

**BEOL CHO LONG- menthol, garlic oil liquid
sunkyung trading**

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.

menthol, garlic oil

digestion

Keep out of reach of children

shake the container for 2 to 3 times before use, apply it evenly

■ if following abnormal 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

purified water

Batylene Glycol

allantoin

SODIUMHYALURONATE

Solubiliser

1.2Hexanediol

Vitamin E

Carbomer

DisodiumEDTA

TEA TREE oil

citrus junos seed oil

Avandula Angustifolia(Lavander) oil

for external use only

How is Aloe vera Amuldy S different?



- 1 Relieve itchiness**
It relieves itchiness by providing moisture and nutrients to the dry skin.
- 2 Contains aloe vera**
Aloe vera helps maintain skin health.
- Aroma herb**
- 3** It relieves skin stimulation, and it does not sting your eyes even if you apply it around your eyes.
- 4 Fast absorption**
Thanks to its fast absorption, it soothes skin quickly. It also purifies skin and reduces stimuli.
- 5 Roll-on applicator**
Roll-on applicator is easy to use and carry.

BEOL CHO LONG

menthol, garlic oil liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:82994-0001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	1.2 g in 100 mL
GARLIC OIL (UNII: 4WG8U28833) (GARLIC OIL - UNII:4WG8U28833)	GARLIC OIL	7.5 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
ALLANTOIN (UNII: 344S277G0Z)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
WATER (UNII: 059QF0K00R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82994-0001-1	20 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/22/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		09/22/2022	

Labeler - sunkyung trading (695235330)

Registrant - sunkyung trading (695235330)

Establishment

Name	Address	ID/FEI	Business Operations
sunkyung trading		695235330	manufacture(82994-0001)

Revised: 9/2022

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