



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

## Public Summary

|                                |   |                                  |
|--------------------------------|---|----------------------------------|
| <b>Summary for ARTG Entry:</b> | 385864  | ETHICAL NUTRIENTS ST JOHN'S WORT |
| <b>ARTG entry for</b>          | Medicine Listed                                 |                                  |
| <b>Sponsor</b>                 | Metagenics (Aust) Pty Ltd                       |                                  |
| <b>Postal Address</b>          | PO Box 675, VIRGINIA BC, QLD, 4014<br>Australia |                                  |
| <b>ARTG Start Date</b>         | 18/03/2022                                      |                                  |
| <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.

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 . ETHICAL NUTRIENTS ST JOHN'S WORT

|                     |                         |                       |            |
|---------------------|-------------------------|-----------------------|------------|
| <b>Product Type</b> | Single Medicine Product | <b>Effective Date</b> | 18/03/2022 |
|---------------------|-------------------------|-----------------------|------------|

#### Permitted Indications

Support healthy stress response in the body  
Decrease/reduce/relieve restlessness/excess nervous energy  
Decrease/reduce/relieve symptoms of stress  
Linked indication - Support healthy emotional/mood balance  
Decrease/reduce/relieve nervous tension/unrest  
Support healthy emotional/mood balance

#### Indication Requirements

Product presentation must not imply or refer to mental illnesses, disorders or conditions.  
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

St John's Wort affects the way many prescription medicines work, including the oral contraceptive pill. Consult your doctor.  
If symptoms persist consult your healthcare practitioner (or words to that effect).

#### Additional Product information

#### Pack Size/Poison information

|                  |                        |
|------------------|------------------------|
| <b>Pack Size</b> | <b>Poison Schedule</b> |
|------------------|------------------------|

#### Components

##### 1 . Formulation 1

**Dosage Form** Capsule, hard  
**Route of Administration**



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**Department of Health**  
Therapeutic Goods Administration

Oral

**Visual Identification**

**Active Ingredients**

|   |               |
|---|---------------|
| <b>Hypericum perforatum herb top flowering Extract dry concentrate standardised</b> | <b>300 mg</b> |
| Equivalent: Hypericum perforatum (Dry)  | 1.725 g       |

**Other Ingredients (Excipients)**

chlorophyllin-copper complex  
colloidal anhydrous silica  
disodium edetate  
gellan gum  
hypromellose  
magnesium stearate  
microcrystalline cellulose  
potassium acetate  
purified water  
sorbitan monolaurate

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