

ANTIBACTERIAL MOISTURIZING HAND LOTION SUN BLOSSOM- benzalkonium chloride lotion

CVS PHARMACY

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

DRUG FACTS

ACTIVE INGREDIENT

BENZALKONIUM CHLORIDE 0.1 PERCENT

PURPOSE

ANTIMICROBIAL

USES

TO HELP REDUCE BACTERIA ON THE SKIN. RECOMMENDED FOR ROUTINE USE.

WARNINGS

FOR EXTERNAL USE ONLY.

WHEN USING THIS PRODUCT

AVOID CONTACT WITH EYES. IF CONTACT OCCURS, RINSE THOROUGHLY WITH WATER.

STOP USING THIS PRODUCT AND ASK DOCTOR IF

IRRITATION OR REDNESS DEVELOPS AND LASTS.

KEEP OUT OF REACH OF CHILDREN

IN CASE OF ACCIDENTAL INGESTION, GET MEDICAL HELP OR CONTACT A POISON CONTROL CENTER IMMEDIATELY.

DIRECTIONS

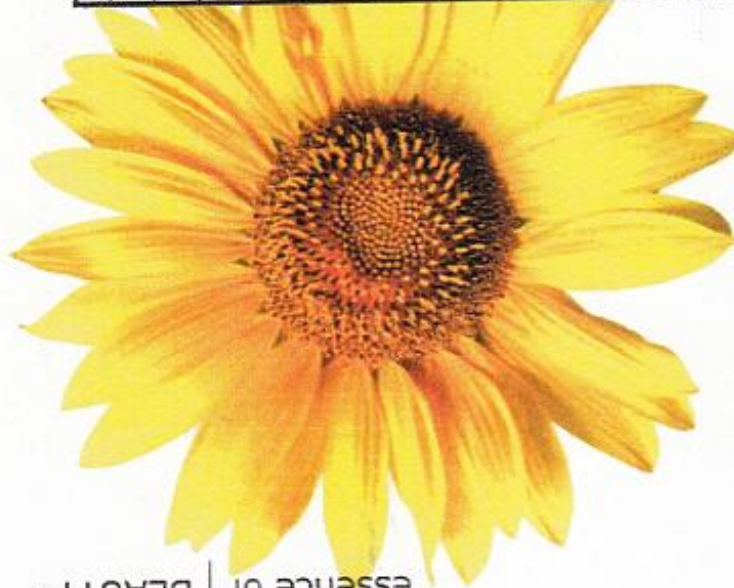
APPLY A SMALL AMOUNT TO PALM AND RUB HANDS TOGETHER THOROUGHLY.

INACTIVE INGREDIENTS

WATER, CYCLOPENTASILOXANE, GLYCINE SOJA (SOYBEAN) OIL, GLYCERIN, CETEARYL ALCOHOL, CETEARETH-20, GLYCERYL STEARATE, DIMETHICONE, CETYL ALCOHOL, STEARYL ALCOHOL, CAPRYLIC/CAPRIC TRIGLYCERIDE, ETHYLHEXYL PALMITATE, DICAPRYLYL CARBONATE, THEOBROMA CACAO (COCOA) SEED BUTTER, FRAGRANCE, HELIANTHUS ANNUUS (SUNFLOWER) SEED EXTRACT, ETHYLHEXYL METHOXYCINNAMATE, BUTYL METHOXYDIBENZOYLMETHANE, ETHYLHEXYL SALICYLATE, PPG-26-BUTETH-26, PEG-40 HYDROGENATED CASTOR OIL, DMDM HYDANTOIN, MANNITOL, CELLULOSE, MICA, TITANIUM OXIDE (CI 77891), IRON OXIDES (CI 77491), IRON OXIDES (CI 77492), RED 40 (CI 16035), YELLOW 5 (CI 19140).

Antibacterial

Antibacterial
moisturizing hand lotion
Softens, kills germs &
leaves hands lightly scented.



essence of | BEAUTY®

SUNBLOSSOM

moisturizing hand lotion

kills 99.99% of germs

2.5 FL OZ (72 mL)

Drug Facts

Active Ingredient
Benzalkonium chloride 0.1% Antimicrobial

Uses ■ To help reduce the bacteria on the skin
■ recommended for routine use

Warnings
■ For external use only.
■ When using this product ■ avoid contact with eyes.
if contact occurs, rinse thoroughly with water.

Stop using this product and ask doctor if
■ irritation or redness develops and lasts.

Keep out of reach of children ■ In case of
accidental ingestion, get medical help or contact
a Poison Control Center immediately.

Directions ■ Apply a small amount to palm
and rub hands together thoroughly.

Inactive Ingredients: water (Aqua), Cyclopentasiloxane,
Glycine Soja (Soybean) Oil, Glycerol, Cetanol Alcohol,
Ceteareth-20, Glyceryl Stearate, Dimethicone, Cetyl
Alcohol, Stearyl Alcohol, Caprylic/Capric Triglyceride,
Ethylhexyl Palmitate, Dicaprylyl Carbonate, Theobroma
Cacao (Cocoa) Seed Butter, Fragrance (Parfum), Helianthus
Annuus (Sunflower) Seed Extract, Ethylhexyl Methoxy-
cinamate, Butyl Methoxybenzoylmethane, Ethylhexyl
Salicylate, PEG-26-Buteth-26, PEG-40 Hydrogenated
Castor Oil, DMDM Hydantoin, Menthyl Cellulose,
Hydroxypropyl Methacrylate, Mica, Titanium Oxide
(CI 77891), Iron Oxides (CI 77491), Iron Oxides
(CI 77492), Red 40 (CI 16035), Yellow 5 (CI 19140).

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or your money back.
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ANTIBACTERIAL MOISTURIZING HAND LOTION SUN BLOSSOM

benzalkonium chloride lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:59 779-339
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM- UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.1 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
CYCLOMETHICONE 5 (UNII: 0THF5PCI0R)	
SOYBEAN OIL (UNII: 241ATL177A)	
GLYCERIN (UNII: PDC6A3C0OX)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
POLYOXYL 20 CETOSTEARYL ETHER (UNII: YRC528SWUY)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
STEARYL ALCOHOL (UNII: 2KR89I4HIY)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
ETHYLHEXYL PALMITATE (UNII: 2865993309)	
DICAPRYLYL CARBONATE (UNII: 609A3V1SUA)	
COCOA BUTTER (UNII: 512OYT1CRR)	
SUNFLOWER SEED (UNII: R9N3379M4Z)	
MICA (UNII: V8A1AW0880)	
OCTINOXATE (UNII: 4Y5P7MUD51)	
AVOBENZONE (UNII: G63QQF2NOX)	
OCTISALATE (UNII: 4X49Y0596W)	
POLYOXYL 40 CASTOR OIL (UNII: 4ERD2076EF)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
MANNITOL (UNII: 3OWL53L36A)	
POWDERED CELLULOSE (UNII: SMD1X3XO9M)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
CHROMIC OXIDE (UNII: X5Z09SU859)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59779-339-03	72 mL in 1 TUBE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	05/18/2011	

Labeler - CVS PHARMACY (062312574)

Registrant - APOLLO HEALTH AND BEAUTY CARE (201901209)

Establishment

Name	Address	ID/FEI	Business Operations
APOLLO HEALTH AND BEAUTY CARE		201901209	manufacture

Revised: 5/2011

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