ULTRA MOISTURIZING ESSENCE- glycerin essence solution
NANONATURE

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

Glycerin (12.3%) - Skin Protectant

INACTIVE INGREDIENT

Water, Olein Acid, Lemon Balm Extract, Olive Oil, Angelica Gigas NAKAI Extract, Chinese Matrimony Vine Extract, Ginseng Extract, Retinol, Tocopherol, Ascobic Acid, Aromatic

PURPOSE

Skin Protectant

WARNINGS

For skin use only.
Do not use
 on puncture wounds on animal bites
When using this product
 avoid contact with eyes
Stop use and ask a doctor if
 you have allergy or skin disease
 rash or irritation develops and lasts
 red spots, swelling, itching, irritation and other abnormalities developsFor skin use only.
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 you have allergy or skin disease
 rash or irritation develops and lasts
 red spots, swelling, itching, irritation and other abnormalities develops

KEEP OUT OF REACH OF CHILDREN

 if swallowed, get medical help or contact a Poison Control Center right away

INDICATIONS & USAGE

 Clean skin thoroughly before using
 Squeeze 1 or 2 pumps onto palm
 Apply lightly to a slightly wet face
 Frequency of usage should accord to skin condition

DOSAGE & ADMINISTRATION

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 Squeeze 1 or 2 pumps onto palm
 Apply lightly to a slightly wet face
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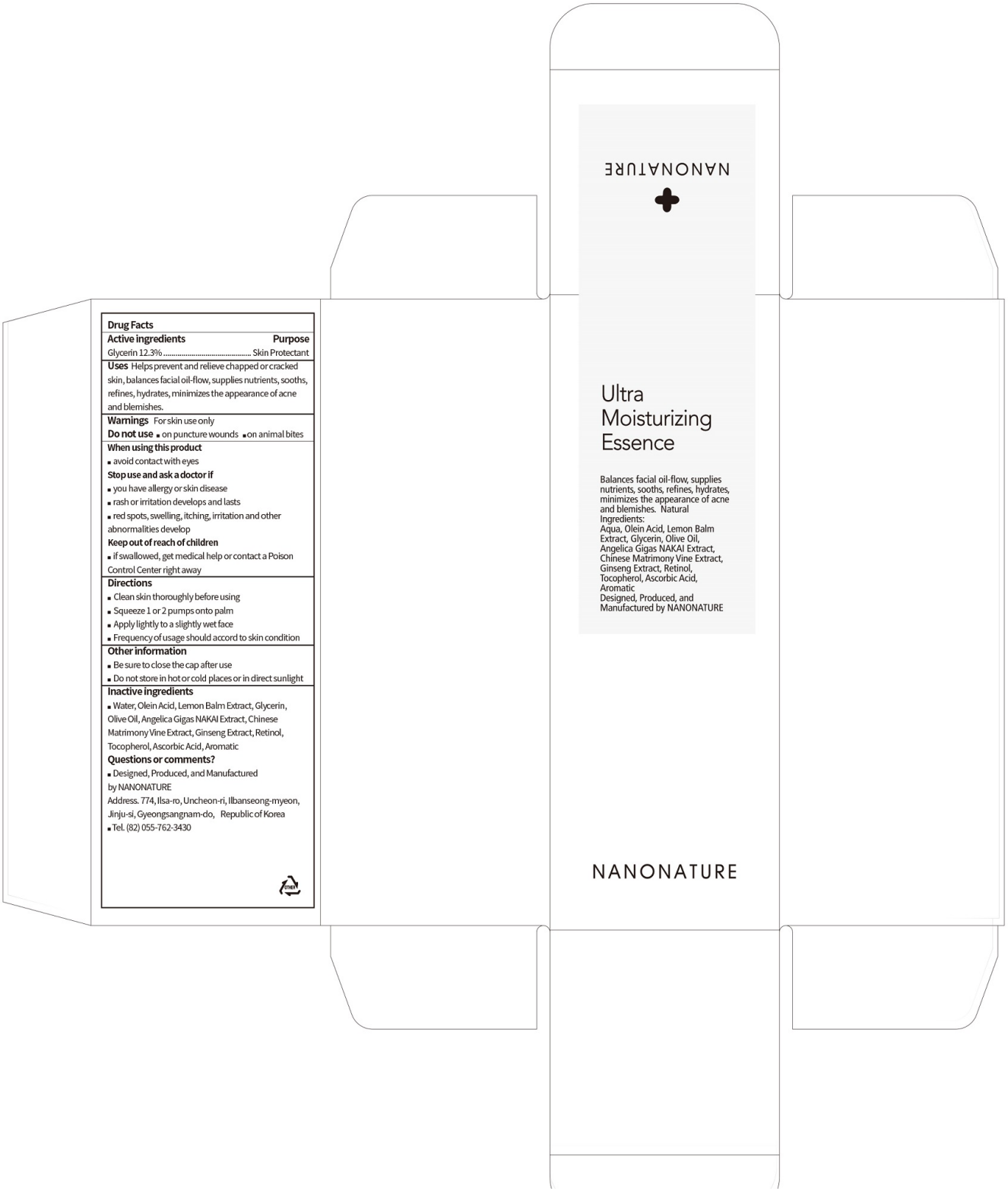
PACKAGE LABEL.PRINCIPAL DISPLAY PANEL



NANONATURE

ULTRA MOISTURIZING ESSENCE

glycerin essence solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71538-002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Glycerin (UNII: PDC6A3C0OX) (GLYCERIN - UNII:PDC6A3C0OX)	Glycerin	12.3 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
Water (UNII: 059QF0KO0R)	
Olive Oil (UNII: 6UYK2W1W1E)	
Tocopherol (UNII: R0ZB2556P8)	
Retinol (UNII: G2SH0XKK91)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:71538-002-01	100 mL in 1 PACKAGE; Type 0: Not a Combination Product	07/17/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		07/17/2017	

Labeler - NANONATURE (689851291)

Registrant - NANONATURE (689851291)

Establishment

Name	Address	ID/FEI	Business Operations

NANONATURE		689851291	manufacture(71538-002)