BANDHA FOR AND MASK- niacinamide, adenosine patch
DKCOSTECH

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 INGREDIENTS

NIACINAMIDE 2.0%
ADENOSINE 0.04%

INACTIVE INGREDIENTS

Water, Dendropanax Morbiferus Leaf Extract, Propanediol, Glycerin, 1,2-Hexanediol, Butylene Glycol, betaine, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Tromethamine, Xanthan Gum, Aloe Barbadensis Leaf Extract, Astragalus Membranaceus Root Extract, Boswellia Serrata Extract, Melia Azadirachta Leaf Extract, Phyllanthus Emblica Extract, Trifolium Pratense (Clover) Flower Extract, Garcinia Cambogia Fruit Extract, Allantoin, Disodium EDTA, Fragrance, Sodium Hyaluronate

PURPOSE

Brightening
Anti-wrinkle

WARNINGS

1. For external use only
2. Avoid contact with eyes.
3. Discontinue use if signs of irritation or rashes appear.
4. Keep out of reach of children.

KEEP OUT OF REACH OF CHILDREN

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Uses

■ Helps brighten skin tone.
■ Helps prevent wrinkle problem.

Directions

1. After cleansing your face,
2. Leave it on for 20 minutes.
3. Gently remove the mask and pat the remaining essence into your face.

QUESTIONS

+8227978088
www.bandha.co.kr

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

BANDHA FOR AND MASK

niacinamide, adenosine patch

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:80623-010
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
NIACINAMIDE (UNII: 25X51I8RD4) (NIACINAMIDE - UNII:25X51I8RD4)	NIACINAMIDE	0.56 g in 28 g
ADENOSINE (UNII: K72T3FS567) (ADENOSINE - UNII:K72T3FS567)	ADENOSINE	0.01 g in 28 g

Inactive Ingredients	
Ingredient Name	**Strength**
Water (UNII: 059QF0KO0R)	
Propanediol (UNII: 5965N8W85T)	
Glycerin (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80623-010-02	10 in 1 CARTON	09/01/2020	
1	NDC:80623-010-01	28 g in 1 POUCH; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		09/01/2020	

Labeler - DKCOSTECH (695504288)

Registrant - DKCOSTECH (695504288)

Establishment

Name	Address	ID/FEI	Business Operations
HANSCOS Co.,Ltd.		688494423	manufacture(80623-010)

Revised: 9/2020 DKCOSTECH