# ULTRA REPAIR- shea butter cream ULAB

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.

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#### **ACTIVE INGREDIENT**

Active ingredients: SHEA BUTTER 4.0%

## **INACTIVE INGREDIENT**

Inactive ingredients: Aqua (WATER), CAPRYLIC/CAPRIC TRIGLYCERIDE, ISOHEXADECANE, PEG-8 BEESWAX, MARIS AQUA (SEA WATER), MACADAMIA INTEGRIFOLIA SEED OIL, CETYL ALCOHOL, POLYACRYLAMIDE, PROPYLENE GLYCOL DIPELARGONATE, PHENOXYETHANOL, GLYCERYL STEARATE, PROPANEDIOL, C13-14 ISOPARAFFIN, HYDROXYACETOPHENONE, THEOBROMA GRANDIFLORUM SEED BUTTER, PROPYLENE GLYCOL, CETETH-20, TOCOPHERYL ACETATE, STEARETH-20, CENTELLA ASIATICA EXTRACT, LAURETH-7, GLYCERIN, PARFUM (FRAGRANCE), TOCOPHEROL, NYMPHAEA ALBA FLOWER EXTRACT, POTASSIUM SORBATE, CITRIC ACID, TROPOLONE, GARDENIA TAHITENSIS FLOWER EXTRACT, CUCURBITA PEPO (PUMPKIN) SEED EXTRACT, SODIUM HYDROXIDE, SODIUM BENZOATE

## PURPOSE

Purpose: Moisturizing and soothing sensitized skin

# **KEEP OUT OF REACH OF CHILDREN**

KEEP OUT OF REACH OF CHILDREN

# **INDICATIONS & USAGE**

Indications & Usage: Apply morning and night

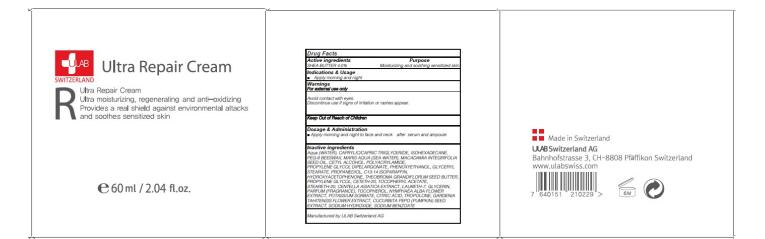
#### WARNINGS

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

#### **DOSAGE & ADMINISTRATION**

Dosage & Administration: Apply morning and night to face and neck after serum and ampoule

# PACKAGE LABEL.PRINCIPAL DISPLAY PANEL



U	LTRA REPA	AIR										
shea butter cream												
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Product Information												
Product T ype			HUMAN OTC DRUG Item Code (Source)			urce)	NDC:71276-090					
Route of Administration			TOPICAL									
A	ctive Ingredie	nt/Active Moi	ety									
							Strength Strength					
SHEA BUTTER (UNII: K49 155WL9 Y) (SHEA BUTTER - UNII:K49 155WL9 Y) SHEA BUTT								2.4 g in 60 mL				
	nactive Ingredi	ents	In and in the Name					Sauce walk				
TA	ATER (UNII: 059Q)		Ingredient Name				1	Strength				
			()									
ISOHEXADECANE (UNII: 918X1OUF1E)												
Р	ackaging											
#	Item Code		Package Description		N	farketing Sta Date	rt N	Aarketing End Date				
1	NDC:71276-090- 02	1 in 1 CARTON			03/0	2/2017						
1	NDC:71276-090- 01	60 mL in 1 BOTT Product	TLE, PUMP; Type 0: Not a Combi	natio n								
_												
Marketing Information												
N	larketing Catego	ry Applicatio	n Number or Monograph Citation		Marketing Start Date		te Mar	Marketing End Date				
ur	approved drug othe	r		0	3/02/2	0 17						

# **Registrant -** ULAB (688976692)

Establishment											
Name	Address	ID/FEI		Business Operations							
ULAB		688976692		relabel(712	76-090)						
Establishment											
	Name	Address	ID/	FEI	<b>Business Operations</b>						
U-LAB Switzerland AG			485958743	}	manufacture(71276-090)						

Revised: 3/2017

ULAB