



Australian Government

Department of Health and Aged Care
Therapeutic Goods Administration

Australian Register of Therapeutic Goods Certificate

Issued to

Prescribing Biochemists Pty Ltd

for approval to supply

PRESCRIBING BIOCHEMISTS THE MACROMOLECULAR SYSTEM FDP 192 (Cancelled)

ARTG Identifier AUST L 46002
ARTG Start Date 27/07/1993
Product Type Listed Medicine

Manufacturer Details	Address	Manufacturing Steps
Manucom Pty Ltd	270 Lahrs Road ORMEAU , QLD , 42 08 Australia	Release for supply
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ARTG Standard Conditions

The above Medicine Listed has been entered on the Register subject to the following conditions:

- The product is intended for use by practitioners and shall not be supplied directly to the public.
- A copy of a signed and dated certificate of analysis, which is not more than six months old, for the first batch of goods manufactured is to be provided to the Head, Listing Treaties & Export Section, Chemical & Non Prescription Drug Branch within six months of the date of supply of the goods.
- Conditions applicable to all therapeutic goods as specified in the document "Standard Conditions Applying to Registered or Listed Therapeutic Goods Under Section 28 of the Therapeutic Goods Act 1989" effective 1 February 1992.
- Conditions applicable to the relevant category and class of therapeutic goods as specified in the document "Standard Conditions Applying to Registered or Listed Therapeutic Goods Under Section 28 of the Therapeutic Goods Act 1989" effective 1 February 1992.

Products Covered by This Entry

1. PRESCRIBING BIOCHEMISTS THE MACROMOLECULAR SYSTEM FDP 192 (Cancelled)

Product Specific Conditions

No specific conditions have been recorded against this entry.

Product Permitted Indications

No permitted indications have been recorded against this entry.

Product Indication Requirements

No indication requirements have been recorded against this entry.

Product Standard Indications

No standard indications have been recorded against this entry.

Product Specific Indications

- For the relief of gastro-intestinal pain, and as an aid in soothing irritated mucous membranes of the gastro-intestinal tract due to digestive upset.

Warnings

No warnings have been recorded against this entry.

Dosage Form

- Powder, oral

Route of Administration

- Oral

Visual Identification

- A grey/green, gently effervescent powder with characteristic peppermint odour and taste.

Additional Product information

Product Formulation(s)

Active Ingredients

	Quantity	Units
calcium carbonate	265	mg/g
Magnesium carbonate - light	265	mg/g
Ulmus rubra	50	mg/g
Zingiber officinale	10	mg/g
Foeniculum vulgare	10	mg/g
Salix alba	20	mg/g
Glycyrrhiza glabra	40	mg/g

Therapeutic Goods Administration
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