



Australian Government
Department of Health
Therapeutic Goods Administration

Public Summary

Summary for ARTG Entry:	133404	REISHI GANO (RG) 270 mg
ARTG entry for	Medicine Listed	
Sponsor	DXN International (Australia) Pty Ltd	
Postal Address	Suite 504 / 159-175 Church Street, Level 5 Office Tower / Westfields Shoppingtown, PARRAMATTA, NSW, 2150 Australia	
ARTG Start Date	4/12/2006	
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.

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 . REISHI GANO (RG) 270 mg

Product Type	Single Medicine Product	Effective Date	11/01/2021
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Permitted Indications

Traditionally used in Chinese medicine to help enhance/promote general health and wellbeing

Indication Requirements

No Indication Requirements included on Record

Standard Indications

No Standard Indications included on Record

Specific Indications

No Specific Indications included on Record

Warnings

If symptoms persist consult a medical practitioner (or healthcare professional) (or words to that effect).

Additional Product information

Pack Size/Poison information

Pack Size	Poison Schedule
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Components

1 . Formulation 1

Dosage Form	Capsule, hard
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Route of Administration	Oral
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Visual Identification

Active Ingredients

Ganoderma lucidum fruiting body Powder	270 mg
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Other Ingredients (Excipients)

disodium edetate

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gellan gum
hypromellose
potable water
potassium acetate

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