

Australian Government

Department of Health and Aged Care

Therapeutic Goods Administration

Public Summary

Summary for ARTG Entry:	220761	CurcuForte
ARTG entry for	Medicine Listed	
Sponsor	Biomedica Nutra	aceuticals Pty Ltd
Postal Address	PO Box 7052, A Australia	LEXANDRIA, NSW, 2015
ARTG Start Date	6/03/2014	
Product Category	Medicine	
Status	Active	
Approval Area	Listed Medicines	3
Conditions		

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 . CurcuForte							
Product Type	Single Medicine Product	Effective Date	4/09/2024				
Permitted Indication	ons						
Antioxidant/Reduce free radicals formed in the body							
Helps reduce/decrease free radical damage to body cells							
Maintain/support general health and wellbeing							
Anti-inflammatory/relieve inflammation							
Decrease/reduce/relieve mild joint inflammation/swelling							
Maintain/support healthy digestive system function							
Maintain/support healthy digestion							
Maintain/support digestive system health							
Indication Requirements							
Label statement: If symptoms persist, talk to your health professional.							
Product presentation must only refer to mild joint symptoms.							
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.							
Standard Indicatio	ons						
No Standard Indications included on Record							
Specific Indications							
No Specific Indications included on Record							
Warnings							
In very rare cases, Curcuma species may harm the liver. Stop use and see a doctor if you have yellowing skin/eyes or unusual: fatigue, nausea, appetite loss, abdominal pain, dark urine, or itching.							
Additional Produc	t information						
Pack Size/Poison	information						
Pack Size		Poison Schedule					

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Components				
1 . Formulation 1				
Dosage Form	Capsule, hard			
Route of Administration	Oral			
Visual Identification				
Active Ingredients				
curcumin			550 mg	
Other Ingredients (Excipie	ents)			
calcium hydrogen phosph	ate dihydrate			
colloidal anhydrous silica				
hypromellose				
purified water				
stearic acid				
tapioca starch				

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