the members so elected shall hold office until the first meeting of the Council following the next annual election.

8. (1) Notice of every election shall be posted to every financial member of the Pharmaceutical Society addressed to his last known place of address.

(2) The notice shall state—

(a) the date of the election; and

(b) the place, time and date (not being less than 14 days nor more than 28 days prior to the date of the election) for receipt of nominations.

Nominations.

9. Every nomination of a candidate at an election shall be lodged with the returning officer in the form of Form 1 in Appendix A to these regulations, signed by the candidate and by not less than three persons qualified to vote at the election.

10. (1) If the number of persons nominated is not greater than the number required to fill the vacancies, the returning officer shall forthwith make a return to the registrar, and declare the person or persons nominated to be elected as members.

(2) If the number of persons so declared to be elected is insufficient to fill the vacancies, the retiring President of the Council shall so report to the Governor in Council, who may thereupon appoint one or more qualified persons to fill such vacancy or vacancies.

11. (1) If the number of persons nominated is greater than the number required to fill the vacancies, a poll shall be taken by the returning officer, who shall cause voting papers to be printed in the form of Form 2 in Appendix A to these regulations.

(2) The returning officer shall send one voting paper, initialed by him, together with one unfastened envelope marked "Ballot Paper", and another unfastened envelope with the name and address of the returning officer printed thereon, by post in a sealed envelope to the address appearing in the register of every pharmaceutical chemist registered by the Council.

(3) A voter shall indicate the candidate or candidates for whom he votes by striking out clearly and distinctly the names of the candidate or candidates for whom he does not vote, but leaving untouched the same number of names on the ballot paper as there are vacancies to be filled.

12. (1) The returning officer shall at the time and at the place appointed for the holding of the ballot proceed, in the presence of the registrar and of the scrutineers (if any) appointed by any of the candidates, to open all the printed envelopes received by him, and to remove the voting papers therefrom and, if satisfied upon making a comparison of each of the signatures on the counterfoils of the voting papers with the signatures of the voters in the Signature Book that each vote has been regularly and properly given, and that no person entitled to vote has voted twice, he shall, after rejecting any informal ballot papers, proceed to ascertain the number of votes cast in favour of the respective candidates.

(2) As soon as conveniently may be thereafter the returning officer shall give notice thereof to the Council, and shall duly declare elected the candidates (not exceeding the number of vacancies) who have received the greatest number of votes or, in the event of an equal number of votes being received by two or more candidates, the candidate or candidates in whose favour he exercises his casting vote or votes.

(3) In the case of an equality of votes, the returning officer shall have a casting vote.

13. Any voting paper on which the names of candidates not struck out does not equal the number of members to be elected, or which has not been signed by the voter, or which is enclosed in an envelope other than the printed one aforesaid, shall be deemed informal and shall not be counted by the returning officer, except that where a candidates withdraws his nomination between the date of nomination and the date of the election, no voting paper shall be deemed to be invalid by reason of a vote cast in favour of that retiring candidate.

14. Any candidate for election desirous of retiring before the day of election shall, not later than seven clear days before the day of election, sign and deliver to the returning officer a notice in the form of Form 3 in Appendix A to these regulations, and if the number of candidates is reduced by such retirement to the number of members to be elected, the returning officer shall declare such remaining members duly elected.

15. The name and address of every successful candidate at any election, and the name and address of the President and the Deputy President upon their election, shall be published by the registrar in the *Government Gazette* within 28 days after the election.

16. Every candidate at an election shall be entitled to appoint by writing addressed and delivered to the returning officer one scrutineer, who shall be entitled to be present while the returning officer is opening and counting voting papers.

17. The omission of the returning officer to send or post to, or the non-receipt of any voting paper by, any voter within the time mentioned therein, or at all, shall not in any manner invalidate or affect the election.

Meetings of the Council.

18. The Council shall meet on the first Tuesday in every month, at such time and place as it shall from time to time appoint, and on such other days and times as the President or any two members may appoint by requisition in writing and delivered to the registrar, if the time so appointed by the President or any two members is sufficient to allow the registrar to summon the members of the Council as provided in regulation 19 of these regulations.

19. The registrar, on receipt of a requisition, referred to in regulation 18 of these regulations, shall summon the members of the Council by posting a letter or postcard to each of the members 48 hours at least before the time so appointed for the meeting.

20. The omission by the registrar to send or post to, or the non-receipt of any notice of any meeting by, any member of the Council within the time aforesaid or at all, shall not in any manner invalidate or affect any meeting.

21. If at the expiration of 30 minutes after the time appointed for the meeting there shall not be a quorum of members present, no business shall be transacted and the meeting shall lapse or may be adjourned by the member or members present to such time and place as he or they may determine.

22. Voting at meetings of the Council shall be by a show of hands: Provided that, if in any particular case any member present shall so request, voting shall be by ballot.

23. No resolution arrived at, or act, matter or thing done, or authorised by any meeting shall be rescinded or amended at any subsequent meeting, unless notice of such intended rescission or amendment shall be given in the notices convening the meeting at which such rescission or amendment is proposed.

24. At every meeting of the Council the business and proceedings and the conduct and management shall be dealt with, carried on, and regulated as provided from time to time by standing orders not inconsistent with the Act or these regulations or, in the absence or silence of such standing orders, as the Council may from time to time determine.

