BELIF THE TRUE CREAM AQUA BOMB SUNSCREEN BROAD SPECTRUM SPF 45-
avobenzone, homosalate, octisalate, octocrylene cream
fmg Co.Ltd

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belif The true cream aqua bomb sunscreen Broad Spectrum SPF 45 (10ml)

ACTIVE INGREDIENTS

Avobenzone 2.7%
Homosalate 9%
Octisalate 4.5%
Octocrylene 9%

PURPOSE

Sunscreen

USES

- 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.

WARNINGS

- For external use only.
- **Do not use** on damaged or broken skin.
- **When using this product** keep out of eyes. Rinse with water to remove.
- **Stop use and ask doctor if** rash occurs.
- **Keep out of reach of children.** If product is swallowed, get medical help or contact a Poison Control Center right away.

DIRECTIONS

- Apply generously and evenly 15 minutes before sun exposure.
- Reapply at least every 2 hours.
- Use a water resistant sunscreen if swimming or sweating.
- **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 10 a.m. - 2 p.m.

- Wear long-sleeved shirts, pants, hats, and sunglasses.

- Children under 6 months: Ask a doctor.

INACTIVE INGREDIENTS

Water, Glycerin,

OTHER INFORMATION

Protect the product in this container from excessive heat and direct sun

QUESTIONS OR COMMENTS?

Contact our Sun Specialist at **1-866-993-4709** or **www.belifusa.com**

DISTR. LG H&H Co., Ltd.
58, Saemunan-ro,
Jongno-gu, Seoul,
Korea

www.belifcosmetic.com

Made in Korea

PRINCIPAL DISPLAY PANEL

belif
believe in truth

The true cream -
aqua bomb sunscreen
Broad Spectrum SPF 45

Lightweight moisturizer and
sun protection in one? Yes!

Hydrates skin with
lightweight texture
---------- Yes

No visible white cast
---------- Yes

Dermatologically tested

10 ml / 0.33 fl. oz.

Drug Facts (continued)

Inactive Ingredients

Water, Glycerin, Dimethicone, Cetearyl Alcohol, Potassium Cetyl Phosphate, Panthenol, Benzotriazolyl Dodecyl P-Cresol, 1,2-Hexanediol, Glyceryl Caprylate, Isododecane, Glyceryl Stearate, Niacinamide, Caprylic/Capric Triglyceride, Acrylates/Polytrimethylsiloxymethacrylate Copolymer, Cetearyl Olivate, Inulin Lauryl Carbamate, Sorbitan Olivate, Sodium Polyacryloyldimethyl Taurate, Magnesium Aluminum Silicate, Tromethamine, Xanthan Gum, Butylene Glycol, Trisodium EDTA, Myristica Fragrans (Nutmeg) Extract, Stellaria Media (Chickweed) Extract, Alchemilla Vulgaris Leaf Extract, Equisetum Arvense Leaf Extract, Urtica Dioica (Nettle) Leaf Extract, Rubus Idaeus (Raspberry) Leaf Extract, Nepeta Cataria Extract, Calendula Officinalis Flower Extract, Avena Sativa (Oat) Kernel Extract, Baptisia Tinctoria Root Extract, Lonicera Japonica (Honeysuckle) Flower Extract, Morinda Citrifolia Fruit Extract, Citrus Aurantium Dulcis (Orange) Peel Oil, Citrus Aurantifolia (Lime) Oil, Pelargonium Graveolens Flower Oil, Rosmarinus Officinalis (Rosemary) Leaf Oil, Limonene, Citronellol, Geraniol, Citral, Linalool

MIX
Packaging
Emballage
FSC FSC® C127945

8 03954 41527 8

52102225

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Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72330-722
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	4.5 g in 50 mL
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	9 g in 50 mL
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	2.7 g in 50 mL
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	9 g in 50 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72330-722-10	1 in 1 BOX	02/29/2024	
1		10 mL in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	02/29/2024	

Labeler - fmg Co.Ltd (690188305)

Registrant - LG H&H Co.,Ltd. (688276187)

Establishment

Name	Address	ID/FEI	Business Operations
fmg Co.Ltd		690188305	manufacture(72330-722)

Revised: 2/2024 fmg Co.Ltd