



Australian Government
Department of Health
Therapeutic Goods Administration

Public Summary

Summary for ARTG Entry:	355433	ACTIVATED PROBIOTICS BIOME ECZEMA
ARTG entry for	Medicine Listed	
Sponsor	Biome Australia Limited	
Postal Address	16 Dover Street, Cremorne, VIC, 3121 Australia	
ARTG Start Date	23/02/2021	
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 . ACTIVATED PROBIOTICS BIOME ECZEMA

Product Type	Single Medicine Product	Effective Date	23/02/2021
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Permitted Indications

- Decrease/reduce/relieve itchy/prickling skin associated with mild eczema/dermatitis
- Decrease/reduce/relieve symptoms of mild eczema/dermatitis
- Helps reduce occurrence of symptoms of eczema/dermatitis

Indication Requirements

- Label statement: If symptoms persist, talk to your health professional.
- Product presentation must only refer to mild eczema.

Standard Indications

No Standard Indications included on Record

Specific Indications

No Specific Indications included on Record

Warnings

If symptoms persist, seek the advice of a healthcare professional.

Additional Product information

Container information

Type	Material	Life Time	Temperature	Closure	Conditions
Sachet	Not recorded	Not recorded	Not recorded	Neither child resistant closure nor restricted flow insert	Not recorded

Pack Size/Poison information

Pack Size	Poison Schedule
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Components

- 1 . Formulation 1

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Dosage Form Powder, oral

Route of Administration Oral

Visual Identification

Active Ingredients

Lactobacillus salivarius ssp salivarius 1 billion organisms/g

Other Ingredients (Excipients)

maltodextrin

silicon dioxide

sorbitol

Vanilla Powder

xylitol

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