25. The Council may adopt by a resolution, a seal as and for the common seal of the Council, and such seal shall at all times be kept in the custody of the registrar, and deposited in the office of the Council.

26. The seal of the Council may be affixed by the registrar, in the presence of any member of the Council, to any instrument or writing, when authorised by a resolution passed for that purpose and entered upon the minutes of the proceedings of the Council, but not otherwise.

27. Every certificate granted under section 22 of the Act shall be signed by the President and registrar of the Council, and shall have affixed thereto the common seal of the Council.

28. The Council may from time to time appoint from amongst themselves such committee or committees, as may be thought fit, and may by resolution at any time abolish any committee so appointed, or modify or extend its power, or regulate its proceedings.

29. The President shall be an *ex officio* member of every such committee and, when present, shall preside.

30. The proceedings of such committee shall, as far as practicable and subject to any resolution, be regulated by the same standing orders which apply to the proceedings of the Council.

31. Minutes of every meeting shall be kept by the registrar, and such minutes, when signed by the chairman of the same or any subsequent meeting, shall be binding and conclusive for all purposes and before all courts of the proceedings at such meeting.

The Registrar.

32. The Council shall appoint a registered pharmaceutical chemist as registrar, who shall be paid by salary and shall hold office subject to one month's notice of termination of engagement by either side.

33. The registrar shall discharge such duties of office as he may be required to discharge by the Act and these regulations, and such further duties as the Council may from time to time determine, and he shall be subject at all times to the direction of the Council.

34. The registrar shall be in attendance at his office at such hours as may from time to time be appointed by the Council and be present at all meetings of the Council and committees, and make a report of all matters that come under his cognisance for the information of the Council and committees. He shall consult the President, or in his absence, the Deputy President on any business requiring attention between the various meetings, and obey the order and direction of the President or Deputy President as the case may be during such time, and he shall be responsible for the safe custody of all documents and property belonging to the Council which shall be under his control.

35. (1) The registrar shall keep and maintain a "Signature Book" in the form of Form 4 in Appendix A to these regulations, and shall enter therein in alphabetical order the name of every pharmaceutical chemist appearing in the register and obtain the signature of every such chemist.

(2) The "Signature Book" shall be produced for the inspection of the returning officer on the day of holding any election as provided by these regulations.

36. (1) The registrar shall receive all fees, fines, subscriptions, donations and other moneys that are due or payable to the Council, and shall give a printed receipt and no other, for the same, in the form approved by the Council, retaining a duplicate of such receipt.

(2) At least once in each month, and more often if required by the Council, the registrar shall pay into some bank appointed by the Council to the credit of an account called "Pharmaceutical Council of Western Australia", the amount of money so received by him.

37. All surplus funds to the credit of the said account, or such parts thereof as may be deemed advisable, shall be invested in such manner and upon such security as shall be authorised by a resolution of the Council.

Payment of Accounts.

38. (1) The registrar shall submit all accounts to the Council at its next monthly meeting, to be passed for payment by resolution of the Council.

(2) Every resolution shall specify the sum or sums of money to be paid and to whom such sum or sums is or are payable.

(3) No account shall be paid, except under the authority of a resolution so passed.

(4) Every account shall be paid by crossed cheque marked "Not negotiable", made payable to the person specified in the resolution passing the account for payment, and be signed by the persons authorised in that behalf from time to time by the Council.

The Honorary Treasurer.

39. (1) The Council shall from time to time appoint a member of the Council to be honorary treasurer, who shall hold office for one year and shall, on the expiration of his term of office, be eligible for re-appointment.

(2) The honorary treasurer shall discharge such duties, in addition to those prescribed by these regulations, as the Council may from time to time determine.

(3) The financial year shall be the period from 1st January to 31st December, and at the meeting held in the month of February the honorary treasurer shall present a statement of accounts covering the financial transactions of the Council during the last preceding financial year and present a balance sheet showing the assets and liabilities of the Council at the end of that financial year.

Auditors.

40. The Council shall in the month of April in each year appoint as auditors two fit and proper persons, who shall be eligible for re-appointment, to hold office until the 31st day of March in the following year.

41. The auditors shall—

(a) inspect the books and accounts of the Council;

(b) examine the annual balance sheet prepared by the honorary treasurer, and certify same if correct;

(c) investigate and examine all contracts, accounts, invoices, books, securities, and vouchers in anywise relating to or concerning the same which may be kept by or in the possession of the honorary treasurer, registrar, or any other person;

(d) examine the bank pass books or bank statements and ascertain that they correspond with the account of the Council in the bank;

(e) present an annual report to the Council stating the result of their inspection and examination.

PART III.—EXAMINATIONS.

42. The Council shall hold such examinations as are prescribed by these regulations at such times as it may deem necessary and shall cause any examination to be conducted by one or more examiners appointed under the Act.

43. The subjects for examinations shall be—

First Year—

Biology.
English Expression II.
Pharmaceutical Chemistry I.
Pharmaceutics I.
Physics I.
Mathematics Pharmaceutical.

