



Australian Government
Department of Health
Therapeutic Goods Administration

Public Summary

Summary for ARTG Entry:	174633	Xuan Fei Tan Chuan Fang A.K.A. Apricot, Perilla & Ginkgo Seed Lung Function I Formula
ARTG entry for	Medicine Listed	
Sponsor	Sun Herbal Pty Ltd	
Postal Address	Unit 5/25 Garema Cct, Kingsgrove, NSW, 2208 Australia	
ARTG Start Date	7/08/2010	
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. Xuan Fei Tan Chuan Fang A.K.A. Apricot, Perilla & Ginkgo Seed Lung Function I Formula

Product Type	Single Medicine Product	Effective Date	10/05/2018
---------------------	-------------------------	-----------------------	------------

Permitted Indications

Traditionally used in Chinese medicine to dispel/expel/extinguish/disperse/clear lung-heat

Traditionally used in Chinese medicine to soothe/descend Qi

Traditionally used in Chinese medicine to disseminate/diffuse lungs/lung-qi

Traditionally used in Chinese medicine to clear/expel/dissolve/resolve Phlegm in/of Lung Dampness - Phlegm Heat pattern

Indication Requirements

Product presentation must not imply or refer to disease in any body organ.

Product presentation must not imply or refer to serious forms of respiratory disorders/diseases, such as: asthma, pneumonia, COAD, COPD, influenza.

If TCM terminology is used on medicine label, label statement: Talk to a TCM practitioner/health professional if you are unsure if this medicine is right for you.

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

Standard Indications

No Standard Indications included on Record

Specific Indications

No Specific Indications included on Record

Warnings

For practitioner dispensing only.

Additional Product information



Australian Government
Department of Health
Therapeutic Goods Administration

Pack Size/Poison information

Pack Size

Poison Schedule

Components

1 . Formulation 1

Dosage Form Capsule, hard

Route of Administration Oral

Visual Identification

Active Ingredients

Aster tataricus root Extract dry concentrate	30.36 mg
Equivalent: Aster tataricus (Dry)	182.16 mg
Biota orientalis branch terminal leafy Extract dry concentrate	40.5 mg
Equivalent: Biota orientalis (Dry)	243 mg
Citrus maxima fruit peel Extract dry concentrate	20.16 mg
Equivalent: Citrus maxima (Dry)	120.96 mg
Ginkgo biloba seed Extract dry concentrate	20.16 mg
Equivalent: Ginkgo biloba (Dry)	120.96 mg
Glycyrrhiza uralensis root Extract dry concentrate	16.86 mg
Equivalent: Glycyrrhiza uralensis (Dry)	101.16 mg
Inula britannica flower Extract dry concentrate	30.36 mg
Equivalent: Inula britannica (Dry)	182.16 mg
Morus alba root bark Extract dry concentrate	30.36 mg
Equivalent: Morus alba (Dry)	182.16 mg
Perilla frutescens seed Extract dry concentrate	30.36 mg
Equivalent: Perilla frutescens (Dry)	182.16 mg
Platycodon grandiflorus root Extract dry concentrate	20.16 mg
Equivalent: Platycodon grandiflorus (Dry)	120.96 mg
Prunus armeniaca seed Extract dry concentrate	30.36 mg
Equivalent: Prunus armeniaca (Dry)	182.16 mg
Scutellaria baicalensis root Extract dry concentrate	30.36 mg
Equivalent: Scutellaria baicalensis (Dry)	182.16 mg

Other Ingredients (Excipients)

Agar

carrageenan

hypromellose

purified water

sodium citrate dihydrate

soluble maize starch

Public Summary

© Commonwealth of Australia. This work is copyright. You are not permitted to re-transmit, distribute or commercialise the material without obtaining prior written approval from the Commonwealth. Further details can be found at <http://www.tga.gov.au/about/website-copyright.htm>.