

## Classification of Medicines

Pursuant to section 106(1) of the Medicines Act 1981, I, Derek Fitzgerald, Acting Group Manager, Medsafe, Ministry of Health, acting under delegated authority, hereby declare the following:

1. The medicines listed in Schedule 1 to this notice are classified as prescription medicines.
2. The medicines listed in Schedule 2 to this notice are classified as restricted medicines.
3. The medicines listed in Schedule 3 to this notice are classified as pharmacy-only medicines.

Every reference to a medicine in this notice applies whether the medicine is synthetic in origin or is from biological or mineral sources.

Unless specific reference is made otherwise, every reference applies also to medicines that are:

- a. preparations and admixtures containing any proportion of any substance listed in the notice.
- b. salts and esters of any substance listed in the notice.
- c. preparations or extracts of biological materials listed in the notice.
- d. salts or oxides of elements listed in the notice.

Unless specific reference is made otherwise, every reference to a medicine applies:

- i. if the medicine is in an injection or eye preparation, to any concentration of that medicine; and
- ii. if the medicine is not in an injection or eye preparation, only if the concentration of the medicine is greater than 10 milligrams per litre or per kilogram.

Where any reference is modified by a statement of the strength of the medicine, the strength is calculated using the free acid, base, alcohol or element unless specifically stated otherwise.

In accordance with section 106(2) of the Act, to the extent that any part of this notice is inconsistent with any provisions of any regulations made under section 105(1)(j) of the Act, the provisions in those regulations cease to have effect while this notice remains in force.

### Schedule 1

#### **Prescription Medicines**

Alanylglutamine

Amethocaine; for internal use; for external use in medicines containing more than 10%; for ophthalmic use except when used in practice by an optometrist registered with the Optometrists and Dispensing Opticians Board; except when containing 2% or less and used topically as a local anaesthetic in practice by a dental therapist, oral health therapist or dental hygienist (without local anaesthetic exclusion on their scope of practice) registered with the Dental Council

Amivantamab

Arbutin; in oral preparations except herbal preparations containing 500 milligrams or less beta-arbutin per recommended daily dose

Articaine; except when used as a local anaesthetic in practice by a dental therapist, oral health therapist or dental hygienist (without local anaesthetic exclusion on their scope of practice) registered with the Dental Council

Asciminib

Atogepant

Avacopan

Avalglucosidase alfa

Belumosudil

Belzutifan

Benzocaine; except when specified elsewhere in this schedule; except in dermal preparations containing 2% or less total anaesthetic substances; except in lozenges containing 30 milligrams or less of total anaesthetic substances per dosage unit; except when containing 20% or less and used topically as a local anaesthetic in practice by a dental therapist, oral health therapist or dental hygienist (without local anaesthetic exclusion on their scope of practice) registered with the Dental Council

Bilastine; except when specified elsewhere in this schedule

Body Protective Compound -157

NEW ZEALAND GAZETTE

Bufexamac  
Cannabidiol; except when elsewhere in the schedule  
Cardarine  
Cariprazine  
Casirivimab  
Cemiplimab  
Cilgavimab  
Ciltacabtagene autoleucel  
Deucravacitinib  
Deutetrabenazine  
Diroximel fumarate  
Dostarlimab  
Edaravone  
Elexacaftor  
Enfortumab vedotin  
Eslicarbazepine  
Estetrol monohydrate  
Etesevimab  
Faricimab  
Felypressin; except when combined with a local anaesthetic and used in practice by a dental therapist, oral health therapist or dental hygienist (without local anaesthetic exclusion on their scope of practice) registered with the Dental Council  
Fenbendazole  
Filgotinib  
Finerenone  
Fostemsavir  
Fruquintinib  
Ganaxolone  
Glecaprevir; except when supplied in combination with pibrentasvir in a manufacturer's original pack that has received consent from the Minister of Health or Director General for treatment of chronic hepatitis C virus infection to people who meet the clinical and eligibility criteria of an approved training programme, when provided by nurses who meet the requirements of the Nursing Council or pharmacists who meet the requirements of the Pharmacy Council  
Glofitamab  
Hyaluronidase  
Icosapent ethyl  
Imdevimab  
Inclisiran  
Infigratinib  
Lemborexant  
Lenacapavir  
Levomefolic acid; for injection.  
Lignocaine; for injection except when used as a local anaesthetic in practice by a nurse whose scope of practice permits the performance of general nursing functions or by a podiatrist registered with the Podiatry Board or by a dental therapist, oral health therapist or dental hygienist (without local anaesthetic exclusion on their scope of practice) registered with the Dental Council; except when containing 2.5% or less and used topically as a local anaesthetic in practice by a dental therapist, oral health therapist or dental hygienist (without local anaesthetic exclusion on their scope of practice) registered with the Dental Council; for ophthalmic use except when used in practice by an optometrist registered with the Optometrists and Dispensing Opticians Board; for oral use except in throat lozenges in medicines containing 30 milligrams or less per dose form; for external use in medicines containing more than 10%; except in throat sprays in medicines containing 2% or less; except when specified elsewhere in this schedule

NEW ZEALAND GAZETTE

Lurbinectedin

Luspatercept

Mavacamten

Mobocertinib

Molnupiravir; except when specified elsewhere in this schedule

Naloxone; except when provided as part of an approved emergency kit for the treatment of opioid overdose; except when supplied as ampoules with needles and syringes, or as a prefilled syringe, by those authorised by regulation 3, 4 or 5 of the Health (Needles and Syringes) Regulations 1998 for the treatment of opioid overdose, and when supplied with instructions for use

