



Australian Government
Department of Health
Therapeutic Goods Administration

Public Summary

Summary for ARTG Entry:	160383	Zinc Forte + C
ARTG entry for	Medicine Listed	
Sponsor	Herbs of Gold Pty Ltd	
Postal Address	PO Box 3143, KIRRAWEE, NSW, 2232 Australia	
ARTG Start Date	24/03/2009	
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 . Zinc Forte + C

Product Type	Single Medicine Product	Effective Date	3/09/2019
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Permitted Indications

- Antioxidant/Reduce free radicals formed in the body
- Antioxidant/Reduce free radicals formed in the body in children
- Maintain/support collagen formation
- Maintain/support collagen formation in children
- Maintain/support collagen health in children
- Maintain/support collagen health
- Maintain/support general health and wellbeing in children
- Maintain/support connective tissue health in children
- Maintain/support connective tissue health
- Aid/assist/helps connective tissue production/formation in children
- Aid/assist/helps connective tissue production/formation
- Maintain/support bone health in children
- Maintain/support bone health
- Maintain/support taste sensation
- Maintain/support taste sensation in children
- Maintain/support immune system health
- Maintain/support immune system health in children
- Maintain/support healthy immune system function
- Maintain/support healthy immune system function in children



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- Maintain/support immune system to fight illness in children
- Maintain/support absorption of dietary (state vitamin/mineral/nutrient) in children
- Maintain/support absorption of dietary (state vitamin/mineral/nutrient)
- Maintain/support sperm health
- Maintain/support sperm motility
- Maintain/support skin health
- Maintain/support skin health in children
- Maintain/support wound healing in children
- Maintain/support wound healing

Indication Requirements

Product presentation must not imply or refer to bone disease or disorders e.g. rheumatoid arthritis, juvenile arthritis, debilitating osteoarthritis, osteoporosis. Note: this requirement is not intended to apply where the indications referring to osteoporosis specified in column 2 of Table 2 of this instrument are also used.

Product presentation must not imply or refer to infertility.

Product presentation must not imply or refer to serious immunological diseases.

Label statement: [Vitamins/minerals/nutrients/dietary supplements] can only be of assistance if dietary intake is inadequate OR [Vitamins/minerals/nutrients/dietary supplements] should not replace a balanced diet (or words to that effect).

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

WARNING: May be dangerous if taken in large amounts or for a long period. OR WARNING: Contains zinc which may be dangerous if taken in large amounts or for a long period (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	Powder, oral
Route of Administration	Oral

Visual Identification

Active Ingredients

betacarotene	56.67 microgram/g
calcium ascorbate dihydrate	333.33 mg/g
Equivalent: ascorbic acid	275.4 mg/g
magnesium amino acid chelate	8.35 mg/g
Equivalent: magnesium	1.67 mg/g
pyridoxine hydrochloride	2 mg/g
Equivalent: pyridoxine	1.66 mg/g
zinc citrate dihydrate	31.15 mg/g
Equivalent: zinc	10 mg/g

Other Ingredients (Excipients)

- citric acid
- colloidal anhydrous silica
- dl-alpha-tocopherol
- Flavour
- iron oxide red

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isomalt
purified water
starch sodium octenyl succinate
Stevia rebaudiana

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