

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

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

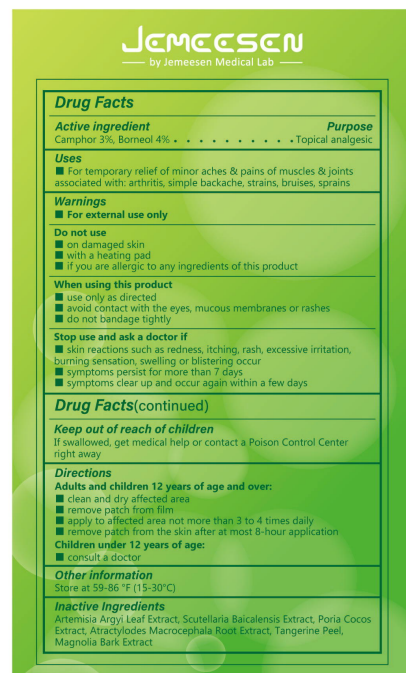
■ remove patch from the skin after at most 8-hour application

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:84445-010
Route of Administration	TRANSDERMAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CAMPBOR, (-)- (UNII: 213N3S8275) (CAMPBOR, (-)- - UNII:213N3S8275)	CAMPBOR, (-)-	0.084 g in 28
BORNEOL (UNII: M89NIB437X) (BORNEOL - UNII:M89NIB437X)	BORNEOL	1.12 g in 28

Inactive Ingredients

Ingredient Name	Strength
ATRACYLDES MACROCEPHALA ROOT (UNII: 08T3N29QJB)	
TANGERINE PEEL (UNII: JU3D414057)	
SCUTELLARIA BAICALENSIS ROOT (UNII: 7J95K7ID2S)	
ARTEMISIA ARGYI LEAF (UNII: 2JYC99Q0WZ)	

FU LING (UNII: XH37TWY5O4)				
MAGNOLIA OBOVATA BARK (UNII: SM9Z2LD5TK)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84445-010-01	28 in 1 BOX; Type 0: Not a Combination Product	08/08/2024	
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug		505G(a)(3)	08/08/2024	

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

Establishment			
Name	Address	ID/FEI	Business Operations
Shenzhen Furuizhilian keji Co., Ltd.		418598613	manufacture(84445-010)