Second Year—

Pharmaceutical Chemistry IIA.
Pharmaceutical Chemistry IIB.
Pharmaceutics II.
Microbiology and Immunology.
Psychology I.
Physiology I.
Biometrics.

Third Year—

Pharmaceutical Chemistry III.
Pharmaceutics III.
Pharmacology.
Pharmacognosy.
Forensic Pharmacy.
Pharmaceutical Management,

but this regulation does not apply to apprentices referred to in paragraph (b) of subsection (1) of section 21 of the Act in respect of whom the provisions regarding examination subjects which apply immediately prior to the commencement of these regulations shall continue to apply.

44. (1) The examiner or examiners, as the case may be, by whom an examination under these regulations is conducted shall certify in writing to the registrar in relation to each candidate who presented himself for the examination, whether the candidate has or has not satisfactorily passed the examination.

(2) The registrar shall submit any certificate given in pursuance of subregulation (1) of this regulation to the Council for approval, and shall thereafter retain each certificate.

45. Applications to sit for any examination shall be lodged with the registrar at such time and place as may from time to time be determined by the Council, shall be in the form approved by the Council and shall be accompanied by the appropriate fees set out in Appendix B to these regulations.

46. A person is not eligible to commence a course of practical training for the purpose of meeting the requirements of subparagraph (i) of paragraph (a) of subsection (1) of section 21 of the Act, unless he satisfies the Council that he has passed all prescribed examinations or such examinations as in the opinion of the Council are substantially equivalent thereto.

Practical Training.

47. (1) For the purposes of section 21 of the Act a course of practical training shall—

(a) occupy not less than 2,000 hours;
(b) be served under the personal supervision of a pharmaceutical chemist;
(c) include instruction and experience in current dispensing practice and pharmaceutical administration;
(d) be undertaken and completed under Articles of Traineeship between the trainee and the pharmaceutical chemist who conducts the pharmacy, or the pharmaceutical chemist in charge of the pharmacy department of a hospital, as the case may require.

(2) Each trainee shall submit his Articles of Traineeship to the Council for registration within 28 days of commencement of his training.

(3) Each trainee shall keep a record in the form of Form 10 in Appendix A to these regulations, showing the practical work carried out by him, and a duplicate copy of the entries made therein shall be submitted to the Council not later than 14 days following the close of each three monthly period.

48. Credit for practical training shall not be granted if—

(a) the period of continuous service with one employer is less than 150 hours;
(b) the record required by regulation 47 of these regulations does not satisfy the Council that the trainee has carried out the course of practical training prescribed by these regulations; or
(c) the Articles of Traineeship have not been approved by the Council.

PART IV.—REGISTRATION OF PHARMACEUTICAL CHEMISTS.

49. The Council shall, for the purpose of paragraph (d) of subsection (1) of section 21 of the Act, recognise the certificates or diplomas of competency as a pharmaceutical chemist or as a chemist and druggist of the Societies, Colleges or Boards of Pharmacy set forth hereunder, namely—

The Pharmacy Board of New Zealand;
The Pharmacy Board of New South Wales;
The Pharmacy Board of Queensland;
The Pharmacy Board of South Australia;
The Pharmacy Board of Tasmania;
The Pharmacy Board of Victoria;
The Pharmaceutical Society of Great Britain;
The Pharmaceutical Society of Ireland;
The Pharmaceutical Society of Northern Ireland.

50. Application for registration shall be made to the registrar in the form of Form 5 or Form 6, as the case may be, in Appendix A to these regulations, signed by the applicant and accompanied by all necessary documents and certificates and the fee prescribed in Appendix B to these regulations.

51. An applicant for registration shall, if requested by the registrar, supply to the Council such information or evidence (oral or in writing), as the Council may from time to time require, and may be required by the registrar to attend in person before the Council for that purpose.

52. The registrar shall, upon the granting of a registration by the Council and payment of the fee prescribed in Appendix B to these regulations, issue to the applicant a certificate in the form in the Third Schedule to the Act.

53. Every pharmaceutical chemist, who changes his place of residence, business, or employment, shall thereupon notify in writing, either personally or by registered letter, such change of address, giving particulars of both the old and new addresses.

PART V.—ANNUAL LICENCES TO PRACTISE.

54. Every registered pharmaceutical chemist desirous of applying to the Council for a licence or renewal of a licence to practise or carry on business as a pharmaceutical chemist shall sign and deliver to the registrar an application in the form of Form 7 in Appendix A to these regulations.

55. The Council shall consider such application at its next meeting and may by resolution grant to the applicant a licence in the form of Form 8 in Appendix A to these regulations, or may refuse the application.

56. The registrar shall forthwith give to an applicant whose application has been refused by the Council notice by registered letter in the form of Form 9 in Appendix A to these regulations.

57. Every pharmaceutical chemist practising or carrying on business as such shall keep his current licence to practise posted in a conspicuous place in a portion of his place of business to which the public have access.

Erasure of Name from the Register.

58. Any pharmaceutical chemist whose name has been ordered by the Council to be erased from the register pursuant to subsection (5) of section 26 of the Act, shall forthwith be informed of such fact by the registrar by registered letter.

PART VI.—REGISTRATION OF PHARMACIES.

