LUBRIDERM DAILY MOISTURE WITH SUNSCREEN BROAD SPECTRUM SPF15- avobenzone, octisalate, octocrylene, oxybenzone lotion
Johnson & Johnson Consumer Inc.

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.

Lubriderm® Daily Moisture LOTION with Sunscreen BROAD SPECTRUM SPF 15

Drug Facts

Active ingredients

Avobenzone (2%), Octisalate (4%)
Octocrylene (3%), Oxybenzone (2.2%)

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 a doctor if rash occurs

- **Keep out of reach of children**. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

For sunscreen use:

- apply liberally 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 of age: Ask a doctor.

Other information

- protect this product from excessive heat and direct sun.

- may stain some fabrics

Inactive ingredients

Water, Glycerin, C12-15 Alkyl Benzoate, Cetyl Alcohol, Glyceryl Stearate, Cetearyl Alcohol, Stearic Acid, Phenoxyethanol, Ceteareth-20, Triethanolamine, Disodium EDTA, Fragrance, Panthenol, Tocopheryl Acetate, Xanthan Gum, Carbomer, Benzalkonium Chloride, Acrylates/C10-30 Alkyl Acrylate Crosspolymer ,Methylisothiazolinone

Questions?

Call toll-free **877-543-8477** or **215-273-8755** (collect) or visit www.lubriderm.com

Distributed by:
JOHNSON & JOHNSON
CONSUMER INC.
Skillman, NJ 08558

PRINCIPAL DISPLAY PANEL - 400 mL Bottle Label

Lubriderm®
dermatologist developed

Daily
Moisture
LOTION

with
Sunscreen
BROAD SPECTRUM SPF 15

24® CLINICALLY SHOWN TO
MOISTURIZE FOR 24 HOURS

CLEAN, NON-GREASY FEEL

13.5 Fl. Oz. (400 mL)

Lubriderm®

dermatologist developed

Daily Moisture LOTION™

with
Sunscreen

BROAD SPECTRUM SPF 15

CLINICALLY SHOWN TO
MOISTURIZE FOR 24 HOURS

CLEAN, NON-GREASY FEEL

13.5 Fl. Oz. (400 mL)

30002228

Lubriderm®

We believe everyone deserves to have healthy, comfortable skin.

LUBRIDERM®, the brand developed by dermatologists, offers specialized formulations with essential nutrients naturally found in healthy skin.

Daily Moisture Lotion with Sunscreen Broad Spectrum SPF 15

DAILY MOISTURE LOTION with Sunscreen Broad Spectrum SPF 15 is scientifically shown to help provide protection from the sun's harmful UVA and UVB rays. This non-greasy, fast-absorbing formula, enriched with Vitamin B5 and skin essential moisturizers, is clinically shown to moisturize for 24 hours for healthier-looking skin.

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Skillman, NJ 08558
Made in Canada

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The raised dots configuration and ribbon design are trademarks.

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LUBRIDERM DAILY MOISTURE WITH SUNSCREEN BROAD SPECTRUM SPF15

avobenzone, octisalate, octocrylene, oxybenzone lotion

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69968-0305
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Avobenzone (UNII: G63QQF2NOX) (Avobenzone - UNII:G63QQF2NOX)	Avobenzone	20 mg in 1 mL
Octisalate (UNII: 4X49Y0596W) (Octisalate - UNII:4X49Y0596W)	Octisalate	40 mg in 1 mL
Octocrylene (UNII: 5A68WGF6WM) (Octocrylene - UNII:5A68WGF6WM)	Octocrylene	30 mg in 1 mL
Oxybenzone (UNII: 95OOS7VE0Y) (Oxybenzone - UNII:95OOS7VE0Y)	Oxybenzone	22 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POLYOXYL 20 CETOSTEARYL ETHER (UNII: YRC528SWUY)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TROLAMINE (UNII: 9O3K93S3TK)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
XANTHAN GUM (UNII: TTV12P4NEE)	
CARBOXYPOLYMETHYLENE (UNII: 0A5MM307FC)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
PANTHENOL (UNII: WV9CM0O67Z)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:69968-0305-3	400 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	07/01/2010	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	07/01/2010	

Labeler - Johnson & Johnson Consumer Inc. (002347102)

Revised: 8/2016 Johnson & Johnson Consumer Inc.