

Australian Government

Department of Health

Therapeutic Goods Administration

Public Summary

Products

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Summary for ARTG Entry:	221301	ViroGuard	
ARTG entry for	Medicine Listed		
Sponsor	FIT-BioCeuticals Limited		
Postal Address	Care of Blackmo Australia	ores Ltd PO Box 1725, Warriewood, NSW, 2102	
ARTG Start Date	14/03/2014		
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.

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.

1 . ViroGuard			
Product Type	Single Medicine Product	Effective Date	23/10/2020
Permitted Indication	ons		
	mune system health		
Vaintain/support he	althy immune system function		
•	rence of facial cold sores		
Maintain/support wo	ound healing		
ndication Require	ments		
Product presentation	on must not imply or refer to serious immu	unological diseases.	
Label statement: If	symptoms persist, talk to your health prof	fessional.	
Standard Indicatio	ns		
No Standard Indicat	tions included on Record		
Specific Indication	IS		
No Specific Indication	ons included on Record		
Warnings			
No Warnings includ	ed on Record		
Additional Product	tinformation		
Pack Size/Poison i	information		
Pack Size		Poison Schedule	
Components			
1. Formulation 1	1		
Dosage Form	Capsule, soft		
Route of Admini	stration Oral		
Visual Identificat	tion		
Active Ingredien	its		
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Echinacea purpurea herb Juice dry	9.44 mg
Equivalent: Echinacea purpurea (Fresh)	425 mg
lysine hydrochloride	530.98 mg
Equivalent: lysine	425 mg
Melissa officinalis leaf Extract dry concentrate	107.14 mg
Equivalent: Melissa officinalis (Dry)	750 mg
zinc amino acid chelate	12.5 mg
Equivalent: zinc	2.5 mg

Other Ingredients (Excipients)

Gelatin
glycerol
hydrogenated vegetable oil
iron oxide black
iron oxide red
iron oxide yellow
lecithin
maltodextrin
purified water
silicon dioxide
Soya Oil
titanium dioxide
yellow beeswax

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