59. A person desirous of applying to the Council for the registration of premises as a pharmacy shall complete, sign and deliver to the registrar an application, accompanied by the requisite fee, in the form of Form 11 in Appendix A to these regulations.

60. The Council shall consider such application and may by resolution grant to the applicant a certificate in the form prescribed, issue a conditional certificate or refuse the application.

61. (1) A pharmacy may not be registered under section 23 of the Act unless the Council is satisfied that—

(a) the premises used or intended to be used as a pharmacy, whether a single unit or portion of larger premises, are self-contained by being walled, floored and ceiled, have no means of access to any adjacent other type of professional or business premises and have a separate entrance from a street or public thoroughfare;

(b) the pharmacy is well lit and adequately ventilated;

(c) the premises and fittings therein, and all equipment, utensils and apparatus used or to be used in dispensing and storing of drugs are, and will be maintained, in clean condition and proper state of efficiency and repair;

(d) the dispensary area has a minimum floor space of 90 square feet and is equipped with a suitable sink having both hot water and cold water connected thereto;

(e) the dispensary is equipped with the basic scale of apparatus and equipment and reference books specified in Appendix C to these regulations;

(f) proper and adequate provision is made for fully recording and maintaining a record of all prescriptions and repeat prescriptions dispensed in the pharmacy.

(2) Notwithstanding the provisions of paragraph (a) of subregulation (1) of this regulation, if at the commencement of these regulations any premises are in an advanced stage of construction as a pharmacy, or are premises in which the business of a pharmaceutical chemist is being carried on, those premises may be registered as a pharmacy if the Council is satisfied that the area of the premises used or intended to be used as a pharmacy is clearly defined and identified.

(3) For the purposes of paragraph (f) of subregulation (1) of this regulation, prescriptions and repeat prescriptions shall be recorded in an appropriate bound prescription book or other system approved by the Council, except that records of National Health Service prescriptions and Repatriation Department prescriptions may be in the form of duplicates kept in sequence in marked monthly order that are readily identifiable and accessible.

62. Where in any case application is made for registration of a pharmacy and circumstances exist which render it impracticable to comply immediately with the requirements of regulation 61 of these regulations, or in the opinion of the Council it is not in public interest to require such compliance, the Council may register the pharmacy subject to such conditions as it may determine.

Service of Notice.

63. The Council may in its discretion and for good cause shown or for repeated or continued breach of the conditions governing registration of a pharmacy, cancel the certificate issued.

64. On receipt of notice of cancellation of registration, the holder shall return the certificate to the Council by hand or registered post, within 7 days.

65. A certificate issued in accordance with regulation 60 of these regulations is not transferable either to a person or to other premises.

PART VII.—MISCELLANEOUS.

66. All books, records and documents which are required to be kept or retained for a prescribed period shall (unless otherwise prescribed), in the case of books or records, be preserved for a period of two years from the date on which the last entry is made therein, and, in the case of any documents, for a period of two years from the date on which it is first received.

67. Any person who contravenes or neglects, refuses, or fails to comply with any provisions of these regulations shall be guilty of an offence.

APPENDIX A—FORMS.

Form 1.

Western Australia.

Pharmacy Act, 1964.

NOMINATION PAPER.

We, the undersigned pharmaceutical chemists of Western Australia, do hereby nominate ..... of .....as a candidate for the office of a member of the Pharmaceutical Council of Western Australia at the election to be held on the..... .....day of.....19.....

.....
Pharmaceutical Chemist.

.....
Pharmaceutical Chemist.

.....
Pharmaceutical Chemist.

And I the abovenamed..... ..... do hereby consent to such nomination.

.....
Pharmaceutical Chemist.

Form 2.

Western Australia.

Pharmacy Act, 1964.

VOTING PAPER.

Candidates for election as members of the Pharmaceutical Council of Western Australia:—

(Arrange in alphabetical order of surname)

Directions.

The voter is to strike out the name of the candidate or candidates for whom he does not intend to vote by drawing a line through the name or names of such candidate or candidates. He must be careful not to leave uncancelled names of more or less than ..... candidates, otherwise this voting paper will be invalid.

The voter must insert his voting paper in the printed envelope, fasten it and post it to the address of the returning officer in time to be received before.....o'clock on the .....day of.....19..... at which time the Ballot will close.

.....

(Perforated)

COUNTERFOIL.

Name of Voter..... Signature of Voter.....
(In Block Letters)

Note.—Counterfoil to be detached, and not enclosed in ballot paper envelope but to be placed loose in envelope addressed to returning officer.

Form 3.

Western Australia.

Pharmacy Act, 1964.

NOTICE OF WITHDRAWAL OF NOMINATION.

We, the undersigned, nominators of.....as a candidate at the election of the Pharmaceutical Council of Western Australia to be held on the.....day of....., 19....., do hereby withdraw the said.....as a candidate.

.....
Pharmaceutical Chemist.

.....
Pharmaceutical Chemist.

.....
Pharmaceutical Chemist.

And I, the said....., do hereby retire from being such a candidate.

.....
Pharmaceutical Chemist.

Form 4.

Western Australia.

Pharmacy Act, 1964.

SIGNATURE BOOK.

Surname.	Christian Names.	No. in Register.	Signature.

Form 5.

Western Australia.

Pharmacy Act, 1964.

