BLACK SNAIL COLLAGEN TONER- snail secretion filtrate, collagen 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

Snail Secretion Filtrate

Water

Betula Alba Juice

Butylene glycol

Methyl Gluceth-20

Sodium Hyaluronate

1,2-Hexanediol

Water

Glycoproteins

Sodium Hyaluronate

Betaine

Butylene glycol

Glycerin

1,2-Hexanediol

Viola Mandshurica Flower Extract

Morus Alba Fruit Extract

Sea Water

Galactomyces Ferment Filtrate

Portulaca Oleracea Extract

PPG-26-Buteth-26

PEG-40 Hydrogenated Castor Oil

Allantoin

Black Soybean Extract

Sesamum Indicum (Sesame)

Seed Extract

Oryza Sativa (Rice) Extract

Theobroma Cacao (Cocoa) Extract

Panax Ginseng Berry Extract

Adenosine

Copper Tripeptide-1

Hydrolyzed Collagen

Carbomer

Triethanolamine

Dipotassium Glycyrrhizate

Ethylhexylglycerin

Parfum

anti-wrinkle and nourish

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

lacksquare 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



BLACK SNAIL COLLAGEN TONER

snail secretion filtrate, collagen liquid

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73590-0014	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
SNAIL, UNSPECIFIED (UNII: 2VIO5GL901) (SNAIL, UNSPECIFIED - UNII:2VIO5GL901)	SNAIL, UNSPECIFIED	60 g in 100 mL	

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

ı	Packaging				
# Item Code Package Description		Marketing Start Date	Marketing End Date		
ı	1	NDC:73590-0014-1	150 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 (73590-0014)	

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