



**Australian Government**  
**Department of Health**  
Therapeutic Goods Administration

## Public Summary

<b>Summary for ARTG Entry:</b>	335200	Pro8-50 Plus
<b>ARTG entry for</b>	Medicine Listed	
<b>Sponsor</b>	Spectrumceuticals Pty Ltd	
<b>Postal Address</b>	10/5 Narabang Way, BELROSE, NSW, 2085 Australia	
<b>ARTG Start Date</b>	25/04/2020	
<b>Product Category</b>	Medicine	
<b>Status</b>	Active	
<b>Approval Area</b>	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.

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 . Pro8-50 Plus

<b>Product Type</b>	Single Medicine Product	<b>Effective Date</b>	25/04/2020
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#### Permitted Indications

Maintain/support general health and wellbeing  
Maintain/support intestinal health  
Maintain/support intestinal good/beneficial/friendly flora  
Maintain/support small intestine good/beneficial/friendly flora  
Maintain/support gastrointestinal system health  
Maintain/support gastrointestinal mucosal membrane health  
Maintain/support immune system health  
Maintain/support healthy immune system function  
Maintain/support healthy gastrointestinal immune function

#### Indication Requirements

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

#### 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

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Pack Size

Poison Schedule

**Components**

**1 . Formulation 1**

**Dosage Form** Capsule, hard

**Route of Administration** Oral

**Visual Identification**

**Active Ingredients**

<b>Bifidobacterium breve</b>	<b>1.35 billion CFU</b>
<b>Bifidobacterium lactis</b>	<b>4.05 billion CFU</b>
<b>Bifidobacterium longum</b>	<b>.45 billion CFU</b>
<b>Lactobacillus acidophilus</b>	<b>7.4 billion CFU</b>
<b>Lactobacillus casei</b>	<b>9.45 billion CFU</b>
<b>Lactobacillus gasseri</b>	<b>1.35 billion CFU</b>
<b>Lactobacillus plantarum</b>	<b>3.15 billion CFU</b>
<b>Lactobacillus rhamnosus</b>	<b>30 billion CFU</b>

**Other Ingredients (Excipients)**

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

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