

**Australian Government** 

## **Department of Health**

# Therapeutic Goods Administration

## **Public Summary**

Summary for ARTG Entry:	322473	NPM BUFFERED VITAMIN C POWDER WITH HESPERIDIN & MINERALS	
ARTG entry for	Medicine Listed		
Sponsor	PremaLife Pty Ltd t/a Natural Vitality Australia		
Postal Address	11 Aldinga Street, BRENDALE, QLD, 4500 Australia		
ARTG Start Date	29/08/2019		
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.

Effective Date

29/08/2019

#### Products

# **1. NPM BUFFERED VITAMIN C POWDER WITH HESPERIDIN & MINERALS**

Product Type

ct Type Single Medicine Product

Permitted Indications

Maintain/support immune system health

Decrease/reduce/relieve common cold duration

Decrease/reduce/relieve symptoms of common cold

#### Indication Requirements

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

Label statement: Adults only, OR Not to be used in children under 2 years of age without medical advice (or words to that effect).

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

#### **Standard Indications**

No Standard Indications included on Record

#### **Specific Indications**

No Specific Indications included on Record

#### Warnings

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

Adults only. OR Not to be used in children under two years of age without medical advice (or words to that effect).

Vitamins can only be of assistance if the dietary vitamin intake is inadequate. OR Vitamin supplements should not replace a balanced diet.

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#### This is not an ARTG Certificate document.



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**Department of Health** Therapeutic Goods Administration

Additional Product information

Pack Size/Poison information	
Pack Size	Poison Schedule
Components	
1 . Formulation 1	
Dosage Form Powder, oral	
Route of Administration Oral	
Visual Identification	
Active Ingredients	
ascorbic acid	400 mg/g
calcium ascorbate dihydrate	100 mg/g
calcium phosphate	50 mg/g
Equivalent: calcium	19.5 mg/g
hesperidin	95 mg/g
magnesium phosphate pentahydrate	95 mg/g
Equivalent: magnesium	19.625 mg/g
zinc sulfate monohydrate	1.45 mg/g
Equivalent: zinc	529 microgram/g
Other Ingredients (Excipients)	
Flavour	
fructose	
glucose monohydrate	
silicon dioxide	
Stevia rebaudiana	

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