JIKIMI PATCH- anti-bacterial patch patch
BM Pharmaceutical Co., Ltd.

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Active Ingredient(s)

Ethyl alcohol 7% w/w……...……. Antimicrobial

Levomenthol 3% w/w……...……. Antifungal

Eucalyptus oil 20% w/w……...…. Antimicrobial

Purpose

To help reduce bacteria.

Use

Apply to affected area as topical aromatic diffusion.

Warnings

For external use only. Flammable. Keep away from heat or flame

Do not use

- Do not use if you are allergic to any ingredients of this product.
- Do not use in children under 6 and pregnant or breast-feeding.

When using this product use only as directed. Use within 2 days after removing the film. Avoid contact with the eyes. In case of contact, rinse eyes thoroughly with water. Avoid contact with broken skin. Do not inhale or ingest.

Stop use and ask a doctor if irritation or rash appears and lasts. It happen headache. Conditions worsen.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

Remove patch from film.
Apply to affected area as topical aromatic diffusion not more than 3 to 4 times daily.

Remove patch at most 24-36 hours application.

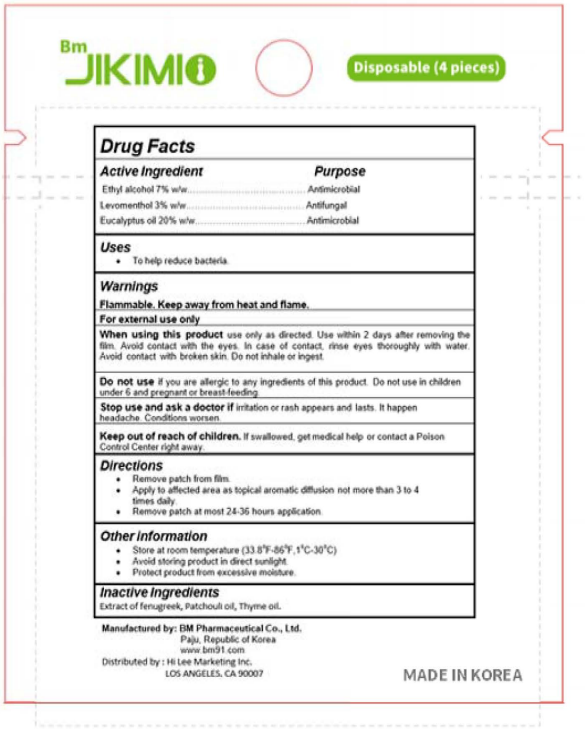
Other information

- Store at room temperature between 1-30oC (33.8-86oF).
- Avoid storing product in direct sunlight.
- Protect product from excessive moisture.

Inactive ingredients

Extract of fenugreek, Patchouli oil, Thyme oil.

Package Label - Principal Display Panel



JIKIMI PATCH

anti-bacterial patch patch

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:76614-100
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
LEVOMENTHOL (UNII: BZ1R15MTK7) (LEVOMENTHOL - UNII:BZ1R15MTK7)	LEVOMENTHOL	0.03
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.07
EUCALYPTUS OIL (UNII: 2R04ONI662) (EUCALYPTUS OIL - UNII:2R04ONI662)	EUCALYPTUS OIL	0.2

Inactive Ingredients	
Ingredient Name	**Strength**
FENUGREEK SEED OIL (UNII: 50LM4E7MG3)	
POGOSTEMON CABLIN LEAF OIL (UNII: F3IN55X5PO)	
THYME OIL (UNII: 2UK410MY6B)	

Product Characteristics			
Color		Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76614-100-10	4 in 1 PACKAGE; Type 0: Not a Combination Product	05/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/01/2020	

Labeler - BM Pharmaceutical Co., Ltd. (694470190)

Registrant - BM Pharmaceutical Co., Ltd. (694470190)

Establishment

Name	Address	ID/FEI	Business Operations
BM Pharmaceutical Co., Ltd.		694470190	manufacture(76614-100)

Revised: 5/2020 BM Pharmaceutical Co., Ltd.