THE SKIN HOUSE HYALURONIC 6000 AMPOULE- sodium hyaluronate liquid
NOKSIBCHO cosmetic 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.

Drug Facts

Sodium Hyaluronate

Water
Sodium Hyaluronate
Phenoxy ethanol
Glycerin
Butylene glycol
Betaine
Alcohol
1,2-Hexanediol
PPG-26-Buteth-26
PEG-40 Hydrogenated Castor Oil
Arginine
Squalane
Camellia Sinensis Leaf Extract
Hydrolyzed Collagen
Dipotassium Glycyrrhizate
Sodium Citrate
Glyceryl Acrylate/Acrylic Acid Copolymer
Propylene Glycol
PVM/MA Copolymer
Carbomer
Xanthan GUM
Solanum Lycopersicum (Tomato) Fruit Extract
Paeonia Lactiflora Bark/Sap Extract
Lonicera Japonica (Honeysuckle) Flower Extract
Myrciaria Dubia Fruit Extract
Euterpe Oleracea Fruit Extract
Ardisia Crispa Extract
Rubus Coreanus Fruit Extract
Phellinus Linteus Extract
Aloe Barbadensis Leaf Extract
Nelumbo Nucifera Leaf Extract
Citric Acid
Adenosine
Disodium EDTA
Panax ginseng culture extract
Caprylyl Glycol
Phaseolus Radiatus Seed Extract
Ethylhexylglycerin
Gardenia Florida Fruit Extract
Parfum

anti-wrinkle function

keep out of reach of the children

apply proper amount and gently massage

For external use only

When using this product

■ if the following symptoms occurs after use, stop use and consult with a skin specialist

red specks, swelling, itching

■ don’t use on the part where there is injury, eczema, or dermatitis

Keep out of reach of children

■ if swallowed, get medical help or contact a person control center immediately

for external use only

THE SKIN HOUSE HYALURONIC 6000 AMPOULE

sodium hyaluronate liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73590-0020
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
HYALURONATE SODIUM (UNII: YSE9PPT4TH) (HYALURONIC ACID - UNII:S270N0TRQY)	HYALURONATE SODIUM	6 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:73590-0020-1	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/14/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		03/14/2020	

Labeler - NOKSIBCHO cosmetic Co., Ltd. (690182175)

Registrant - NOKSIBCHO cosmetic Co., Ltd. (690182175)

Establishment

Name	Address	ID/FEI	Business Operations
NOKSIBCHO cosmetic Co., Ltd.		690182175	manufacture(73590-0020)

Revised: 3/2020 NOKSIBCHO cosmetic Co., Ltd.