Public Summary

Summary for ARTG Entry:	82614 LYMPHOMYOSOT N
ARTG entry for	Medicine Listed
Sponsor	Brauer Professional Pty Ltd
Postal Address	PO Box 174, GLEN OSMOND, SA, 5064 Australia
ARTG Start Date	1/05/2002
Product Category	Medicine
Status	Active
Approval Area	Listed Medicines

Conditions

Colouring agents used in listed medicine for ingestion, other than those listed for export only under section 25 of the Act, shall be only those included in the list of 'Colourings permitted in medicines for oral use'.

The sponsor shall keep records relating to this listed medicine as are necessary to: (a) Expedite recall if necessary of any batch of the listed medicine, (b) Identify the manufacturer(s) of each batch of the listed medicine. Where any part of or step in manufacture in Australia of the listed medicine is sub-contracted to a third party who is not the sponsor, copies of relevant Good Manufacturing Practice agreements relation to such manufacture shall be kept.

The sponsor shall retain records of the distribution of the listed medicine for a period of five years and shall provide the records or copies of the records to the Complementary Medicines Branch, Therapeutic Goods Administration, upon request.

The sponsor of the listed medicine must not, by any means, intentionally or recklessly advertise the medicine for an indication other than those accepted in relation to the inclusion of the medicine in the Register.

All reports of adverse reactions or similar experiences associated with the use or administration of the listed medicine shall be notified to the Head, Office of Product Review, Therapeutic Goods Administration, as soon as practicable after the sponsor of the goods becomes aware of those reports. Sponsors of listed medicines must retain records of such reports for a period of not less than 18 months from the day the Head, Office of Product Review is notified of the report or reports.

The sponsor shall not supply the listed medicine after the expiry date of the goods.

Where a listed medicine is distributed overseas as well as in Australia, product recall or any other regulatory action taken in relation to the medicine outside Australia which has or may have relevance to the quality, safety or efficacy of the goods distributed in Australia, must be notified to the National Manager Therapeutic Goods Administration, immediately the action or information is known to the sponsor.

Products

1 . LYMPHOMYOSOT N

Product Type Single Medicine Product **Effective Date** 12/07/2019

Permitted Indications

Traditionally used in Homoeopathic medicine to aids/assists natural body cleansing/detoxification processes

Traditionally used in Homoeopathic medicine to temporarily relieve mild fluid retention

Traditionally used in Homoeopathic medicine to maintain/support blood circulation/flow to the peripheral areas of the body (legs, hands and feet)

Traditionally used in Homoeopathic medicine to maintain/support healthy lymphatic system

Traditionally used in Homoeopathic medicine to maintain/support healthy immune system function

Indication Requirements

Product presentation must only refer to detoxification in relation to natural body processes.

Product presentation must not imply or refer to cardiovascular or renal conditions.

Product presentation must not imply or refer to serious cardiovascular conditions.

Product presentation must not imply or refer to serious immunological diseases.

Product presentation must not imply or refer to drugs/alcohol.

Product presentation must not imply or refer to disease in any body organ, in particular the kidney or liver.

Label statement: If fluid retention persists, seek medical advice (or words to that effect).

Product presentation must only refer to mild fluid retention.

Standard Indications

No Standard Indications included on Record

Specific Indications

No Specific Indications included on Record

Warnings

Homoeopathic product/preparation or medicine (or words to that effect)

Contains ethanol or contains alcohol.

If fluid retention persists, seek medical advice (or words to that effect).

If symptoms persist consult your healthcare practitioner (or words to that effect).

Additional Product information

Pack Size/Poison information

Pack Size	Poison Schedule

Components

1 . Formulation 1

Dosage Form Liquid, multipurpose

Route of Administration Oral

Visual Identification

Active Ingredients

calcium hydrogen phosphate dihydrate (Homeopathic)	**50 mg/g**
Equisetum hiemale whole plant (Homeopathic)	**50 mg/g**
European garden spider (Homeopathic)	**50 mg/g**
ferrous iodide (Homeopathic)	**100 mg/g**
Fumaria officinalis herb flowering (Homeopathic)	**50 mg/g**
Gentiana lutea root and rhizome (Homeopathic)	**50 mg/g**
Geranium robertianum herb flowering (Homeopathic)	**100 mg/g**
levothyroxine sodium (Homeopathic)	**50 mg/g**
Myosotis arvensis herb flowering (Homeopathic)	**50 mg/g**
Nasturtium officinale herb flowering (Homeopathic)	**100 mg/g**
Pinus sylvestris shoot (Homeopathic)	**50 mg/g**
Scrophularia nodosa herb flowering (Homeopathic)	**50 mg/g**
Smilax aristolochiifolia root and rhizome (Homeopathic)	**50 mg/g**
sodium sulfate (Homeopathic)	**50 mg/g**
Teucrium scorodonia herb flowering (Homeopathic)	**50 mg/g**
Veronica officinalis herb flowering (Homeopathic)	**50 mg/g**

Other Ingredients (Excipients)

ethanol

purified water

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