ULTRA REJUVE 7S AMPOULE- glycerin liquid
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: GLYCERIN 0.97%

INACTIVE INGREDIENT

Inactive ingredients: AQUA (WATER), PROPYLENE GLYCOL, PANTHENOL, PENTYLENE GLYCOL, PHENOXYETHANOL, HYDROLYZED WHEAT PROTEIN, CAPRYLYL/CAPRYL GLUCOSIDE, CENTELLA ASIATICA EXTRACT, ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER, PARFUM (FRAGRANCE), SACCHAROMYCES LYSATE, DISODIUM EDTA, TRITICUM VULGARE (WHEAT) SEED EXTRACT, ALGAE EXTRACT, SODIUM HYDROXIDE, o-CYMEN-5-OL, ETHYLHEXYLGLYCERIN, DISODIUM SUCCINATE, GLUTAMIC ACID, GLYCINE, THREONINE, VALINE, CITRIC ACID, POTASSIUM SORBATE, SODIUM BENZOATE, CI 19140 (FD&C YELLOW N°5)

PURPOSE

Purpose: Skin rebalancing and nourishing

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

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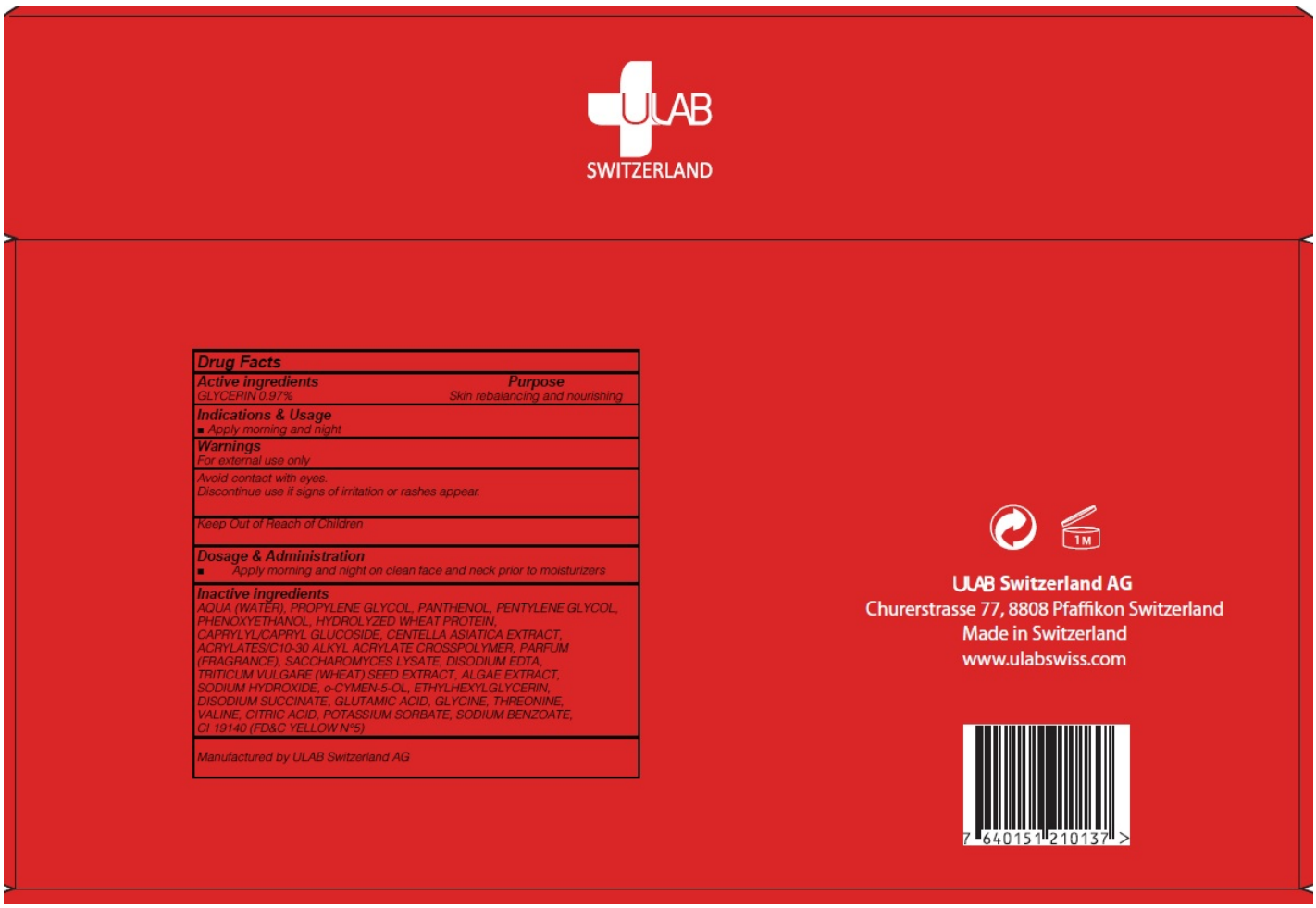
INDICATIONS & USAGE

Indications & Usage: Apply morning and night

DOSAGE & ADMINISTRATION

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

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL



ULTRA REJUVE 7S AMPOULE

glycerin liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71276-120
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
GLYCERIN (UNII: PDC6A3C0OX) (GLYCERIN - UNII:PDC6A3C0OX)	GLYCERIN	0.04 g in 5 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**

1	NDC:71276-120-02	5 in 1 CARTON	03/02/2017	
1	NDC:71276-120-01	5 mL in 1 VIAL; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		03/02/2017	

Labeler - ULAB (688976692)

Registrant - ULAB (688976692)

Establishment

Name	Address	ID/FEI	Business Operations
ULAB		688976692	relabel(71276-120)

Establishment

Name	Address	ID/FEI	Business Operations
U-LAB Switzerland AG		485958743	manufacture(71276-120)

Revised: 3/2017 ULAB