



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

## MACROMOLECULAR SYSTEM BF CHOLINE Tablet (Cancelled)

**ARTG Identifier** AUST L 19354  
**ARTG Start Date** 2/10/1991  
**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:

- Conditions applicable to the relevant category and class of therapeutic goods as specified in the document "Conditions Applying to Therapeutic Goods Accepted for Registration or Listing as Goods Currently Supplied at the Commencement of the Therapeutic Goods Act 1989"

### Products Covered by This Entry

#### 1. MACROMOLECULAR SYSTEM BF CHOLINE Tablet (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

- This product accepted for registration/listing as 'currently supplied' at the time of commencement of the Act. Indications held in ARTG paper records. (Old code)

### Product Specific Indications

No specific indications have been recorded against this entry.

### Warnings

No warnings have been recorded against this entry.

### Dosage Form

- Tablet, film coated

### Route of Administration

- Oral

### Visual Identification

- Pale brown, speckled film coated, round 11.2mm diameter biconvex tablet.

### Additional Product information

**Product Formulation(s)**

**Active Ingredients**

	Quantity	Units
ascorbic acid	125	mg
hesperidin	100	mg
lecithin	200	mg
d-alpha-tocopheryl acid succinate	50	mg

Therapeutic Goods Administration  
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