AN ADC SP INTENSIVE MOISTURE ESSENCE- niacinamide cream AN Co Ltd.

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 Ingredient: Niacinamide 2.0%

INACTIVE INGREDIENT

Inactive Ingredients: Water, Dipropylene Glycol, Phytosqualane, Glycerin, Phenyl Trimethicone, Panthenol, Cyclopentasiloxane/PEG-10 Dimethicone/Vinyl Dimechicone Cross Polymer, Cyclopentasiloxane, Hydrogenated Lecithin, 1,2-Hexanediol, Polyglyceryl-3 Methylglucose Distearate, Polyacrylamide/C13-14 Isoparaffin/ Laureth-7, CeramideNP/Hydrogenated Polydecene/Stearic Acid/ButyrospermumParkii(Shea) Butter/Ceteareth-20//Glyceryl Citrate/Lactate/Linoleate/Oleate Butylene glycol, Ethoxydiglycol, Trimethylpentanediol/Adipic Acid Copolymer, Sodium Polyacrylate, Phenoxyethanol, Beta-Glucan, Tocopheryl Acetate, Carbomer, Triethanolamine, Fragrance, Sodium Hyaluronate, Sodium Stearoyl Glutamate, Xanthan Gum, Disodium EDTA, Copper Tripeptide-1, Acetyl Hexapeptide-8, Palmitoyl Pentapeptide-4

PURPOSE

Purpose: Skin Protectant

WARNINGS

Precautions on use: 1. Stop using the product if there are any of the following abnormal symptoms appearing after use and consult with your dermatologist as continued use can worsen the symptoms. A) If there are red spots, swelling, itchiness, or irritation B) If the above symptoms appear around the skin to which the product has been applied after being exposed to direct sunlight. 2. Precautions on storage and handling A) Make sure to close the cap after use. B) Keep out of the reach of children. C) Do not keep the product in a hot or cold place or a place getting direct sunlight. 3. Do not apply the product to any parts of the skin with wound, eczema, or dermatitis.

KEEP OUT OF REACH OF CHILDREN

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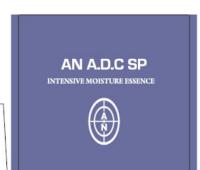
Suggested use

Suggested use: Nourish the skin with anti-wrinkle eye cream and apply an appropriate amount of the essence to the whole face and neck gently and evenly.

DOSAGE & ADMINISTRATION

Dosage & administration: Apply 3~4g of essence to face evenly.

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL



AN ADC SP

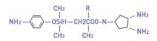
Intensive Moisturizing Essence Functional whitening cosmetics

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(주)에이앤 www.alinationss.com 본사: 경기도 수원시 장안구 팔달로 259번길35 AN빌딩 고객센터 : 1899-6859 공장 : 충청북도 영동군 영동황간로 1452-248

MADE IN KOREA



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- inflation

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 * Apply 1--2g of cream to the skin around eye once a day.

본 제품에 이상이 있을 경우 공정거래위원회 고시 "소비자 분쟁 해결 기준" 에 의해 보상해 드립니다.







INTENSIVE MOISTURE ESSENCE

All Nations A.D.C Special Program

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NET WT. 55q



Drug Facts

Active Ingredient: Niacinamide2,0%

Inactive Ingredients: Water, Dipropylene Glycol, Phytosqualane, Glycerin, Phenyl Trimethicone, Panthenol, Cyclopentasiloxane/PEG-10 Dimethicone/Vinyl Direchicone Cross Polymer, Cyclopentasiloxane, Hydrogenated Lecithin, 1,2-Hexanediol, Polyglyceryl-3 MethylglucoseDistearate, Polyacrylamide/Ct3-14 Isoparaffin/ Laureth-7, CeramideNP/Hydrogenated Polydecene/Stearic Acid/BulyrospermumParkii(Shea) Butter/Ceteareth-20//Givcervi Citrate/Lactate/Linoleate/Oleate Butylene glycol, Ethoxydiglycol, Trimethylpentanediol/Adipic Acid Copolymer, Sodium Polyacrylate Phenoxyethanol, Beta-Glucan, Tocopheryl Acetate, Carbomer, Triethanolamine, Fragrance, Sodium Hyaluronate, Sodium Stearoyl Glutamate, Xanthan Gum, Disodium EDTA, Copper Tripeptide-1, Acetyl Hexapeptide-8, Palmitoyl Pentapeptide-4

AN ADC SP INTENSIVE MOISTURE ESSENCE

niacinamide cream

Product Information

HUMAN OTC DRUG NDC:69153-080 Product Type Item Code (Source)

TOPICAL Route of Administration

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Niacinamide (UNII: 25X5118RD4) (NIACINAMIDE - UNII:25X5118RD4)	Niacinamide	1.10 g in 55 g

Inactive Ingredients

Ingredient Name	Strength

Water (UNII: 059QF0KO0R)	
Dipropylene Glycol (UNII: E107L85C40)	

ı	Packaging			
	# Item Code	Package Description	Marketing Start Date	Marketing End Date
	1 NDC:69153-080- 02	1 in 1 CARTON	05/01/2016	
	1 NDC:69153-080- 01	55 g 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		05/01/2016	

Labeler - AN Co Ltd. (688448454)

Registrant - AN Co Ltd. (688448454)

Establishment			
Name	Address	ID/FEI	Business Operations
AN Co Ltd.		688448454	manufacture(69153-080)

Revised: 6/2016 AN Co Ltd.