Public Summary

Summary for ARTG Entry:	82616	RENEEL
ARTG entry for	Medicine Listed	
Sponsor	Brauer Professional Pty Ltd	
Postal Address	PO Box 174, GLEN OSMOND, SA, 5064 Australia	
ARTG Start Date	1/05/2002	
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.

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 . RENEEL

Product Type Single Medicine Product **Effective Date** 10/07/2019

Permitted Indications

Traditionally used in Homoeopathic medicine to aids/assists natural kidney cleansing/detoxification processes

Traditionally used in Homoeopathic medicine to maintain/support kidney function

Traditionally used in Homoeopathic medicine to kidney Tonic

Indication Requirements

Product presentation must only refer to detoxification in relation to natural body processes.

Product presentation must not imply or refer to kidney disease.

Product presentation must not imply or refer to drugs/alcohol.

Product presentation must not imply or refer to disease in any body organ, in particular the kidney or liver.

Standard Indications

No Standard Indications included on Record

Specific Indications

No Specific Indications included on Record

Warnings

Contains ethanol or contains alcohol.

Homoeopathic product/preparation or medicine (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

Berberis vulgaris root bark (Homeopathic)	**100 mg/g**
Cantharides (Homeopathic)	**100 mg/g**
Equisetum hiemale whole plant (Homeopathic)	**100 mg/g**
Populus tremuloides stem bark inner (Homeopathic)	**50 mg/g**
Populus tremuloides leaf (Homeopathic)	**50 mg/g**
Serenoa repens fruit (Homeopathic)	**100 mg/g**
Solidago virgaurea flower (Homeopathic)	**100 mg/g**

Other Ingredients (Excipients)

ethanol absolute
purified water