



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

## Public Summary

<b>Summary for ARTG Entry:</b>	222465	METAGENICS RESVERATROL HEALTHY AGEING COMPLEX
<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 Australia	
<b>ARTG Start Date</b>	14/04/2014	
<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.

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 . METAGENICS RESVERATROL HEALTHY AGEING COMPLEX

<b>Product Type</b>	Single Medicine Product	<b>Effective Date</b>	29/01/2020
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### Permitted Indications

- Antioxidant/Reduce free radicals formed in the body in elderly individuals
- Antioxidant/Reduce free radicals formed in the body
- Helps reduce/decrease free radical damage to body cells
- Maintain/support general health and wellbeing in aging individuals
- Traditionally used in Chinese medicine to maintain/support healthy blood circulation
- Helps maintain/support healthy blood sugar/glucose in healthy individuals
- Maintain/support cardiovascular system health in healthy individuals
- Maintain/support healthy cardiovascular system function in elderly individuals
- Maintain/support healthy cardiovascular system function in healthy individuals
- Traditionally used in Chinese medicine to maintain/support healthy immune system function
- Maintain/support healthy immune system function in healthy individuals

### Indication Requirements

- Product presentation must not imply or refer to serious cardiovascular conditions.
- Product presentation must not imply or refer to serious immunological diseases.
- Product presentation must not imply or refer to lowering or raising blood sugar/glucose levels from outside of the normal healthy range.

### Standard Indications

No Standard Indications included on Record

### Specific Indications



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Therapeutic Goods Administration

No Specific Indications included on Record

**Warnings**

If symptoms persist consult your healthcare practitioner (or words to that effect).

**Additional Product information**

**Pack Size/Poison information**

**Pack Size**

**Poison Schedule**

**Components**

**1 . Formulation 1**

**Dosage Form**                      Tablet, film coated

**Route of Administration**      Oral

**Visual Identification**

**Active Ingredients**

<b>Curcuma longa root Extract dry concentrate</b>	<b>152 mg</b>
Equivalent: Curcuma longa (Dry)	3.8 g
<b>quercetin</b>	<b>200 mg</b>
<b>Reynoutria japonica root Extract dry concentrate standardised</b>	<b>600 mg</b>
Equivalent: Reynoutria japonica (Dry)	60 g

**Other Ingredients (Excipients)**

Carnauba Wax  
chlorophyllin-copper complex  
colloidal anhydrous silica  
croscarmellose sodium  
hypromellose  
macrogol 8000  
magnesium stearate  
microcrystalline cellulose  
sodium starch glycollate type A

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