



Australian Government
Department of Health
Therapeutic Goods Administration

Public Summary

Summary for ARTG Entry:	283961	Mito-Charge Powder
ARTG entry for	Medicine Listed	
Sponsor	RN Labs Pty Ltd	
Postal Address	18 / 93 Rivergate Place, MURARRIE, QLD, 4172 Australia	
ARTG Start Date	22/12/2016	
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 . Mito-Charge Powder

Product Type	Single Medicine Product	Effective Date	29/08/2019
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Permitted Indications

Maintain/support energy levels
Relieve weariness/tiredness/fatigue/feeling of weakness
Helps enhance/promote general health and wellbeing
Maintain/support healthy cardiovascular system function
Aid/assist/helps post exercise recovery
Helps enhance/improve/promote/increase physical/exercise performance
Maintain/support nervous system health
Help establish/restore/reset sleep-wake cycle (circadian rhythm)

Indication Requirements

Label statement: If symptoms persist, talk to your health professional.
Product presentation must not imply or refer to chronic fatigue syndrome.
Product presentation must not imply or refer to mental illnesses, disorders or conditions.
Product presentation must not imply or refer to serious cardiovascular conditions.

Standard Indications

No Standard Indications included on Record

Specific Indications

No Specific Indications included on Record

Warnings

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The onus is on the reader to verify the current accuracy of the information on the document subsequent to the date shown.

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Produced at 23.01.2022 at 11:05:50 AEDT

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Vitamins can only be of assistance if the dietary vitamin intake is inadequate. OR Vitamin supplements should not replace a balanced diet.
(If medicine contains one sugar) contains [insert name of sugar] OR (If medicine contains two or more sugars) Contains sugars [or words to that effect].

Additional Product information

Pack Size/Poison information

Pack Size

Poison Schedule

Components

1 . Formulation 1

Dosage Form Powder

Route of Administration Oral

Visual Identification

Active Ingredients

Biotin	.13 mg/g
calcium pantothenate	8.9 mg/g
Equivalent: pantothenic acid	8 mg/g
calcium pyruvate	89.6 mg/g
Equivalent: calcium	13.5 mg/g
magnesium citrate	85 mg/g
Equivalent: magnesium	13.5 mg/g
nicotinamide	6.7 mg/g
propionyllevocarnitine hydrochloride	134.5 mg/g
ribose	618 mg/g
taurine	6.7 mg/g

Other Ingredients (Excipients)

colloidal anhydrous silica

malic acid

Steviol glycosides

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