Nirmatrelvir; except when specified elsewhere in this schedule

Nitrofurantoin; except when supplied for oral use containing 100mg per dose unit when sold in a pack of 10 solid dosage units to a woman aged 16–65 years for the first-line empiric treatment of an uncomplicated urinary tract infection by a registered pharmacist who has successfully completed the Pharmaceutical Society of New Zealand training in the treatment of urinary tract infections

Onasemnogene abeparvovec

Opicapone

Osilodrostat

Patisiran

Pegcetacoplan

Pegvaliase

Pemigatinib

Pibrentasvir; except when supplied in combination with glecaprevir in a manufacturer's original pack that has received consent from the Minister of Health or Director General for treatment of chronic hepatitis C virus infection to people who meet the clinical and eligibility criteria of an approved training programme, when provided by nurses who meet the requirements of the Nursing Council or pharmacists who meet the requirements of the Pharmacy Council

Ponesimod

Prilocaine; for injection except when used as a local anaesthetic in practice by a dental therapist, or oral health therapist or dental hygienist (without local anaesthetic exclusion on their scope of practice) registered with the Dental Council; except when specified elsewhere in this schedule except when containing 2.5% or less and used topically as a local anaesthetic in practice by a dental therapist, oral health therapist or dental hygienist (without local anaesthetic exclusion on their scope of practice) registered with the Dental Council

Recombinant varicella zoster virus glycoprotein E antigen; except when administered for the prevention of herpes zoster (shingles) to a person 50 years of age or over by a registered pharmacist who has successfully completed the Vaccinator Foundation Course (or any equivalent training course approved by the Ministry of Health) and who complies with the immunisation standards of the Ministry of Health (but excluding a vaccinator who has completed the Provisional Vaccinator Foundation Course)

Regdanvimab

Relugolix

Ripretinib

Risdiplam

Ritonavir; except when specified elsewhere in this schedule

Sacituzumab govitecan

Selinexor

Selumetinib

Somapacitan

Sotorasib

Sotrovimab

Teneligliptin

Tepotinib

Tirzepatide

Tislelizumab

Tixagevimab

Trabectedin

Trastuzumab deruxtecan

Trifarotene

Vericiguat

Vosoritide

Zanubrutinib

Zinc; except for internal use in medicines containing 25 milligrams or less per recommended daily dose; except for internal use in medicines containing 50 milligrams or less and more than 25 milligrams per recommended daily dose in packs that have received the consent of the Minister or the Director-General to their distribution as general sale medicines, when sold in the manufacturer's original pack and when labelled with a statement that the product may be dangerous if taken in large amounts or for long periods; except for external use except when specified elsewhere in this schedule; except in parenteral nutrition replacement preparations

## Schedule 2

### **Restricted Medicine**

Adrenaline; in medicines containing 1% or less except in medicines for injection containing 0.02% or less; except in medicines for injection containing 0.1% for use in practice in an emergency by a dental therapist, an oral health therapist or a dental hygienist registered with the Dental Council

Cannabidiol; when supplied, in medicines with dosing instructions for 150 milligrams or less per day and containing not more than 4.5 grams, when sold in the manufacturer's original pack that has received consent from the Minister or Director-General, for adults aged 18 years and over, by a registered pharmacist

Choline Salicylate; in medicines containing 10% or less and in packs sizes of 15 grams or less when indicated for use in children under 18 months of age

Ibuprofen 300 milligrams in powder form; for oral use in powder form containing 300 milligrams per dose with a recommended daily dose of not more than 1.2 grams and sold in the manufacturers original packs containing not more than 12 dose units, and labelled for use in adults and children over 12 years of age

Methenamine hippurate

Molnupiravir; for use in the treatment of COVID-19

Nirmatrelvir; for use in the treatment of COVID-19

Pholcodine; in medicines for oral use containing not more than 15 milligrams of pholcodine per solid dosage unit or per dose of liquid with a maximum daily dose not exceeding 100 milligrams of pholcodine, when combined with 1 or more active ingredients in such a way that the substance cannot be recovered by readily applicable means, or in a yield that would constitute a risk to health, when sold in a pack approved by the Minister or the Director-General for distribution as a restricted medicine

Ritonavir; for use in the treatment of COVID-19

## Schedule 3

### **Pharmacy-only Medicines**

Bilastine; for oral use

Folic acid; For oral use in medicines containing more than 500 micrograms per recommended daily dose. When the medicine contains a combination of folic acid, folinic acid or levomefolic acid, the medicine must not provide more than a combined total of 500 micrograms of folic acid, folinic acid and levomefolic acid per maximum recommended daily dose

Folinic acid; For oral use in medicines containing more than 500 micrograms per recommended daily dose. When the medicine contains a combination of folic acid, folinic acid or levomefolic acid, the medicine must not provide more than a combined total of 500 micrograms of folic acid, folinic acid and levomefolic acid per maximum recommended daily dose

Levomefolic acid; for oral use in medicines containing more than 500 micrograms per recommended daily dose. When the medicine contains a combination of folic acid, folinic acid or levomefolic acid, the medicine must not provide more than a combined total of 500 micrograms of folic acid, folinic acid and levomefolic acid per maximum recommended daily dose

Dated this 16th day of April 2024.

DEREK FITZGERALD, Acting Group Manager, Medsafe, Ministry of Health.