

**DEWYTREE URBAN SHADE HYALURONIC LIGHTWEIGHT SUNSCREEN-  
octocrylene, homosalate, octisalate, avobenzone cream  
DEWYTREE CO. ,LTD**

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**73071-101 DEWYTREE URBAN SHADE HYALURONIC LIGHTWEIGHT SUNSCREEN**

Octocrylene 9.8%

Homosalate 7.2%

Octisalate 4.8%

Avobenzone 3.0%

Sunscreen

■ Helps prevent sunburn. ■ if used as directed with other sun protection measures ( **see *Directions***), decreases the risk of skin cancer and early skin aging caused by the sun.

■ For external use only.

Do not use ■ on damaged or broken skin.

Stop use and ask a doctor if ■ rash occurs.

When using this product ■ keep out of eyes. Rinse with water to remove.

■ Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center.

***Directions***

■ Apply liberally 15 minutes before sun exposure. ■ Reapply at least every 2 hours. ■ Children under 6 months of age: Ask a doctor. ■ **Sun Protection Measures.** Spending time in the sun increases your risk of skin cancer and early skin aging. ■ To decrease this risk, regularly use a sunscreen with a Broad Spectrum SPF value of 15 or higher and other sun protection measures including: ■ limit time in the sun, especially from 10a.m.-2 p.m. wear long-sleeve shirts, pants, hats, and sunglasses.

■ protect this product in this container from excessive heat and direct sun.

Water(Aqua), Isohexadecane, Dipropylene Glycol, Silica, Niacinamide, Cetearyl Glucoside, Sorbitan Olivatate, Benzyl Glycol, Panthenol, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Arginine, Cetearyl Alcohol, Xanthan Gum, Pentylene Glycol, Adenosine, Ethylhexylglycerin, Mentha Piperita (Peppermint) Oil, Squalane, Ganoderma Lucidum (Mushroom) Stem Extract, Butylene Glycol, 1,2-Hexanediol, Hydrolyzed Hyaluronic Acid, Hydrolyzed Sodium Hyaluronate, Dimethylsilanol Hyaluronate, Sodium Hyaluronate, Hyaluronic Acid, Potassium Hyaluronate, Sodium Hyaluronate Crosspolymer, Hydroxypropyltrimonium Hyaluronate, Sodium Hyaluronate Dimethylsilanol, Sodium Acetylated Hyaluronate



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octocrylene, homosalate, octisalate, avobenzone cream

## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:73071-101
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
<b>OCTISALATE</b> (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	2.4 g in 50 mL
<b>HOMOSALATE</b> (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	3.6 g in 50 mL
<b>OCTOCRYLENE</b> (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	4.9 g in 50 mL
<b>AVOBENZONE</b> (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	1.5 g in 50 mL

**Inactive Ingredients**

Ingredient Name	Strength
<b>ARGININE</b> (UNII: 94ZLA3W45F)	
<b>SODIUM ACETYLATED HYALURONATE</b> (UNII: WN66R7GL93)	
<b>ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER (60000 MPA.S)</b> (UNII: 8Z5ZAL5H3V)	
<b>SQUALANE</b> (UNII: GW89575KF9)	
<b>CETEARYL ALCOHOL</b> (UNII: 2DMT128M1S)	
<b>NIACINAMIDE</b> (UNII: 25X51I8RD4)	
<b>ADENOSINE</b> (UNII: K72T3FS567)	
<b>BUTYLENE GLYCOL</b> (UNII: 3XUS85K0RA)	
<b>DIMETHYSILANOL HYALURONATE</b> (UNII: Z853O1D4TE)	
<b>PANTHENOL</b> (UNII: WW9CM0O67Z)	
<b>DIPROPYLENE GLYCOL</b> (UNII: E107L85C40)	
<b>SILICA</b> (UNII: ETJ7Z6XBU4)	
<b>PENTYLENE GLYCOL</b> (UNII: 50C1307PZG)	
<b>ETHYLHEXYLGLYCERIN</b> (UNII: 147D247K3P)	
<b>1,2-HEXANEDIOL</b> (UNII: TR046Y3K1G)	
<b>SORBITAN OLIVATE</b> (UNII: MDL271E3GR)	
<b>BENZYL GLYCOL</b> (UNII: 06S8147L47)	
<b>GANODERMA LUCIDUM STEM</b> (UNII: U8PA41532G)	
<b>XANTHAN GUM</b> (UNII: TTV12P4NEE)	
<b>SODIUM HYALURONATE</b> (UNII: YSE9PPT4TH)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>ISOHEXADECANE</b> (UNII: 918X1OUF1E)	
<b>MENTHA PIPERITA (PEPPERMINT) OIL</b> (UNII: AV092KU4JH)	
<b>HYALURONIC ACID</b> (UNII: S270N0TRQY)	
<b>CETEARYL GLUCOSIDE</b> (UNII: 09FUA47KNA)	

**Product Characteristics**

<b>Color</b>	white	<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>		<b>Imprint Code</b>	
<b>Contains</b>			

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73071-101-01	1 in 1 BOX	03/26/2025	

<b>1</b>	NDC:73071-101-02	50 mL in 1 TUBE; Type 0: Not a Combination Product		
<b>Marketing Information</b>				
<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>		<b>Marketing Start Date</b>	<b>Marketing End Date</b>
OTC Monograph Drug	M020		03/26/2025	

**Labeler -** DEWYTREE CO. ,LTD (688358542)

<b>Establishment</b>			
<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
reBom Co., Ltd.		695951708	manufacture(73071-101)

Revised: 10/2025

DEWYTREE CO. ,LTD