

Public Summary

Summary for ARTG Entry: 225162 PSC 200

ARTG entry for Medicine Listed

Sponsor Tirsel Ptv Ltd

Postal Address PO Box 1430, FRANKSTON, VIC, 3199

Australia

ARTG Start Date 30/06/2014

Product category Medicine

Status Active

Approval area Listed Medicines

Conditions

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.,

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'.

Products

1. PSC 200

Product Type Single Medicine Product Effective date 30/06/2014

Warnings

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

For practitioner dispensing only.

Standard Indications

Treatment of dry or inflamed skin conditions. [Warning S required],

For the symptomatic relief of dry skin.,

May assist in the management of dry skin.

Specific Indications

No Specific Indications included on Record

Additional Product information

Pack Size/Poison information

Pack Size Poison Schedule

Components

1. Formulation 1

Dosage Form Tablet, film coated

Route of Administration

Visual Identification

Active Ingredients

Allium sativum 50 mg
Equivalent: Allium sativum (Fresh) 1 g

Page 1 of 2

Oral



Arctium lappa	7.5 mg
Equivalent: Arctium lappa (Dry)	30 mg
Artemisia annua	7.5 mg
Equivalent: Artemisia annua (Dry)	30 mg
Astragalus membranaceus	32 mg
Equivalent: Astragalus membranaceus (Dry)	160 mg
Berberis vulgaris	36.7 mg
Equivalent: Berberis vulgaris (Dry)	220.2 mg
Echinacea purpurea	30 mg
Equivalent: Echinacea purpurea (Fresh)	1.35 g
Glycyrrhiza glabra	3 mg
Equivalent: Glycyrrhiza glabra (Dry)	12 mg
Hydrastis canadensis	10 mg
Equivalent: Hydrastis canadensis (Dry)	40 mg
Juglans nigra	230 mg
Panax ginseng	14 mg
Equivalent: Panax ginseng (Dry)	140 mg
Taraxacum officinale	40 mg
Equivalent: Taraxacum officinale (Dry)	160 mg
Thymus vulgaris	9.8 mg
Equivalent: Thymus vulgaris (Dry)	68.6 mg

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