(Regulation 50.)

APPLICATION TO BE REGISTERED AS A PHARMACEUTICAL CHEMIST.

I, ..., of .....
being of (or over) the age of 21 years and—

(i) having served for a period of at least four years as an apprentice in the business of..... a chemist and druggist (or a pharmaceutical chemist) of....., in the State of.....(or in the Dominion of New Zealand), in the keeping of open shop for the compounding and dispensing of prescriptions of legally qualified medical practitioners;

or

(ii) having completed a prescribed course of practical training of not less than 2,000 hours duration with a pharmaceutical chemist or chemists in accordance with prescribed conditions and in premises approved by the Council,

and having passed—

(i) all the examinations prescribed by the Pharmacy Act, 1964 regulations;

or

(ii) .....
.....

do hereby apply to the Pharmaceutical Council of Western Australia, pursuant to section 21 of the Pharmacy Act, 1964, to be registered by the said Council as a pharmaceutical chemist.

I annex hereto the necessary supporting documents, namely:—

.....
.....
.....
.....

.....
Signature of Applicant.

To the Registrar,
Pharmaceutical Council of Western Australia,
Technical College, Perth.

Form 6.

Western Australia.

Pharmacy Act, 1964.

(Regulation 50.)

APPLICATION FOR REGISTRATION AS A PHARMACEUTICAL CHEMIST.

I, (a) ..... of .....,
being of (or over) the age of 21 years, and being the holder of the certificate (or diploma) of competency as a pharmaceutical chemist (or as a chemist and druggist) of the (c)....., do hereby apply to the Pharmaceutical Council of Western Australia, pursuant to subsection (1) (d) of section 21 of the Pharmacy Act, 1964, to be registered by the said Council as a pharmaceutical chemist.

.....
Signature of Applicant.

To the Registrar,
Pharmaceutical Council of Western Australia,
Technical College, Perth.

(a) Name of Applicant in full. (b) Address of Applicant. (c) Name of Society, College or Board of Pharmacy recognised by the regulations.

Form 7.

Western Australia.

Pharmacy Act, 1964.

(Regulation 54.)

APPLICATION FOR A LICENCE (OR RENEWAL OF A LICENCE) TO PRACTISE AS A PHARMACEUTICAL CHEMIST.

I, ..., of .....,
in the State of Western Australia, duly registered on the.....
.....19....., by the Pharmaceutical Council of Western Australia as a pharmaceutical chemist, do hereby apply to the said Council for a licence to practise as a pharmaceutical chemist for the year ending the 30th day of June, 19.....

Dated this.....day of....., 19.....

.....
Signature of Applicant.

To the Registrar,
Pharmaceutical Council of Western Australia,
Technical College, Perth.

Form 8.

Western Australia.

Pharmacy Act, 1964.

(Regulation 55.)

ANNUAL LICENCE TO PRACTISE AS A PHARMACEUTICAL CHEMIST.

I, the undersigned, on behalf of the Pharmaceutical Council of Western Australia, do hereby certify that.....
of....., duly registered by the said Council as a pharmaceutical chemist, is licensed to practise or carry on business as such within the State of Western Australia until the 30th day of June, 19.....

Dated at.....this.....day of.....
19.....

....
Registrar President of the Pharmaceutical Council of Western Australia.

Form 9.

Western Australia.

Pharmacy Act, 1964.

(Regulation 56.)

NOTICE OF REFUSAL TO GRANT LICENCE.

I hereby give you notice that at a meeting of the Pharmaceutical Council of Western Australia, held at Perth on.....your application for a licence to practise as a pharmaceutical chemist was refused by the Council.

.....

Registrar

Pharmaceutical Council of Western Australia.

Form 10.

Western Australia.

Pharmacy Act, 1964.

(Regulation 47.)

RECORD OF PRACTICAL TRAINING.

I certify that the above student has completed practical training under my supervision in my/our premises, which are approved by the Pharmaceutical Council of Western Australia.

Name and Address of Approved Premises	Date of Commencement	Date of Completion	Number of Hours	Signature of Chemist in Charge
.....	.....	.....	.....	.....
.....	.....	.....	.....	.....
.....	.....	.....	.....	.....

.....

Progressive Total of Hours

Each three months a duplicate copy of the entries made herein shall be submitted to the Council within 14 days of the close of each period.

Form 11.

Western Australia.

Pharmacy Act, 1964.

(Regulation 59.)

APPLICATION FOR A CERTIFICATE OF REGISTRATION AS A PHARMACY.

1. Plan of Premises	2. Application and Declaration	3. Ext. of Regulations Appendix C	4. Appendix D

Form 12.

Western Australia.

Pharmacy Act, 1964.

(Regulation 60.)

CERTIFICATE OF REGISTRATION AS A PHARMACY.

Pharmaceutical Council of Western Australia.

This is to certify that the premises known as.....
..... situated at .....
and at which the business of a Pharmaceutical Chemist is carried on by/or under the managership of.....are registered under section 23 of the Pharmacy Act, 1964, as a Pharmacy until the 30th day of June, 19.....

.....19..... .....
Registrar.

APPENDIX B—FEES.

Sect. 47 (c).

