THE SKIN HOUSE VITAL BRIGHT TONER- niacinamide 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

Niacinamide

Water
Butylene glycol
Sodium Hyaluronate
Cetyl Ethylhexanoate
Cyclomethicone
Stearic Acid
Isohexadecane
Butylene Glycol Dicaprylate/Dicaprate
Cetearyl Alcohol
Beeswax
Sorbitan Stearate
Squalane
1,2-Hexanediol
Glycerin
Arachidyl Alcohol
Behenyl Alcohol
Arachidyl Glucoside
Hydrolyzed Collagen
Polysorbate 60
Glyceryl Stearate
Dimethicone
Limnanthes Alba (Meadowfoam) Seed Oil
Tocopheryl Acetate
Solanum Lycopersicum (Tomato) Fruit Extract
Paeonia Lactiflora Extract
Lonicera Japonica (Honeysuckle) Flower Extract
Arginine
Allantoin
Panthenol
Disodium EDTA
Snail Secretion Filtrate
Panax ginseng callus culture extract
Bambusa vulgaris Callus Culture extracts
Aloe vera Callus Culture extracts
Leontopodium alpinum extract
Thymus Vulgaris (Thyme) Extract
Buddleja Davidii Extract
Ascorbic Acid
Phenoxyethanol
Ethylhexylglycerin
Parfum

skin brightening

keep out of reach of the children

apply proper amount to the skin

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

THE SKIN HOUSE VITAL BRIGHT TONER

niacinamide liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73590-0029
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 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:73590-0029-1	130 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/08/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		03/08/2020	

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(73590-0029)

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