Public Summary

Summary for ARTG Entry:	68615	NERVOUS SYSTEM NERVE TONIC
ARTG entry for	Medicine Listed	
Sponsor	Brauer Natural Medicine Pty Ltd	
Postal Address	PO Box 234, TANUNDA, SA, 5352 Australia	
ARTG Start Date	8/04/1999	
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'.

Sponsors must confirm the absence of aristolochic acids, in all medicines containing herbal material derived from any of the following plant genera - Akebia, Asarum, Bragantia, Clematis, Cocculus, Diploclisia, Menispermum, Saussurea, Sinomenium, Stephania, Vladimiria.
The confirmation must be undertaken by chemical analysis using Liquid Chromatography Mass Spectrometry (LC-MS). The methodology used should adhere to best practice according to contemporary scientific literature.
Confirmatory evidence is to be provided to the Director of Listing Compliance, Complementary and OTC Medicines Branch, prior to supply of each batch in Australia. The evidence submitted to the TGA is to include the certificate of analysis, all relevant details of the methodology, such as analytical method validation data, and the raw results.
All supporting evidence must be approved by the TGA prior to supply of the batch in Australia.

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 . NERVOUS SYSTEM NERVE TONIC

Product Type	Single Medicine Product	**Effective Date**	26/08/2019

Permitted Indications

Traditionally used in Homoeopathic medicine to decrease/reduce/relieve symptoms of stress

Traditionally used in Homoeopathic medicine to decrease/reduce/relieve nervous tension/unrest

Traditionally used in Homoeopathic medicine to decrease/reduce/relieve symptoms of mild anxiety

Traditionally used in Homoeopathic medicine to nerve tonic

Traditionally used in Homoeopathic medicine to soporific/induces sleep

Traditionally used in Homoeopathic medicine to decrease/reduce/relieve sleeplessness

Indication Requirements

Label statement: If symptoms persist, talk to your health professional.

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

Product presentation must only refer to mild anxiety.

Standard Indications

No Standard Indications included on Record

Specific Indications

No Specific Indications included on Record

Warnings

Homoeopathic product/preparation or medicine (or words to that effect)

Contains ethanol or contains alcohol.

Contains sorbates' (or word to this effect) if medicine contains two or more sorbate sources OR 'Contains [insert the approved name of sorbate source used]' (or words to this effect) if medicine contains one sorbate source.

(If medicine contains one sugar) contains [insert name of sugar] OR (If medicine contains two or more sugars) Contains sugars [or words to that effect].

Additional Product information

Pack Size/Poison information

Pack Size	Poison Schedule

Components

1 . Formulation 1

Dosage Form	Oral Liquid
Route of Administration	Oral
Visual Identification	

Active Ingredients

Anamirta cocculus fruit (Homeopathic)	**50 nanolitre/mL**
Equivalent: Picrotoxin	25 ng/mL
Chamaelirium luteum root and rhizome (Homeopathic)	**50 nanolitre/mL**
Passiflora incarnata herb flowering (Homeopathic)	**5 microlitre/mL**
phosphoric acid (Homeopathic)	**60 nanolitre/mL**
sepia (Homeopathic)	**50 nanolitre/mL**
silver nitrate (Homeopathic)	**1.67 microlitre/mL**
Strychnos ignatii seed (Homeopathic)	**5 microlitre/mL**
Equivalent: strychnine (of Strychnos spp.)	1.8 ng/mL
Strychnos nux-vomica seed (Homeopathic)	**1.67 microlitre/mL**
Equivalent: strychnine (of Strychnos spp.)	0 picogram/mL
zinc (Homeopathic)	**50 nanolitre/mL**

Other Ingredients (Excipients)

ascorbic acid
ethanol absolute
potassium sorbate
purified water
Ribes nigrum
Rosa canina
sucrose

Public Summary