	£	s.	d.
For Conducting an Election	5	0	0
For Third Year Examination	5	0	0
For Single Subjects	2	10	0
For Fourth Year Examination	7	10	0
For Single Subjects	5	0	0
Maximum as for complete examination in each case.			
For Registration of Articles of Traineeship	1	0	0
For Transfer of Articles of Traineeship		10	0
For Registration of Pharmaceutical Chemist	5	0	0
For Certified Copy of Certificate of Registration	1	0	0
For Annual Licence to Practice	2	10	0
For Certificate of Competency for Registration elsewhere than in Western Australia	1	0	0
For Registration of Pharmacies (including the issue of Certificate)	5	0	0

APPENDIX C.

Pharmacy Act, 1964.

(Regulation 61.)

BASIC SCALE OF APPARATUS, EQUIPMENT AND REFERENCE BOOKS REQUIRED BY A REGISTERED PHARMACY.

Autoclave: Small autoclave or approved substitute	1 only
Beakers:	
100 ml.	1 only
250 ml.	1 only
Bunsen Burner or equivalent or alternative method of heating, such as an electric hotplate	1 only
Funnels:	
glass or plastic 2 in.	1 only
glass or plastic 5 in.	1 only
sintered glass No. 3	1 only
vulcanite two piece	1 only
Hood for aseptic dispensing (approved type)	1 only

Measures Graduated (dispensing glass):

Imperial:

2 fl. dr. graduated to minims	1 only
1 fl. oz.	1 only
2 fl. oz.	1 only
4 fl. oz.	1 only
10 fl. oz.	1 only

Metric:

10 ml.	1 only
20 ml.	1 only
50 ml.	1 only
100 ml.	1 only
200 ml.	1 only
1 litre	1 only

Mortars and Pestles:

Glass 3 in.	1 only
Wedgewood Asstd.	2 only

*Poison Cupboard 1 only

Refrigerator: A refrigerator for storage of biological preparations, insulin, heparin and other preparations at temperatures required by The British Pharmacopoeia 1 only

Stirring Rods: Glass and Vulcanite—2 each different sizes.

Scales:

Dispensing Beam type to weigh to 2 oz. or 50 gm.	1 only
Counter Beam type to weigh to 2 lb. or 1 kg.	1 only

Sieve: Not coarser than 60 mesh 1 only

Slabs Ointment: 10 in. x 10 in. (minimum size) 1 only

Spatulas:

Stainless Steel 7 in.	2 only
Stainless Steel 10 in.	1 only
Vulcanite 4 in.	1 only
Vulcanite 6 in.	1 only

Thermometer Chemical 0°-200° C. 1 only

Tripod and Asbestos Gauze (if burner used for heating) 1 only

Water Bath 1 only

Water Pump: Some type of small vacuum pump to work from a tap and to provide negative pressure to filter eye drops through the sintered glass funnel.

Weights:

Apothecaries—$\frac{1}{2}$-6 grs. and $\frac{1}{2}$ scruple to 4 drachms inclusive	1 set
Avoirdupois—$\frac{1}{2}$ oz.-1 lb. inclusive	1 set
Metric—5 mgm.-500 gm. inclusive	1 set

Books:

Current copy of British Pharmacopoeia and Addenda.
Current copy of British Pharmaceutical Codex and Addenda.
Current copy of Australian Pharmaceutical Formulary.
Current copy of Martindale's Extra Pharmacopoeia Vol. 1.
Current copy of P.P. Guide.

* "Poisons Cupboard" means a substantially made cupboard provided with an effective locking device, and having the word "Poison" conspicuously painted on the outside of the cupboard. Such cupboard shall be securely fastened to a portion of the premises and not used for any purpose other than the storage of Poisons.

FACTORIES AND SHOPS ACT, 1963-1964.

Department of Labour,
Perth, 23rd June, 1965.

HIS Excellency the Governor in Executive Council, acting pursuant to the provisions of the Factories and Shops Act, 1963-1964, and a recommendation in that regard, has been pleased to make the regulations set forth in the Schedule hereunder.

T. H. BURGESS,
Chief Inspector of Factories.

Schedule.

Regulations.

1. In these regulations the Factories and Shops (Rostered Extraordinary Trading Hours) Regulations, 1964, made under the provisions of the Factories and Shops Act, 1963-1964, and published in the *Government Gazette on the* 30th December, 1963, and amended from time to time by notices published in the *Government Gazette,* are referred to as the principal regulations. Principal regulations.

2. The principal regulations are amended by adding immediately after Part XIII the following part:—

Part XIV.

Division 1.

Zone No. 14—Narrogin District.

All that portion of the State contained within a radius of five miles from P.O., Narrogin.

Division 2.

Shops in Zone No. 14.

Description of Shops in Zone No. 14—Inclusive Dates during which Shop is required to be Kept Open during Rostered Extraordinary Trading Period for Extraordinary Trading Times.

Subdivision (i).

Mols Motors, 4 Kipling Street, Narrogin—28th June, 1965, to 4th July, 1965.

Berson's Service Station, 111 Federal Street, Narrogin—5th July, 1965, to 11th July, 1965.

Young's Ampol Service Station, Clayton Road, Narrogin—12th July, 1965, to 18th July, 1965.

