

ANTIBACTERIAL PE FILM- titanium dioxide film
DH Investment 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](#).

titanium dioxide

polyethylene, copper

Antibacterial

Keep out of reach of children

apply proper amount where needed

For external use only

When using this product

■ if following abnormal symptoms persist, discontinue use

Irritation around the eyes, ears, mucous membranes, including the mouth, under the skin irritation and rashes

■ Stop immediately and consult a doctor if you experience

1) Hypersensitivity symptoms such as erythema, itching and dermatitis.

2) Skin Irritation

3) Following Instructions when using medication

(1) For external use only (Do not use internally)

(2) Avoid getting into the eyes (if contact occurs, wash well with clean water)

■ Do not use the product for a long time in the same area as swelling, inflammation or sickness may occur due to absorption through the skin.

It is not recommended to use this one areas that have been medically treated with a cast or bandage.

For external use only

Drug Facts

Active Ingredients

Titanium dioxide -----

Purpose

antibacterial

Uses

■ antibacterial

Warning

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Directions

■ apply proper amount where needed

Other Information

- read the directions and warnings before use

- avoid freezing and excessive heat above 40 degree C (104 degree F)

Inactive Ingredients

polyethylene, copper

Questions or Comments? Call XXXXX or email: XXXXXX

ANTIBACTERIAL PE FILM

titanium dioxide film

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79234-0001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name		Basis of Strength	Strength	
TITANIUM DIOXIDE (UNII: 15FIX9 V2JP) (TITANIUM DIOXIDE - UNII:15FIX9 V2JP)		TITANIUM DIOXIDE	0.3 g in 100 g	
Inactive Ingredients				
Ingredient Name			Strength	
COPPER (UNII: 789U1901C5)				
HIGH DENSITY POLYETHYLENE (UNII: UG00KM4WR7)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79234-0001-1	1 g in 1 PACKAGE; Type 0: Not a Combination Product	06/27/2020	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
unapproved drug other		06/27/2020		

Labeler - DH Investment Co Ltd (694798202)

Registrant - DH Investment Co Ltd (694798202)

Establishment

Name	Address	ID/FEI	Business Operations
DH Investment Co Ltd		694798202	manufacture(79234-0001)

Revised: 6/2020

DH Investment Co Ltd