

Australian Government

Department of Health

Therapeutic Goods Administration

Public Summary

Summary for ARTG Entry: 237699 Thuja Cream

ARTG entry for Medicine Listed

Sponsor Martin & Pleasance Pty Ltd

Postal Address PO Box 2007, SOUTH MELBOURNE, VIC, 3205

Australia

ARTG Start Date 14/05/2015
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.

Products

1 . Thuja Cream

Product Type Single Medicine Product Effective Date 22/01/2020

Permitted Indications

Traditionally used in Western herbal medicine to helps decrease/reduce/relieve symptoms of nail mild fungal infection

Traditionally used in Western herbal medicine to decrease/reduce/relieve symptoms of mild, superficial skin fungal infections

Traditionally used in Western herbal medicine to helps reduce/relieve warts

Indication Requirements

Label statement: If symptoms persist, talk to your health professional.

Product presentation must not imply or refer to serious infections.

Product presentation must only refer to mild fungal infection.

Standard Indications

No Standard Indications included on Record

Specific Indications

No Specific Indications included on Record

Warnings

Contains ethanol or contains alcohol.

Additional Product information

Pack Size/Poison information

Pack Size Poison Schedule

Components

1 . Formulation 1

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Produced at 31.08.2021 at 01:53:16 AEST



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Dosage FormCreamRoute of AdministrationTopical

Visual Identification

Active Ingredients

Thuja occidentalis leaf Extract liquid 100 microlitre/g

Equivalent: Thuja occidentalis (Dry) 20 mg/g

Other Ingredients (Excipients)

Carnauba Wax

Castor Oil

cetearyl glucoside

cetostearyl alcohol

citric acid monohydrate

gluconolactone

glycerol

glyceryl caprylate

hydrogenated castor oil

medium chain triglycerides

p-anisic acid

purified water

Rice bran oil

self-emulsifying glyceryl monostearate

sodium hydroxide

sodium stearoyl glutamate

white beeswax

xanthan gum

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