DPC DUAL MASTER VITAMIN C- niacinamide, adenosine 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, ADENOSINE

Prunus Mume Flower Water
Hyaluronic Acid
Butylene Glycol
Water
Theanine
Milk Lipids
Glutathione
Hydrogenated Lecithin
Ceramide NP
1,2-Hexanediol
Glycerin
Panthenol
Propanediol
Benzyl Glycol
Ethylhexylglycerin
Raspberry Ketone
Hydrolyzed Hyaluronic Acid
Sodium Hyaluronate
Sodium Hyaluronate Crosspolymer
Lactococcus Ferment Lysate
Sodium Benzoate
Hydrogenated Lecithin
Polyglycerin-3
PCA
PEG-240/HDI Copolymer Bis-Decyltetradeceth-20 Ether
Malus Domestica Fruit Cell Culture Extract
Caprylyl Glycol
Tropolone
Octyldodeceth-16
Arginine
Sodium Acetylated Hyaluronate
Hydrolyzed Glycosaminoglycans
Citric Acid
Tremella Fuciformis Polysaccharide
Tuber Melanosporum Extract
Astragalus Membranaceus Root Extract
Rhizobian Gum
Copper PCA
Dipotassium Glycyrrhizate
Betaine
Acetyl Hexapeptide-8
PVP
Fullerenes

Fragrance

Anti-Wrinkle

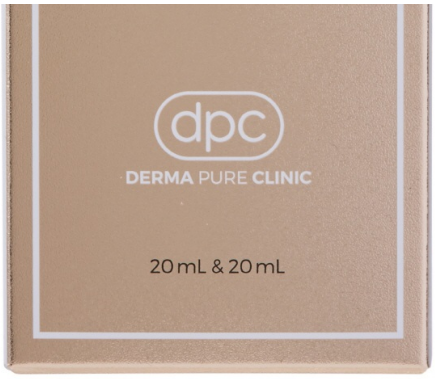
Whitening

keep out of reach of the children

Apply a proper amount on the skin and spread evenly.

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 DUAL MASTER VITAMIN C

niacinamide, adenosine liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71673-0005
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ADENOSINE (UNII: K72T3FS567) (ADENOSINE - UNII:K72T3FS567)	ADENOSINE	0.04 g in 100 mL
NIACINAMIDE (UNII: 25X51I8RD4) (NIACINAMIDE - UNII:25X51I8RD4)	NIACINAMIDE	2 g in 100 mL

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

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:71673-0005-1	40 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	08/23/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		08/23/2017	

Labeler - MSCO (689039838)

Registrant - MSCO (689039838)

Establishment

Name	Address	ID/FEI	Business Operations
MSCO		689039838	label(71673-0005)

Establishment

Name	Address	ID/FEI	Business Operations
ATEC Co., Ltd.		689276681	manufacture(71673-0005)

Revised: 8/2017 MSCO