DOCTORRELIEF- menthol, methyl salicylate and camphor (synthetic) gel Biotanico Inc.

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.

DoctorRelief

Actives

Topical analgesic

Indication

Uses

Temporary relief minor aches and pains of muscles and joints associated with:

■ Arthritis ■ Bruises ■ Simple backache ■ Strains ■ Sprains

Warnings

For external use only

When using this product:

- Use only as directed
- Avoid contact with eyes ,mucous membranes or rashes
- Do not apply to broken, wounded or damaged skin
- Do not bandage or cover with any type of wrap except clothing
- Do not use with a heating pad or apply external heat
- Discontinue use at least 1 hour before a bath or shower and do not use immediately after a bath or shower

Questons

Stop use and ask a doctor if

- Rash, itching or excessive skin irritation develops
- Condition worsens or does not improve after regular use
- Symptoms persist for more than 7 days or clear up and occur again within a few days

OTC-Keep out of reach of childrens

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center immediately

Dosages

Directions

Adult and children 12 years of age and over

■ Clean and dry affected area

- Open pouch and remove patch. If desired, cut patch to size
- Peel off protective backing and apply sticky side to affected area
- Apply to affected area not more than 3 times daily
- Wash hands with soap after applying patch

Children under 12 years of age

■ Consult a doctor

Warning and Precautions

■ This product may cause allergic reaction in some individuals.

Ask Doctor

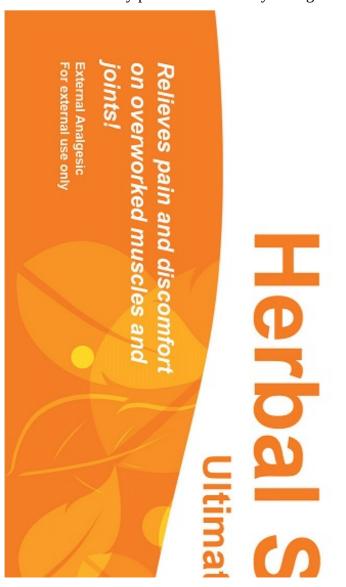
Ask a doctor if symptoms persist for more than 7 days or clear up and occur again within a few days

Inactives

Trichosanthes Cucumenina seed, etc.

Storage and handling section

■ Store in cool dry place from directly sunlight











Herbal Soothe

Ultimate Relief Patch

EXTERNAL PAIN RELIEVING BANDAGE

Drug Facts	
Active ingredients	Purpose
Menthol 60mg/patch	Topical analgesic
Wintergreen Oil 20mg/patch	Topical analgesic
Camphor 10mg/patch	Topical analgesic
Uses temporary relief for minor aches and pains of	muscles and joints associated with:

■Arthritis ■Bruises ■Simple backache ■Strains ■Sprains

Warnings

For external use only

When using this product

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- Apply to affected area no more than 3 times daily
- Wash hands with soap after applying patch

Children under 12 years of age

Consult a doctor before use

Cautions

- Do not apply on ulcerative, or suppurative skin area
- This product may cause allergic reaction in some individuals

Other information

- Store in cool dry place away from direct sunlight
- Re-seal package after opening

Other ingredients Trichosanthes kirilowi radix, Rheum officinale rhizoma, Angelica

BACK

dahurica radix, Phellodendron chinense cortex, Curcuma longa rhizoma, Arisaema erubescens rhizoma, Glycyrrhiza uralensis radix, Magnolia officinalis cortex, Citrus reticulata pericarpium, Atractylodes lancea rhizoma, Natural rubber, Silicon dioxide, Lanolin, Zinc oxide, Indigo Carmine (FD&C Blue #2), FD&C Yellow #4, FD&C Yellow #5, FD&C Red #40, Q.S. Sesame Oil to 1g

Imported by: KPC Products, Inc. 16 Goddard CA 92618, USA

Manufactured by: Biotanico Inc. No. 1 Nanke 1st Rd., Xinshi Dist. Tainan City, 74147 Taiwan















