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.

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

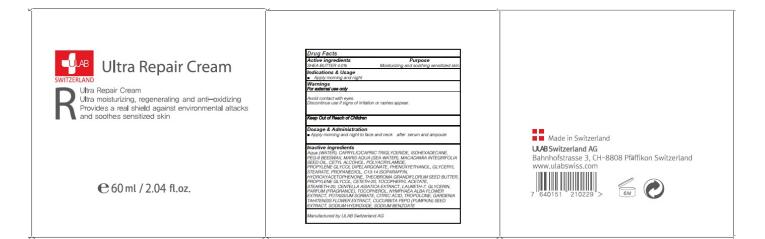
WARNINGS

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	Business Operations						
U-LAB Switzerland AG			485958743	}	manufacture(71276-090)						

Revised: 3/2017

ULAB