NYMPHSYN EYE- glycerin 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: GLYCERIN 0.97%

INACTIVE INGREDIENT

Inactive ingredients: AQUA (WATER), DICAPRYLYL ETHER, BUTYLENE GLYCOL, STEARIC ACID, GLYCERYL STEARATE, PEG-100 STEARATE, ETHYLHEXYL PALMITATE, PHENOXYETHANOL, CARBOMER, CI 77891 (TITANIUM DIOXIDE), SODIUM POTASSIUM ALUMINUM SILICATE, CETEARYL ALCOHOL, ETHYLHEXYLGLYCERIN, CAFFEINE, POTASSIUM CETYL PHOSPHATE, SODIUM HYDROXIDE, DISODIUM EDTA, MICA, PEG-20 STEARATE, PPG-25-LAURETH-25, SILICA, PEG-8, TRITICUM VULGARE (WHEAT) PROTEIN, PENTYLENE GLYCOL, PALMITOYL DIPEPTIDE-5 DIAMINOBUTYROYL HYDROXYTHREONINE, PALMITOYL DIPEPTIDE-5 DIAMINOHYDROXYBUTYRATE, TOCOPHEROL, SILICA DIMETHYL SILYLATE, ASCORBYL PALMITATE, NICOTIANA SYLVESTRIS LEAF CELL CULTURE, CAPRYLYL GLYCOL, ASCORBIC ACID, CITRIC ACID, SODIUM HYALURONATE, HEXYLENE GLYCOL, SODIUM DEHYDROACETATE, SORBIC ACID, UNDECYLENIC ACID

PURPOSE

Purpose: Restore firmness and tonicity of eye contour area

WARNINGS

Warnings: For external use only Discontinue use if signs of irritation or rashes appear.

KEEP OUT OF REACH OF CHILDREN

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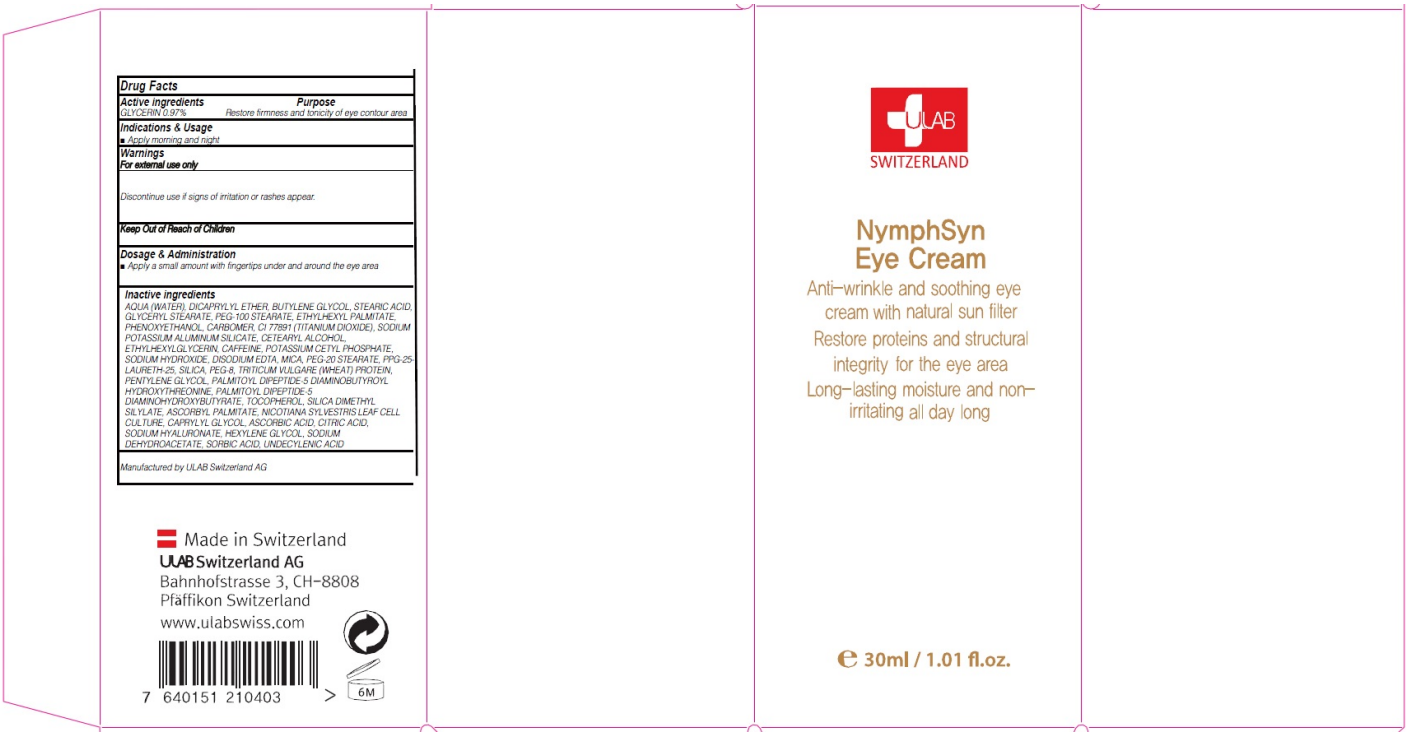
INDICATIONS & USAGE

Indications & Usage: Apply morning and night

DOSAGE & ADMINISTRATION

Dosage & Administration: Apply a small amount with fingertips under and around the eye area

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL



NYMPHSYN EYE

glycerin cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71276-040
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
GLYCERIN (UNII: PDC6A3C0OX) (GLYCERIN - UNII:PDC6A3C0OX)	GLYCERIN	0.29 g in 30 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
DICAPRYLYL ETHER (UNII: 77JZM5516Z)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:71276-040-02	1 in 1 CARTON	03/02/2017	
1	NDC:71276-040-01	30 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
unapproved drug other		03/02/2017	

Labeler - ULAB (688976692)

Registrant - ULAB (688976692)

Establishment

Name	Address	ID/FEI	Business Operations
ULAB		688976692	relabel(71276-040)

Establishment

Name	Address	ID/FEI	Business Operations
U-LAB Switzerland AG		485958743	manufacture(71276-040)

Revised: 3/2017 ULAB