Jeffery's Caltex Service Station, Federal Street, Narrogin—19th July, 1965, to 25th July, 1965.

R. A. Reilly and Co. Pty. Ltd., Federal Street, Narrogin—26th July, 1965, to 1st August, 1965.

S. J. & U. N. McEntee, Federal Street, Narrogin—2nd August, 1965, to 8th August, 1965.

Elliots (Narrogin), Federal Street, Narrogin—9th August, 1965, to 15th August, 1965.

Dorsett Motors Holdings, Federal Street, Narrogin—16th August, 1965, to 22nd August, 1965.

R. A. Reilly and Co. Pty. Ltd., Egerton Street, Narrogin—23rd August, 1965, to 29th August, 1965.

Jeffery's Caltex Service Station, Federal Street, Narrogin—30th August, 1965, to 5th September, 1965.

R. A. Reilly and Co. Pty. Ltd., Federal Street, Narrogin—6th September, 1965, to 12th September, 1965.

Berson's Service Station, 111 Federal Street, Narrogin—13th September, 1965, to 19th September, 1965.

Young's Ampol Service Station, Clayton Road, Narrogin—20th September, 1965, to 26th September, 1965.

S. J. & U. N. McEntee, Federal Street, Narrogin—27th September, 1965, to 3rd October, 1965.

Elliots (Narrogin), Federal Street, Narrogin—4th October, 1965, to 10th October, 1965.

Mols Motors, 4 Kipling Street, Narrogin—11th October, 1965, to 17th October, 1965.

R. A. Reilly and Co. Pty. Ltd., Egerton Street, Narrogin—18th October, 1965, to 24th October, 1965.

Subdivision (ii).

R. A. Reilly and Co. Pty. Ltd., Federal Street, Narrogin—3rd and 4th July, 1965.

S. J. & U. N. McEntee, Federal Street, Narrogin—10th and 11th July, 1965.

Elliots (Narrogin), Federal Street, Narrogin—17th and 18th July, 1965.

Mols Motors, 4 Kipling Street, Narrogin—24th and 25th July, 1965.

Berson's Service Station, 111 Federal Street, Narrogin—31st July and 1st August, 1965.

R. A. Reilly and Co. Pty. Ltd.., Egerton Street, Narrogin—7th and 8th August, 1965.

Jeffery's Caltex Service Station, Federal Street, Narrogin—14th and 15th August, 1965.

Young's Ampol Service Station, Clayton Road, Narrogin—21st and 22nd August, 1965.

Mols Motors, 4 Kipling Street, Narrogin—28th and 29th August, 1965.

S. J. & U. N. McEntee, Federal Street, Narrogin—4th and 5th September, 1965.

Elliots (Narrogin), Federal Streeet, Narrogin—11th and 12th September, 1965.

Dorsett Motors Holdings, Federal Street, Narrogin—18th and 19th September, 1965.

Jeffery's Caltex Service Station, Federal Street, Narrogin—25th and 26th September, 1965.

R. A. Reilly and Co. Pty. Ltd., Egerton Street, Narrogin—2nd and 3rd October, 1965.

Dorsett Motors Holdings, Federal Street, Narrogin—9th and 10th October, 1965.

Berson's Service Station, 111 Federal Street, Narrogin—16th and 17th October, 1965.

Young's Ampol Service Station, Clayton Road, Narrogin—23rd and 24th October, 1965.

Division 3.

Extraordinary Trading Times for Shops in Zone No. 14.

The extraordinary trading times during which shops described in Subdivisions (i) and (ii) of Division 2 of this Part are permitted and required to be kept open during rostered extraordinary trading periods for extraordinary trading times are as follows:—

Subdivision (i).

7 p.m. to 10 p.m. each Monday, Tuesday, Wednesday, Thursday, Friday;
1 p.m. to 10 p.m. each Saturday; and
7 a.m. to 10 p.m. each Sunday.

Subdivision (ii).

1 p.m. to 10 p.m. on Saturday; and
7 a.m. to 10 p.m. on the next succeeding Sunday.

ABATTOIRS ACT, 1909-1964.

Department of Agriculture,
South Perth, 21st June, 1965.

HIS Excellency the Governor in Executive Council, acting pursuant to the provisions of the Abattoirs Act, 1909-1964, has been pleased to make the regulations set out in the schedule hereunder, with effect on and after the 1st July, 1965.

F. L. SHIER,
Acting Director of Agriculture.

Schedule.

Regulations.

Principal regulations. 1. In these regulations the regulations made under the provisions of the Abattoirs Act, 1909-1964, published in the *Government Gazette* on the 14th April, 1938, and amended from time to time thereafter by notices published in the *Government Gazette*, are referred to as the principal regulations.

Reg. 19A revoked. 2. Regulation 19A of the principal regulations is revoked.

ABATTOIRS ACT, 1909-1964.

Department of Agriculture,
South Perth, 22nd June, 1965.

HIS Excellency the Governor in Executive Council, acting pursuant to the provisions of the Abattoirs Act, 1909-1964, has been pleased to make the regulations set out in the schedule hereunder, with effect on and after the 1st July, 1965.

F. L. SHIER,
Acting Director of Agriculture.

Schedule.

Regulations.

