

**Australian Government** 

## **Department of Health**

# Therapeutic Goods Administration

### **Public Summary**

Summary for ARTG Entry:	159395	D3 Capsules
ARTG entry for	Medicine Listed	
Sponsor	FIT-BioCeuticals	s Limited
Postal Address	Care of Blackmo Australia	pres Ltd PO Box 1725, Warriewood, NSW, 2102
ARTG Start Date	17/02/2009	
Product Category	Medicine	
Status	Active	
Approval Area	Listed Medicines	S

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.

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. D3 Capsule	25		
Product Type	Single Medicine Product	Effective Date	12/06/2020
Permitted Indicati	ons		
Maintain/support be	one health		
Aids/assists health	y bone development/growth/building		
Maintain/support be	one mass/density/integrity		
Maintain/support be	one strength		
Help maintain/supp	port bone mineralisation		
Maintain/support he	ealthy cardiovascular system function		
Maintain/support he	ealthy immune system function		
Maintain/support al	bsorption of dietary (state vitamin/mineral/nutrie	nt)	
Maintain/support (s	state vitamin/mineral/nutrient) levels in the body		
Maintain/support ne	ervous system function		
Indication Require	ements		
	Vitamins/minerals/nutrients/dietary supplements /nutrients/dietary supplements] should not repla		

Product presentation must not imply or refer to bone disease or disorders e.g. rheumatoid arthritis, juvenile arthritis, debilitating osteoarthritis, osteoporosis. Note: this requirement is not intended to apply where the indications referring to osteoporosis specified in column 2 of Table 2 of this instrument are also used.

If product is indicated for supplementation, Label statement: [Vitamins/minerals/nutrients/dietary supplements] can only be of assistance if dietary intake is inadequate OR [Vitamins/minerals/nutrients/dietary supplements] should not replace a balanced diet (or words to that effect).

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

Product presentation must not imply or refer to serious immunological diseases.

Product presentation must not imply or refer to serious cardiovascular conditions.

#### **Standard Indications**

No Standard Indications included on Record

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### This is not an ARTG Certificate document.

The onus is on the reader to verify the current accuracy of the information on the document subsequent to the date shown. Visit www.tga.gov.au for contact information



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Therapeutic Goods Administration

Specific Indications

No Specific Indications included on Record Warnings

No Warnings included on Record

Additional Product information

Container information								
Туре	Material		Life Time	Temperature	Closure	Conditions		
Multiple containers	Not recorded		Not recorded	Not recorded	Not recorded	Not recorded		
Pack Size/Poison information								
Pack Size Poison Schedule								
Components								
1. Formulation 1								
Dosage Form	C	Capsule, soft						
Route of Administr	ation (	Oral						
Visual Identification	n							
Active Ingredients								
colecalciferol	ciferol				25 microgram			
Other Ingredients (Excipients)								
Gelatin								
glycerol								
Soya Oil								
titanium dioxide								

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