



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

## Public Summary

<b>Summary for ARTG Entry:</b>	203612	CardioNutrients Forte
<b>ARTG entry for</b>	Medicine Listed	
<b>Sponsor</b>	FIT-BioCeuticals Limited	
<b>Postal Address</b>	Care of Blackmores Ltd PO Box 1725, Warriewood, NSW, 2102 Australia	
<b>ARTG Start Date</b>	4/12/2012	
<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 . CardioNutrients Forte

<b>Product Type</b>	Single Medicine Product	<b>Effective Date</b>	30/09/2019
---------------------	-------------------------	-----------------------	------------

#### Permitted Indications

Antioxidant/Reduce free radicals formed in the body  
Maintain/support energy levels  
Maintain/support energy production  
Maintain/support physical endurance/capacity/stamina  
Maintain/support general health and wellbeing  
Helps maintain/support healthy cholesterol  
Maintain/support cardiovascular system health  
Maintain/support muscle function

#### Indication Requirements

Product presentation must not imply or refer to serious cardiovascular conditions.  
If product is indicated for weight loss, label statement: When used in conjunction with a program of reduced intake of dietary calories and increased physical activity.  
Product presentation must not imply or refer to chronic fatigue syndrome.  
Product presentation must not imply or refer to lowering or raising blood cholesterol levels from outside of the normal healthy range

#### Standard Indications

No Standard Indications included on Record

#### Specific Indications

No Specific Indications included on Record

Public Summary



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

**Warnings**

Do not take while on warfarin therapy without medical advice.

This medicine contains selenium which is toxic in high doses. A daily dose of 150 micrograms for adults of selenium from dietary supplements should not be exceeded.

**Additional Product information**

**Pack Size/Poison information**

**Pack Size** **Poison Schedule**

**Components**

**1 . Formulation 1**

**Dosage Form** Capsule, soft

**Route of Administration** Oral

**Visual Identification**

**Active Ingredients**

<b>alpha lipoic acid</b>	<b>100 mg</b>
<b>magnesium orotate</b>	<b>400 mg</b>
Equivalent: magnesium	25.6 mg
<b>selenomethionine</b>	<b>166 microgram</b>
Equivalent: selenium	67 microgram
<b>ubidecarenone</b>	<b>100 mg</b>

**Other Ingredients (Excipients)**

Gelatin

glycerol

iron oxide red

lecithin

purified water

Rice bran oil

yellow beeswax

© Commonwealth of Australia. This work is copyright. You are not permitted to re-transmit, distribute or commercialise the material without obtaining prior written approval from the Commonwealth. Further details can be found at <http://www.tga.gov.au/about/website-copyright.htm>.

Public Summary