DAILY MOISTURIZING WITH COLLOIDAL OATS- cetyl alcohol 0.80% lotion
Shanghai Lantern Cosmetic Co

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.

ETYL ALCOHOL 0.80%

PURPOSE

Skin Protectant

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

INDICATION AND USAGE

Relieves dry, itchy skin and provides hydration.

DIRECTIONS

Adults and children over two years of age. Apply up to 3-4 times a day.

Chilldren under two years of age, consult a physician.

WARNING

- For external use only - hands
- Stop use and contact a doctor if condition worsens and does not improve in seven days.
- Avoid contact with eyes. In case of contact with eyes, flush thoroughly with water

INACTIVE INGREDIENTS

WATER

GLYCERIN

DISTEARYLDIMONIUM CHLORIDE

PETROLATUM

ISOPROPYL PALMITATE

AVENA SATIVA (OAT) KERNEL FLOUR

BENZYL ALCOHOL

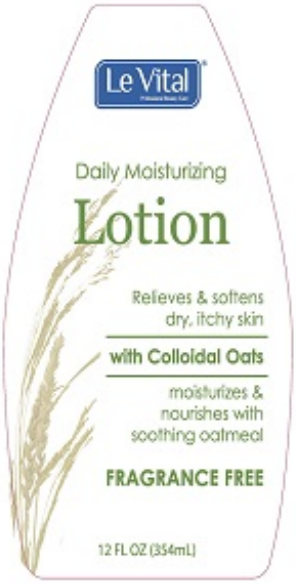
AVENA SATIVA (OAT) KERNEL OIL

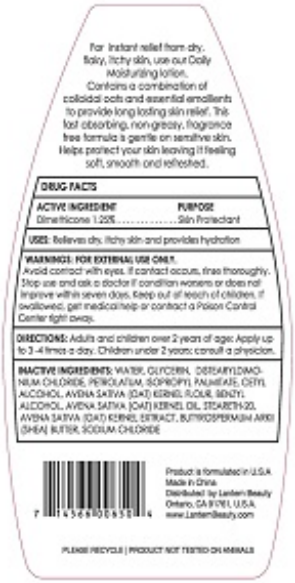
STEARETH-20

AVENA SATIVA (OAT) KERNEL EXTRACT

BUTYROSPERMUM PARKII (SHEA BUTTER)

SODIUM CHLORIDE





DAILY MOISTURIZING WITH COLLOIDAL OATS

cetyl alcohol 0.80% lotion

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50306-010
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CETYL ALCOHOL (UNII: 936JST6JCN) (CETYL ALCOHOL - UNII:936JST6JCN)	CETYL ALCOHOL	80 mg in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
AVENA SATIVA FLOWERING TOP (UNII: MA9CQJ3F7F)	
STEARETH-20 METHACRYLATE (UNII: A268NZ57NB)	
SODIUM CHLORIDE NA-22 (UNII: VMP9781061)	
WATER (UNII: 059QF0KO0R)	
.ALPHA.-(.ALPHA.-AMINOPROPYL)BENZYL ALCOHOL (UNII: S8TT5K3C8Y)	
GLYCERIN (UNII: PDC6A3C0OX)	
DISTEARYLDIMONIUM CHLORIDE (UNII: OM9573ZX3X)	

SHEA BUTTER (UNII: K49155WL9Y)	
PETROLATUM (UNII: 4T6H12BN9U)	
ISOPROPYL PALMITATE (UNII: 8CRQ2TH63M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50306-010-51	354 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/16/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/16/2016	

Labeler - Shanghai Lantern Cosmetic Co (421252043)

Establishment

Name	Address	ID/FEI	Business Operations
Shanghai Lantern Cosmetic Co		421252043	manufacture(50306-010)

Revised: 5/2016 Shanghai Lantern Cosmetic Co