1972/96

THE THERAPEUTIC DRUGS (PERMITTED SALES) REGULATIONS 1972

ARTHUR PORRITT, Governor-General

ORDER IN COUNCIL

At the Government House at Wellington this 1st day of May 1972

Present:

HIS EXCELLENCY THE GOVERNOR-GENERAL IN COUNCIL

PURSUANT to the Food and Drug Act 1969, His Excellency the Governor-General, acting by and with the advice and consent of the Executive Council, hereby makes the following regulations.

REGULATIONS

1. **Title and commencement**—(1) These regulations may be cited as the Therapeutic Drugs (Permitted Sales) Regulations 1972.

(2) These regulations shall come into force on the 1st day of June 1972.

2. **Interpretation**—In these regulations, "the Act" means the Food and Drug Act 1969.

3. **Permitted sales**—Subject to sections 12 and 14 and subsection (1) of section 18 of the Act, and to any other enactment, the sale of a therapeutic drug by any person is permitted, for the purposes of subsection (2) of the said section 18, if—

(a) The drug is named or described in Part I or Part II of the Schedule to these regulations; or

(b) The drug is a food as well as a drug; or

(c) The drug is a cosmetic, dentifrice, or detergent, within the natural and ordinary meaning of those words; or

(d) The drug was available in New Zealand and could be lawfully sold by any person at the 31st day of March 1970:

Provided that paragraph (d) of this regulation shall cease to have effect with the close of the 31st day of March 1973.

4. **Revocation**—The Therapeutic Drugs (Permitted Sales) Regulations 1970* are hereby revoked.

*S.R. 1970/53

SCHEDULE

Reg. 3 (a)

THERAPEUTIC DRUGS THAT MAY BE SOLD FREELY

Part I

Acetic acid
Aromatic ammonia solution
Aspirin tablets
Camphor
Camphor liniment
Cascara liquid extract
Cascara tablets
Castor oil
Cod liver oil
Compound benzoin tincture
Compound effervescent powder
Compound effervescent powder (double strength)
Compound figs syrup
Compound liquorice powder
Compound magnesium carbonate tablets
Compound sodium bicarbonate tablets
Dilute ammonia solution
Eucalyptus oil
Glycerin
Hamamelis water (distilled witch hazel)
Heavy magnesium carbonate
Herbs, in dry uncompounded form, packed under the traditional or botanical name
Hydrous wool fat
Light magnesium carbonate
Liquid senna extract
Liquid paraffin
Liquid paraffin emulsion
Magnesium hydroxide mixture
Magnesium hydroxide tablets
Magnesium sulphate (Epsom salts)
Malt extract with cod liver oil
Senna fruit
Senna leaf
Sodium sulphate (Glauber's Salt)
Sulphur ointment
Sulphur, refined
Weak iodine solution
White liniment
White soft paraffin
Yellow soft paraffin
Zinc cream
Zinc ointment
Zinc and castor oil ointment

SCHEDULE—*continued*

Part II

Alka-Seltzer
Andrews Health Salt
Bonnington's Irish Moss
Califig
De Witts Antacid Powder
Nixoderm
Rawleigh's Anti-Pain Oil
Rawleigh's Camphor Balm
Rawleigh's Cherry Flavour Cough Syrup
Rawleigh's Cold Tablets
Rawleigh's Compound Mustard Ointment
Rawleigh's Cough Control
Rawleigh's Cough Tablets
Rawleigh's Healing Salve
Rawleigh's Laxative Tablets
Rawleigh's Liniment
Rawleigh's Liquid Antiseptic and Mouthwash
Rawleigh's Medicated Ointment
Rawleigh's Medicated Vapour
Rawleigh's Penetrating Lanolin Rub
Rawleigh's Pleasant Relief
Rawleigh's Ready Relief
Rawleigh's Ru-Mex-Ol Compound
Rawleigh's Stainless Vapor Balm

P. J. BROOKS,
Clerk of the Executive Council.

EXPLANATORY NOTE

This note is not part of the regulations, but is intended to indicate their general effect.

These regulations, which come into force on the 1st day of June 1972, specify the therapeutic drugs that may be sold freely in New Zealand (subject to sections 12, 14, and 18 (1) of the Food and Drug Act 1969).

Issued under the authority of the Regulations Act 1936.
Date of notification in *Gazette*: 4 May 1972.
These regulations are administered in the Department of Health.