OXY STEM M SERUM- dimethicone cream ULAB

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.

ACTIVE INGREDIENT

Active ingredients: DIMETHICONE 2.0%

INACTIVE INGREDIENT

Inactive ingredients: AQUA (WATER), Propylene Glycol, Glycerin, PEG-7 Glyceryl Cocoate, Water (aqua), Pentylene Glycol, Tocopheryl Acetate, Hydrolyzed Pea, PEG-40 Hydrogenated Castor Oil, Trideceth-9, Glycerin, Retinyl Palmitate, Carbomer, Triethanolamine, Fructose, Urea, Citric Acid, Phenoxyethanol, Ethylhexylglycerin, Sodium Hydroxide, Glycoproteins, Glutamic Acid, Valine, Threonine, Maltose, Sodium PCA, Sodium Chloride, Sodium Lactate, Trehalose, Allantoin, Sodium Hyaluronate, Glucose

PURPOSE

Purpose: Anti-fatigue and lifting Serum

WARNINGS

Warnings: For external use only Avoid contact with eyes. Discontinue use if signs of irritation or rashes appear.

KEEP OUT OF REACH OF CHILDREN

KEEP OUT OF REACH OF CHILDREN

INDICATIONS & USAGE

Indications & Usage: Apply morning and night

DOSAGE & ADMINISTRATION

Dosage & Administration: Apply morning and night on clean face and neck prior to moisturizer

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL







Energy—boosting and anti-fatigue Serum for skin recovery.
Ensure high energy levels and optimize keratinocyte.
Enhance self—regeneration and skin tightening.

€50ml / 1.69 fl.oz.

OXY STEM M SERUM

dimethicone cream

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71276-070	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
DIMETHICO NE (UNII: 92RU3N3Y1O) (DIMETHICONE - UNII:92RU3N3Y1O)	DIMETHICONE	1.0 g in 50 mL	

Inactive Ingredients			
Ingredient Name	Strength		
WATER (UNII: 059QF0KO0R)			
Propylene Glycol (UNII: 6DC9Q167V3)			

	Packaging			
1	# Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:71276-070- 02	1 in 1 CARTON	03/02/2017	
	NDC:71276-070- 01	50 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part347	03/02/2017	

Labeler - ULAB (688976692)

Registrant - ULAB (688976692)

Establishment				
Name	Address	ID/FEI	Business Operations	
ULAB		688976692	relabel(71276-070)	

Establishment				
Name	Address	ID/FEI	Business Operations	
U-LAB Switzerland AG		485958743	manufacture(71276-070)	

Revised: 3/2017 ULAB