

Australian Government

Department of Health

Therapeutic Goods Administration

Public Summary

Summary for ARTG Entry: MolyZinc 338074

ARTG entry for Medicine Listed

Sponsor Interclinical Laboratories Pty Ltd

Postal Address PO Box 6474, ALEXANDRIA, NSW, 2015

18/06/2020 ARTG Start Date **Product Category** Medicine Status Active

Listed Medicines Approval Area

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.

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. MolyZinc

Product Type Effective Date 18/06/2020 Single Medicine Product

Permitted Indications

Antioxidant/Reduce free radicals formed in the body

Maintain/support collagen formation

Maintain/support energy levels

Maintain/support healthy eye function

Maintain/support eve health

Maintain/support healthy eyesight/vision

Maintain/support general health and wellbeing

Maintain/support hair health

Maintain/support healthy teeth

Maintain/support nail health/strength/thickness

Maintain/support connective tissue health

Aid/assist/helps connective tissue production/formation

Maintain/support bone health

Aid/assist healthy red blood cell production

Helps maintain/support haemoglobin formation/synthesis

Maintain/support blood capillary health

Maintain/support blood vessel health

Maintain/support immune system health

Maintain/support healthy immune system function

Aid/assist/helps protein synthesis in the body

Maintain/support absorption of dietary (state vitamin/mineral/nutrient)

Helps prevent dietary (state vitamin/mineral/nutrient) deficiency

Aid/assist/helps metabolism of (state vitamin/mineral/nutrient)

Page 1 of 3



Australian Government

Department of Health

Therapeutic Goods Administration

Maintain/support cognitive function/mental function in healthy adults

Maintain/support nerve conduction

Aid/assist/helps synthesis of neurotransmitters

Maintain/support nervous system health

Maintain/support nervous system function

Maintain/support female reproductive system health

Maintain/support preconception health in healthy females

Maintain/support preconception health in healthy males

Maintain/support reproductive system health in males

Maintain/support healthy reproductive hormones

Maintain/support sperm health in healthy males

Maintain/support sperm production in healthy males

Maintain/support testosterone level

Maintain/support skin health

Maintain/support skin integrity/structure

Maintain/support skin regeneration

Maintain/support wound healing

Indication Requirements

Product presentation must not imply or refer to serious cardiovascular conditions.

Product presentation must not imply or refer to circulatory disorders/diseases/conditions e.g. thrombosis.

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

Product presentation must not imply or refer to mental illnesses, disorders or conditions.

Product presentation must not imply or refer to hormone imbalances.

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.

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 chronic fatigue syndrome.

If directed to women, Label statement: Advise your doctor of any medicine you take during pregnancy, particularly in your first trimester.

Standard Indications

No Standard Indications included on Record

Specific Indications

No Specific Indications included on Record

Warnings

Vitamins and minerals can only be of assistance if dietary intake is inadequate OR Vitamin and/or mineral supplements should not replace a balanced diet.

Additional Product information

Pack Size/Poison information

Pack Size Poison Schedule

Components

1 . Formulation 1

Dosage Form Capsule, hard

Route of Administration Oral

Visual Identification

Active Ingredients

ascorbic acid 30 mg

molybdenum trioxide150 microgramEquivalent: molybdenum100 microgram

Page 2 of 3

Produced at 31.08.2021 at 04:50:08 AEST



Australian Government

Department of Health

Therapeutic Goods Administration

pyridoxal 5-phosphate monohydrate1.61 mgEquivalent: pyridoxine1.03 mgzinc glycinate monohydrate17.7 mgEquivalent: zinc5 mg

Other Ingredients (Excipients)

calcium hydrogen phosphate dihydrate colloidal anhydrous silica hypromellose magnesium stearate microcrystalline cellulose

© 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.