

Australian Government

## **Department of Health and Aged Care**

Therapeutic Goods Administration

## **Public Summary**

Summary for ARTG Entry:	316500	ACTIVATED PROBIOTICS BIOME PRENATAL+
ARTG entry for	Medicine Listed	
Sponsor	Biome Australia I	limited
Postal Address	192/194 Johnsto Australia	n Street, Collingwood, VIC, 3066
ARTG Start Date	15/04/2019	
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

Product Type	Single Medicine Pro	oduct	Effective Date	19/10/2023	
Permitted Indica	tions				
Helps maintains/s	upport healthy foetal CNS	/brain development			
Enhance/promote	healthy foetal developme	nt			
Maintain/support	healthy pregnancy				
Helps enhance/pr	omote maternal health				
Help to prevent ne	eural tube defects such as	spina bifida and/or ane	ncephaly		
Helps enhance/pr	omote preconception heal	th			
Indication Requi	rements				
Product presenta	tion must not imply or refe	r to infertility.			
If directed to won	nen, Label statement: Adv	se your doctor of any m	nedicine you take during pre	gnancy, particularly in yo	our first trimester.
Label statement:	If you are concerned about	it the health of yourself	or your baby, talk to your he	alth practitioner.	
Indication can on		nat contain folic acid as	an active ingredient and the	recommended daily dos	se of the medicine provides a
minimum of 400	when trying to conceive a				nclude at least one of the followin veeks before conception and durin
minimum of 400 label statements: the first trimester	when trying to conceive a of pregnancy.				
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closure nor restricted flow insert

Pack Size/Poison information	n	
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 breve	•	5 billion CFU
colecalciferol		10 microgram
cyanocobalamin		2.6 microgram
folic acid		.5 mg
Lactobacillus crispatus		1 billion CFU
Lactobacillus fermentum		1 billion CFU
Lactobacillus plantarum		5 billion CFU
Lactobacillus rhamnosus		5 billion CFU
potassium iodide		.2 mg
Equivalent: iodine		150 microgram
Other Ingredients (Excipie	ents)	
glyceryl palmito-stearate		
hypromellose		
maltodextrin		
silicon dioxide		

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