Australian Government

Department of Health
Therapeutic Goods Administration

Public Summary

Summary for ARTG Entry:	172165	BLOOMS MSM 750 WITH GINGER & DEVIL'S CLAW

ARTG entry for	Medicine Listed
Sponsor	Phytologic Holdings Pty Limited
Postal Address	PO Box 6193, Alexandria, NSW, 2015 Australia
ARTG Start Date	10/06/2010
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 . BLOOMS MSM 750 WITH GINGER & DEVIL'S CLAW

Product Type	Single Medicine Product	**Effective Date**	12/02/2021

Permitted Indications

Maintain/support general health and wellbeing

Anti-inflammatory/relieve inflammation

Analgesic/Anodyne/relieve pain

Decrease/reduce/relieve mild rheumatic aches and pains

Decrease/reduce/relieve symptoms of mild arthritis/mild osteoarthritis

- Linked indication - Decrease/reduce/relieve mild joint aches and pains
- Linked indication - Decrease/reduce/relieve mild joint stiffness
- Linked indication - Decrease/reduce/relieve mild joint pain/soreness
- Linked indication - Decrease/reduce/relieve mild joint inflammation/swelling

Helps maintain/support joint cartilage health

Decrease/reduce/relieve mild joint inflammation/swelling

Maintain/support joint mobility/flexibility

Decrease/reduce/relieve mild joint stiffness

Maintain/support healthy digestion

Decrease/reduce/relieve symptoms of indigestion/dyspepsia

Relieve digestive discomfort

Maintain/support immune system health

Maintain/support healthy immune system function

Decrease/reduce/relieve muscle pain/ache/soreness after exercise

Aid/assist/helps post exercise recovery

Maintain/support skin health

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 only refer to mild joint symptoms.

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.

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 immunological diseases.

Product presentation must only refer to mild rheumatic aches/pains.

Product presentation must not imply or refer to gastro oesophageal reflux disease.

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	Capsule, hard
Route of Administration	Oral
Visual Identification	

Active Ingredients

dimethyl sulfone	**750 mg**
Harpagophytum procumbens root Extract dry concentrate standardised	**10 mg**
Equivalent: Harpagophytum procumbens (Dry)	25 mg
Zingiber officinale root Extract dry concentrate	**40 mg**
Equivalent: Zingiber officinale (Dry)	1 g

Other Ingredients (Excipients)

hypromellose
magnesium stearate
maltodextrin
purified water
silicon dioxide
sorbitol