



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 CIOCHEMISTS CHLORELLIN (Cancelled)**

**ARTG Identifier** AUST L 45914

**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:

- 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 CIOCHEMISTS CHLORELLIN (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 temporary relief of vitamin B6 deficiencies and gastro-intestinal disturbances.

##### **Warnings**

No warnings have been recorded against this entry.

##### **Dosage Form**

- Oral Liquid

##### **Route of Administration**

- Oral

**Visual Identification**

- A dark green/brown, opaque liquid with characteristic orange odour and taste.

**Additional Product information****Product Formulation(s)****Active Ingredients**

	<b>Quantity</b>	<b>Units</b>
<b>Apium graveolens</b>	20	microlitre/mL
<b>pyridoxine hydrochloride</b>	250	microgram/mL

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Therapeutic Goods Administration  
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