Public Summary

Summary for ARTG Entry:	328696 MULTIBIOTIC
ARTG entry for	Medicine Listed
Sponsor	Medlab Pty Ltd
Postal Address	PO Box 6452, ALEXANDRIA, NSW, 2015 Australia
ARTG Start Date	16/01/2020
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 . MULTIBIOTIC

Product Type Single Medicine Product **Effective Date** 16/01/2020

Permitted Indications

Decrease/reduce/relieve diarrhoea in children

Relief of symptoms of medically diagnosed Irritable Bowel Syndrome

Help reduce occurrence of symptoms of medically diagnosed Irritable Bowel Syndrome

Maintain/support intestinal health

Maintain/support digestive system health

Helps restore good/beneficial/friendly intestinal/gut/bowel flora

Decrease/reduce/relieve abdominal pain/discomfort

Maintain/support healthy immune system function

Maintain/support skin health

Indication Requirements

Product presentation must only refer to medically diagnosed IBS.

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

Label statement: Seek medical advice if diarrhoea persists for more than: 6 hours in infants under 6 months, 12 hours in children under 3 years, 24 hours in children aged 3 to 6 years or 48 hours in adults and children over 6 years (or words to that effect).

Label statement: If symptoms persist, worsen or episodes become more frequent talk to your medical practitioner.

Label statement: If symptoms persist or worsen talk to your medical practitioner.

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

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

If diarrhoea persists for more than 6 hours in infants under 6 months, 12 hours in children under 3 years, 24 hours in children aged 3-6 years or 48 hours in adults and children over 6 years, seek medical advice (or words to that effect).

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

Additional Product information

Container information

Type	Material	Life Time	Temperature	Closure	Conditions
Multiple containers	Not recorded	Not recorded	Not recorded	Not recorded	Not recorded

Pack Size/Poison information

Pack Size	Poison Schedule

Components

1 . Formulation 1

Dosage Form	Capsule, hard
Route of Administration	Oral
Visual Identification	

Active Ingredients

Bifidobacterium animalis ssp lactis	**3 billion CFU**
Bifidobacterium bifidum	**500 million CFU**
Bifidobacterium breve	**1.75 billion CFU**
Lactobacillus acidophilus	**3.75 billion CFU**
Lactobacillus plantarum	**1.575 billion CFU**
Lactobacillus rhamnosus	**9 billion CFU**
Streptococcus thermophilus	**1.5 billion CFU**

Other Ingredients (Excipients)

colloidal anhydrous silica
disodium edetate
gellan gum
hypromellose
magnesium stearate
maltodextrin
potable water
potassium acetate

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