Principal regulations. 1. In these regulations the regulations made under the provisions of the Abattoirs Act, 1909-1964, published in the *Government Gazette* on the 14th April, 1938, and amended from time to time thereafter by notices published in the *Government Gazette*, are referred to as the principal regulations.

Reg. 19 substituted. 2. The principal regulations are amended by substituting for regulation 19 the following regulation:—

19. The fees to be charged for slaughtering of stock at the abattoirs (inclusive of inspection and 24 hours' free storage in the chilling rooms) shall be as follows:—

	Per Head. s.	d.
(i) Cattle—		
From 201-249 lb. dressed weight	35	9
From 250-400 lb. dressed weight	42	9
From 401-600 lb. dressed weight	50	0
Over 600 lb. dressed weight	57	6
(ii) Calves—		
Up to 100 lb. dressed weight	9	6
From 101-150 lb. dressed weight	12	9
From 151-200 lb. dressed weight	27	0
(iii) Sheep	5	6
(iv) Lambs	5	0

	Per Head. s. d.
(v) Pigs—	
Suckers—up to 22 lb. dressed weight	3 3
From 23-110 dressed weight	12 3
From 111-179 lb. dressed weight	15 0
Over 179 lb. dressed weight	17 9

Extra Charges.

	Per Head. s. d.
(i) Bulls 300 lb. and over (chilled weight and genuine stags	7 3
(ii) Tubercular and/or gangrenous cattle	7 3
(iii) Tubercular injured or septic calves	3 8
	Per 100. s. d.
(iv) Rams and genuine stags	132 1
(v) Ram lambs 50 lb. and over (chilled weight)	132 1
(vi) Injured, maggoty, daggy, downer, objectionably cippled, objectionably wet or dirty sheep or lambs	66 0
(vii) Full wool sheep	66 0
(viii) Sheep over 62 lb. (chilled weight)	66 0

Agistment Charges.

The fees to be charged for agistment of live-stock at the abattoirs (after the first 24 hours) shall be as follows:—

	Per Head Per Day. s. d.
(i) Cattle (based on 16 lb. hay per head per day)	2 0
(ii) Sheep, lambs and pigs (based on 2 lb. chaff per head per day for sheep and 1½ lb. crushed wheat per head per day for pigs)	6

MINES REGULATION ACT, 1946-1961.

Mines Department,
Perth, ..., 1965.

HIS Excellency the Governor in Executive Council, acting pursuant to the provisions of the Mines Regulation Act, 1946-1961, has been pleased to make the regulations set forth in the schedule hereunder.

A. H. TELFER,
Under Secretary for Mines.

Schedule.

Regulations.

Principal Regulations.

1. In these regulations the Mines Regulation Act Regulations made under the Mines Regulation Act, 1946-1961, reprinted pursuant to the Reprinting of Regulations Act, 1954, and published as so reprinted in the *Government Gazette* on the 3rd May, 1965, are referred to as the principal regulations.

2. The principal regulations are amended, by substituting for regulation 100 the following regulation:— Reg. 100 substituted.

100. (1) A person shall not travel, in a shaft or winze, in or on any cage, skip, carriage, receptacle or platform, carrying, timber, pipes, rails, ore, waste rock or similar material, or tools, unless authorised in writing by the District Inspector of Mines so to do.

(2) This regulation does not apply to—

(a) a person repairing a shaft;

(b) subject to subregulation (3), a platman;

(c) a person carrying small tools in a suitable bag or container; or

(d) a surveyor travelling with his instruments.

(3) A platman shall not—

(a) travel with any material that is insecurely fastened; or

(b) travel upwards with drill steel, pipes or material of similar form.

3. The principal regulations are amended, by substituting for regulation 101 the following regulation:— Reg. 101 substituted.

101. (1) The Manager shall ensure that a securely fastened gate, or gates, is or are used on every cage in which a person is travelling, and that provision is made to prevent the feet of any person from protruding outside the cage.

(2) At change of shift, every person appointed by the manager for that purpose is responsible for the safe ingress and egress of men, and for the proper fastening of the gate or gates.

Police Act Amendment Act (No. 2), 1964.

PROCLAMATION

WESTERN AUSTRALIA, TO WIT, DOUGLAS ANTHONY KENDREW, Governor. [L.S.] } By His Excellency Major-General Sir Douglas Anthony Kendrew, Knight Commander of the Most Distinguished Order of Saint Michael and Saint George, Companion of the Most Honourable Order of the Bath, Commander of the Most Excellent Order of the British Empire, Companion of the Distinguished Service Order, Governor in and over the State of Western Australia and its Dependancies in the Commonwealth of Australia.

WHEREAS it is enacted by section 2 of the Police Act Amendment Act (No. 2), 1964, that the Act shall come into operation on a date to be fixed by Proclamation: Now, therefore I, the Governor, acting with the advice and consent of the Executive Council, do hereby fix the 1st day of July, 1965, as the day on which the Police Act Amendment Act (No. 2), 1964 shall come into operation.

Given under my hand and the Public Seal of the said State, at Perth, this 23rd day of June, 1965.

By His Excellency's Command,

J. F. CRAIG,
Minister for Police.

GOD SAVE THE QUEEN ! ! !

98630/6/65

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