GOUT RELIEVER SYNERGY- gout reliever synergy oil PHYTOPIA CO., LTD.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Drug facts Methyl salicylate 50mg Menthol 20mg Analgesic in the treatment of gout pain. Helps relieve the condition of pain and stiffness. Drug facts Methyl salicylate 50mg Menthol 20mg Analgesic in the treatment of gout pain. Helps relieve the condition of pain and stiffness. Dosage and Usage Adults 5ml maximum per day Apply on skin and gentle massage around painful area. Dosage and Usage Adults 5ml maximum per day

Other ingredients

Arnica montana, peppermint, olive oil, grapefruit oil, eucalyptus oil, matricaria recutita, juniper berry oil, borage oil

Warnings

For external use only. Stop use and ask doctor if rash occurs. Keep out reach of children. Protect this product from excessive heat and direct sun.

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Gout reliever synergy

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Analgesic in the treatment of gout pain

Use

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Product Information				
Product T ype	HUMAN OTC DRUG	Item Code (So	Item Code (Source)	
Route of Administration	TOPICAL			
Active Ingredient/Active M	Aoiety			
Ingredient Name		Ba	sis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (M	MENT	HOL	20 mg in 500 mg	

	Strength				
UNIPER BERRY O					
MATRICARIA RECU	J TITA (UNII: G0R4UBI2ZZ)				
ARNICA MONTANA (UNII: O80TY208ZW)					
BORAGE OIL (UNI	: F8XAG1755S)				
GAULTHERIA PRO	CUMBENS LEAF (UNII: 2125M16OWN)				
PEPPERMINT OIL (UNII: AV092KU4JH)					
GRAPEFRUIT OIL (UNII: YR377U58W9)					
EUCALYPTUS OIL	(UNII: 2R04ONI662)				
OLIVE OIL (UNII: 6	UYK2W1W1E)				
Product Chara	teristics				
Color	brown (Bottle Color)	Score			
Shape		Size			
Flavor		Imprint Code			
Contains					
Contains					
Contains					
Packaging	Package Description	Marketing Start Date	Marketing End Date		
Packaging # Item Code	Package Description 500 mg in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	Date	U		
Packaging # Item Code NDC:70470-1113-	500 mg in 1 BOTTLE, GLASS; Type 0: Not a Combination	Date	U		
Packaging # Item Code NDC:70470-1113-	500 mg in 1 BOTTLE, GLASS; Type 0: Not a Combination	Date	U		
<pre>Packaging # Item Code NDC:70470-1113- 5</pre>	500 mg in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	Date	U		
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Labeler - PHYTOPIA CO., LTD. (656940504)

Registrant - PHYTOPIA CO., LTD. (656940504)

Establishment							
Name	Address	ID/FEI	Business Operations				
PHYTOPIA CO., LTD.		656940504	manufacture(70470-1113)				

Revised: 10/2016

PHYTOPIA CO., LTD.