TINNITUS RELINF PATCHES- camphor, borneol patch
Shenzhen Furuizhilian keji Co., Ltd.

TINNITUS RELINF PATCHES

Camphor 3%

Borneol 4%

Topical analgesic

For temporary relief of minor aches & pains of muscles & joints associated with: arthritis, simple backache, strains, bruises, sprains

For external use only

on damaged skin
with a heating pad
if you are allergic to any ingredients of this product

use only as directed
avoid contact with the eyes, mucous membranes or rashes.
do not bandage tightly

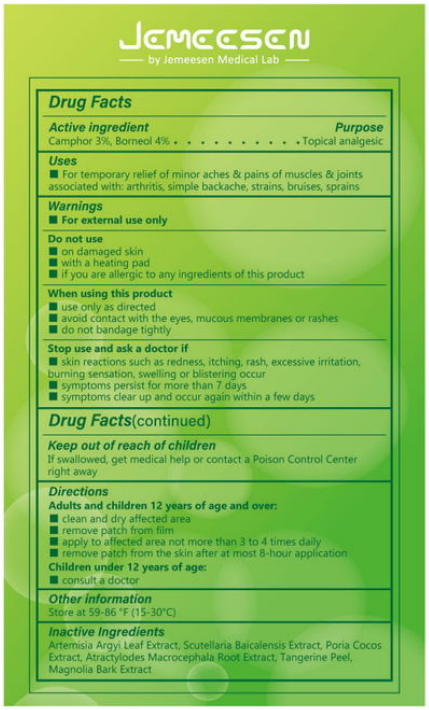
■ skin reactions such as redness, itching, rash, excessive irritation,burning sensation, swelling or blistering occur
■ symptoms persist for more than 7 days
■ symptoms clear up and occur again within a few days

If swallowed, get medical help or contact a Poison Control Center right away.

Adults and children 12 years of age and over:
■ clean and dry affected area
■ remove patch from film
■ apply to affected area not more than 3 to 4 times daily
■ remove patch from the skin after at most8-hour application
Children under 12 years of age:
consult a doctor

Artemisia Argyi Leaf Extract, Scutellaria Baicalensis Extract, Poria Cocos Extract, Atractylodes Macrocephala Root Extract, Tangerine Peel, Magnolia Bark Extract

彩盒尺寸:65×15×120mm
材质:300G铜版纸+覆亮膜
工艺:见彩盒标注



TINNITUS RELINF PATCHES

camphor, borneol patch

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:84445-007
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BORNEOL (UNII: M89NIB437X) (BORNEOL - UNII:M89NIB437X)	BORNEOL	2.4 g in 60 g
CAMPHOR, (-)- (UNII: 213N3S8275) (CAMPHOR, (-)- - UNII:213N3S8275)	CAMPHOR, (-)-	1.8 g in 60 g

Inactive Ingredients	
Ingredient Name	**Strength**
FU LING (UNII: XH37TWY5O4)	
TANGERINE PEEL (UNII: JU3D414057)	
MAGNOLIA OBOVATA BARK (UNII: SM9Z2LD5TK)	
SCUTELLARIA BAICALENSIS ROOT (UNII: 7J95K7ID2S)	
ATRACTYLODES MACROCEPHALA ROOT (UNII: 08T3N29QJB)	

ARTEMISIA ARGYI LEAF (UNII: 2JYC99Q0WZ)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84445-007-01	60 g in 1 BOX; Type 0: Not a Combination Product	07/17/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	07/17/2024	

Labeler - Shenzhen Furuizhilian keji Co., Ltd. (418598613)

Establishment

Name	Address	ID/FEI	Business Operations
Shenzhen Furuizhilian keji Co., Ltd.		418598613	manufacture(84445-007)

Revised: 7/2024 Shenzhen Furuizhilian keji Co., Ltd.