



Australian Government
Department of Health
 Therapeutic Goods Administration

Public Summary

Summary for ARTG Entry:	79905	SEAZEST SEA CUCUMBER POWDER 500 mg capsule bottle
ARTG entry for	Medicine Listed (Export Only)	
Sponsor	Reef Organics Pty Ltd	
Postal Address	PO Box 712N, NORTH CAIRNS, QLD, 4870 Australia	
ARTG Start Date	14/08/2001	
Product Category	Medicine	
Status	Active	
Approval Area	Export only Medicines	

Conditions

Except where the sponsor has been contracted by an overseas party to manufacturer 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

Products

1 . SEAZEST SEA CUCUMBER POWDER 500 mg capsule bottle

Product Type	Single Medicine Product	Effective Date	3/07/2002
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Permitted Indications

No Permitted Indications included on Record

Indication Requirements

No Indication Requirements included on Record

Standard Indications

No Standard Indications included on Record

Specific Indications

For the relief of the pain and swelling associated with arthritis and rheumatism.

Warnings

No Warnings included on Record

Additional Product information

Container information

Type	Material	Life Time	Temperature	Closure	Conditions
Bottle	Not recorded	3 Years	Not recorded	Not recorded	Not recorded

Pack Size/Poison information

Pack Size	Poison Schedule
Export	Not scheduled. Not considered by committee

Components

1 . capsule

Dosage Form	Capsule, hard
Route of Administration	Oral

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Visual Identification Greenish-brown coloured powder in clear capsules.

Active Ingredients

Sea cucumber 500 mg

Other Ingredients (Excipients)

colloidal anhydrous silica
Gelatin
magnesium stearate
microcrystalline cellulose
purified water
sodium laureth sulfate

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