



Australian Government
Department of Health and Aged Care
 Therapeutic Goods Administration

Public Summary

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|--------------------------------|---|----------------------|
| Summary for ARTG Entry: | 282024 | BioActivated Calcium |
| ARTG entry for | Medicine Listed | |
| Sponsor | Biomedica Nutraceuticals Pty Ltd | |
| Postal Address | PO Box 7052, ALEXANDRIA, NSW, 2015 Australia | |
| ARTG Start Date | 2/11/2016 | |
| 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.

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 . BioActivated Calcium

| | | | |
|---------------------|-------------------------|-----------------------|------------|
| Product Type | Single Medicine Product | Effective Date | 18/11/2022 |
|---------------------|-------------------------|-----------------------|------------|

Permitted Indications

- Maintain/support collagen formation
- Maintain/support general health and wellbeing
- Maintain/support healthy teeth
- Maintain/support bone health
- Aids/assists healthy bone development/growth/building
- Maintain/support bone strength
- Help maintain/support bone mineralisation
- A diet deficient in calcium can lead to osteoporosis in later life. Calcium may help prevent osteoporosis when dietary intake is inadequate
- Vitamin D helps calcium absorption (or words of like intent) and a diet deficient in calcium can lead to osteoporosis in later life
- Helps maintain/support joint cartilage health
- Antispasmodic/spasmolytic
- Maintain/support healthy muscle contraction function
- Maintain/support muscle health
- Maintain/support muscle function
- Helps prevent dietary (state vitamin/mineral/nutrient) deficiency
- Maintain/support (state vitamin/mineral) within normal range

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 any form of arthritis or osteoarthritis unless qualified as mild.

Product presentation must not imply or refer to serious musculoskeletal or neurological conditions.

Indication can only be used for medicines that contain calcium as an active ingredient and the recommended daily dose of the medicine must provide at least 290 milligrams of elemental calcium.

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).

Indication only for use for medicines that contain vitamin D as an active ingredient. The medicines may only contain a maximum recommended daily dose

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of 25 micrograms or less of vitamin D and as a minimum, also contain at least 25% of the RDI in the recommended daily dose of vitamin D.
 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

No Warnings included on Record

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

| | |
|-------------------------------------|------------------------|
| ascorbic acid | 9.58 mg/g |
| borax | 2.205 mg/g |
| Equivalent: boron | 250 microgram/g |
| calcium citrate | 197.47 mg/g |
| Equivalent: calcium | 41.67 mg/g |
| calcium phosphate | 157.66 mg/g |
| Equivalent: calcium | 58.3 mg/g |
| colecalfiferol | .0016 mg/g |
| colloidal anhydrous silica | 7.13 mg/g |
| Equivalent: silicon | 3.33 mg/g |
| hydroxyapatite | 33.333 mg/g |
| Equivalent: calcium | 8.33 mg/g |
| lysine hydrochloride | 41.65 mg/g |
| Equivalent: lysine | 33.34 mg/g |
| magnesium citrate | 36.1 mg/g |
| Equivalent: magnesium | 5.83 mg/g |
| magnesium phosphate tribasic | 165 mg/g |
| Equivalent: magnesium | 34.17 mg/g |
| manganese amino acid chelate | 4.17 mg/g |
| Equivalent: manganese | 417 microgram/g |
| phytomenadione | 6.7 microgram/g |
| zinc citrate dihydrate | 3.11 mg/g |
| Equivalent: zinc | 1 mg/g |

Other Ingredients (Excipients)

- Acacia
- dl-alpha-tocopherol
- Flavour
- glycine
- Guar Gum
- maize starch
- maltodextrin

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medium chain triglycerides

rice starch

silicon dioxide

sucrose

thaumatin

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