



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

HAO697 (Cancelled)

ARTG Identifier AUST L 68914
ARTG Start Date 3/05/1999
Product Type Listed Medicine

Manufacturer Details	Address	Manufacturing Steps
Manucom Pty Ltd	270 Lahrs Road ORMEAU , QLD , 42 08 Australia	Manufacture of dosage form Packaging and labelling Testing microbial Testing chemical and physical 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:

- Except where the sponsor has been contracted by an overseas party to manufacture the goods and that party will be responsible for placing the goods on the market in countries other than Australia, the sponsor must have and shall retain, while the goods remain listed, evidence necessary to substantiate and support the accuracy of the indications in relation to the listed goods, and upon the request of the Head, Office of Prescription Medicines Authorisation Branch, Therapeutic Goods Administration, shall produce such evidence to this officer.
- 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.
- The conditions applying to these goods when they are exported from Australia are given below:
- 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 July 1995, other than condition No. 8 and condition No. 9.
- 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 July 1995, other than condition No.11
- 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 July 1995.
- 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 July 1995.
- The Ginkgo biloba leaf extract used in the manufacture of this medicine must comply with the requirement of Identification Test B of the monograph Powdered Ginkgo Extract in the United States Pharmacopeia 32 - National Formulary 27 (USP32-NF27) as in force or existing from time to time. This condition does not apply to powdered or dried leaf.

Products Covered by This Entry

1. HAO697 (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

- Aids or assists in the maintenance of peripheral circulation. [Or words to that effect] (Old code)
- May assist blood circulation.

Product Specific Indications

- herbal antioxidant, protects capillaries.

Warnings

No warnings have been recorded against this entry.

Dosage Form

- Tablet, film coated

Route of Administration

- Oral

Visual Identification

- Light brown speckled oval biconcave film coated tablet

Additional Product information

Product Formulation(s)

Active Ingredients

	Quantity	Units
Vitis vinifera	30	mg
Equivalent: Vitis vinifera (Dry)	3.6	g
Ginkgo biloba	40	mg
Equivalent: Ginkgo biloba (Dry)	2.4	g
Vaccinium myrtillus	10	mg
Equivalent: Vaccinium myrtillus (Dry)	1	g
d-alpha-tocopheryl acid succinate	85	mg
sodium ascorbate	73.12	mg
ascorbic acid	85	mg
zinc amino acid chelate	37.5	mg
betacarotene	5	mg