Product of Taiwan

For external use only External Analgesic

Relieves pain and discomfort

FRONT

Herbal Soothe

Ultimate Relief Patch

EXTERNAL PAIN RELIEVING BANDAGE

Drug Facts Purpose Active ingredients Purpose Menthol 60mg/patch... Topical analgesic Wintergreen Oil 20mg/patch... Topical analgesic Camphor 10mg/patch... Topical analgesic

Uses temporary relief for minor aches and pains of muscles and joints associated with:

■Arthritis ■Bruises ■Simple backache ■Strains ■Sprains

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Other ingredients Trichosanthes kirilowi radix, Rheum officinale rhizoma, Angelica dahurica radix, Phellodendron chinense cortex, Curcuma longa rhizoma, Arisaema erubescens rhizoma, Glycyrrhiza uralensis radix, Magnolia officinalis cortex, Citrus reticulata pericarpium, Atractylodes lancea rhizoma, Natural rubber, Silicon dioxide, Lanolin, Zinc oxide, Indigo Carmine (FD&C Blue #2), FD&C Yellow #4, FD&C Yellow #5, FD&C Red #40, Q.S. Sesame Oil to 1g

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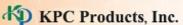














BACK



DOCTORRELIEF

menthol, methyl salicylate and camphor (synthetic) gel

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:57842-7733
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
Menthol (UNII: L7T10 EIP3A) (Menthol - UNII:L7T10 EIP3A)	Menthol	60 mg		
METHYL SALICYLATE (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII:O414PZ4LPZ)	METHYL SALICYLATE	20 mg		
Camphor (synthetic) (UNII: 5TJD82A1ET) (Camphor (synthetic) - UNII:5TJD82A1ET)	Camphor (synthetic)	10 mg		

Inactive Ingredients	
Ingredient Name	Strength
TRICHO SANTHES KIRILO WII ROOT (UNII: V409 XGE0 TS)	125 mg
RHEUM OFFICINALE ROOT (UNII: A47JP5ZG9M)	62 mg
ANGELICA DAHURICA ROOT (UNII: 1V63N2S972)	63 mg
PHELLODENDRON CHINENSIS BARK (UNII: 2866QMZ434)	63 mg
TURMERIC (UNII: 856 YO 1Z64F)	63 mg
ARISAEMA ERUBESCENS ROOT (UNII: 3E8P1PU8SY)	25 mg
GLYCYRRHIZA URALENSIS (UNII: 42B5YD8F0K)	25 mg
MAGNOLIA OFFICINALIS BARK (UNII: 5M609NV974)	25 mg
TANGERINE (UNII: KH3E3096OO)	25 mg
ATRACTYLODES LANCEA ROOT (UNII: CAZ6282J2O)	25 mg
ZINC OXIDE (UNII: SOI2LOH54Z)	20 mg
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	0.05 mg
SESAME OIL (UNII: QX10HYY4QV)	
SILICON DIO XIDE (UNII: ETJ7Z6 XBU4)	
LANOLIN (UNII: 7EV65EAW6H)	
FD&C BLUE NO. 2 (UNII: L06K8R7DQK)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
YELLOW OB (UNII: QE535CBH7S)	

Color yellow (FD&C YELLOW NO. 5) , blue (FD&C BLUE NO. 2) , red (FD&C RED NO. 40) Score Shape Size Flavor Imprint Code	Product Characteristics		
	Color	yellow (FD&C YELLOW NO. 5), blue (FD&C BLUE NO. 2), red (FD&C RED NO. 40)	Score
Flavor Imprint Code	Shape		Size
	Flavor		Imprint Code
Contains	Contains		

Packaging				
# Item Code	Package Description	Marketing Start Date	Marketing End Date	
1 NDC:57842-7733-1	10 in 1 BOX; Type 0: Not a Combination Product	0 1/0 3/20 18		
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph final	part346	0 1/0 3/20 18		

Labeler - Biotanico Inc. (658835665)

Registrant - Biotanico Inc. (658835665)

Establishment			
Name	Address	ID/FEI	Business Operations
Biotanico Inc.		658835665	manufacture(57842-7733)

Revised: 1/2018 Biotanico Inc.