DPC AURA BOOSTER MASK- niacinamide liquid
MSCO

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.

Drug Facts

NIACINAMIDE

Water
Glycerin
Butylene Glycol
Sodium Hyaluronate
1,2-Hexanediol
Allantoin
Dipotassium Glycyrrhizate
Trehalose
Hydroxyethylcellulose
Carbomer
PEG-60 Hydrogenated Castor Oil
Glyceryl Acrylate/Acrylic Acid Copolymer
Propylene Glycol
PVM/MA Copolymer
Panthenol
Diamond powder
Simmondsia Chinensis (Jojoba) Seed Oil
Anthemis Nobilis Flower Extract
Jasminum Officinale (Jasmine) Flower Water
Lilium Tigrinum Extract
Arnica Montana Flower Extract
Arginine
Chlorphenesin
Disodium EDTA
Glycyrrhiza Glabra (Licorice) Root Extract
Zingiber Officinale (Ginger) Root Extract
Schizandra Chinensis Fruit Extract
Coptis Japonica Root Extract
Camellia Sinensis Leaf Extract
Caprylyl Glycol
Citrus Grandis (Grapefruit) Seed Extract
Acorus Calamus Root Extract
Perilla Ocymoides Leaf Extract
Fragrance

Skin Protectant - Intensive Whitening

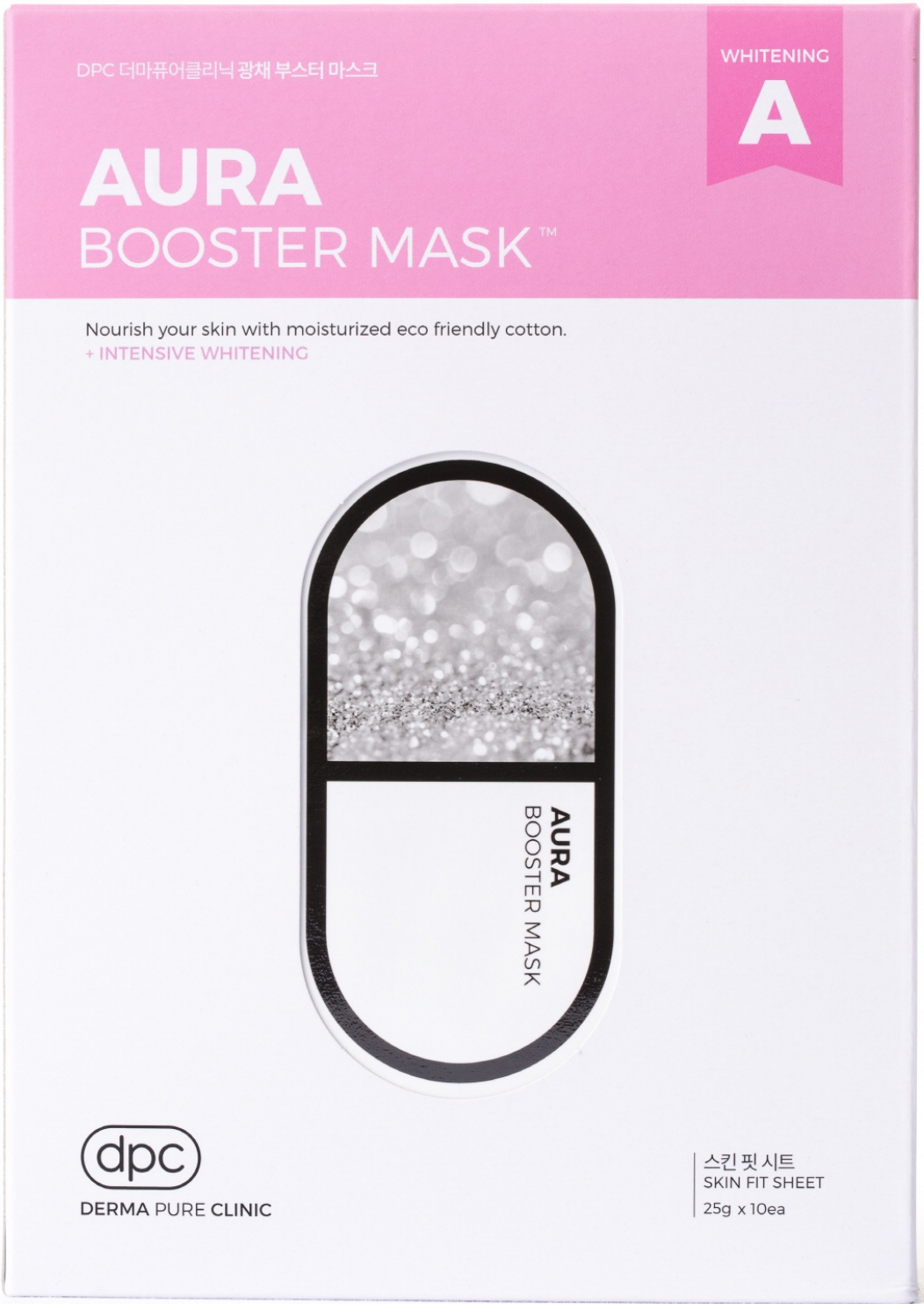
keep out of reach of the children

1. After cleansing wipe water and apply skin lotion on face.
2. Apply the mask evenly over the face for 10-20 minutes and remove.
3. Pad the left over residue lightly to absorb.

For external use only.
Avoid contact with eyes. Not for human consumption.

Discontinue use if irritation occurs.
If irritation persists, consult a physician.

for external use only



DPC AURA BOOSTER MASK

niacinamide liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71673-0001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
NIACINAMIDE (UNII: 25X51I8RD4) (NIACINAMIDE - UNII:25X51I8RD4)	NIACINAMIDE	2 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
ALLANTOIN (UNII: 344S277G0Z)	
WATER (UNII: 059QF0KO0R)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:71673-0001-1	5 in 1 PACKAGE	07/01/2017	
1		25 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		07/01/2017	

Labeler - MSCO (689039838)

Registrant - MSCO (689039838)

Establishment

Name	Address	ID/FEI	Business Operations
MSCO		689039838	label(71673-0001)

Establishment

Name	Address	ID/FEI	Business Operations
CNF CO.,LTD.		689852175	manufacture(71673-0001)

Revised: 8/2017 MSCO