



**Australian Government**  
**Department of Health and Aged Care**  
Therapeutic Goods Administration

## Public Summary

<b>Summary for ARTG Entry:</b>	350474	Pain Relief PEA
<b>ARTG entry for</b>	Medicine Listed	
<b>Sponsor</b>	Herbs of Gold Pty Ltd	
<b>Postal Address</b>	PO Box 3143, KIRRAWEE, NSW, 2232 Australia	
<b>ARTG Start Date</b>	2/12/2020	
<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.

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 . Pain Relief PEA

<b>Product Type</b>	Single Medicine Product	<b>Effective Date</b>	2/12/2020
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#### Permitted Indications

Analgesic/Anodyne/relieve pain  
Decrease/reduce/relieve mild joint aches and pains  
Decrease/reduce/relieve symptoms of mild arthritis/mild osteoarthritis  
Decrease/reduce/relieve mild joint pain/soreness  
Decrease/reduce/relieve mild nerve pain/neuralgia

#### Indication Requirements

Label statement: If symptoms persist, talk to your health professional.  
Product presentation must not imply or refer to bone disease or disorders e.g. rheumatoid arthritis, juvenile arthritis, debilitating osteoarthritis, osteoporosis.  
Product presentation must only refer to mild joint symptoms.  
Product presentation must only refer to mild nerve pain/neuralgia.

#### Standard Indications

No Standard Indications included on Record

#### Specific Indications

No Specific Indications included on Record

#### Warnings

Not to be used for more than 21 consecutive days.  
Adults only.  
The medicine may interact with other prescription analgesic medicines, please consult your healthcare practitioner before use.

#### Additional Product information

#### Pack Size/Poison information

<b>Pack Size</b>	<b>Poison Schedule</b>
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#### Components

1 . Formulation 1



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**Dosage Form** Capsule, hard

**Route of Administration** Oral

**Visual Identification**

**Active Ingredients**

**palmidrol** 149.9953 mg

**Other Ingredients (Excipients)**

calcium hydrogen phosphate dihydrate

colloidal anhydrous silica

disodium edetate

dl-alpha-tocopheryl acetate

gellan gum

hypromellose

lecithin

Lime Oil Coldpressed

magnesium stearate

medium chain triglycerides

microcrystalline cellulose

Olive Oil

PEG-35 castor oil

potable water

potassium acetate

silicon dioxide

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Public Summary