AN ADC SP INTENSIVE WRINKLE- dimethicone cream AN Co Ltd.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

ACTIVE INGREDIENT

Active Ingredient: Dimethicone 1.3%

INACTIVE INGREDIENT

Inactive Ingredients: Water, Butylene Glycol, Cyclopentasiloxane/PEG-10 Dimethicone/Vinyl Dimechicone Cross Polymer, Dipropylene Glycol, Olive Oil Glycereth-8 Esters, Panthenol, Cetyl Ethylhexanoate, Phytosqualane, 1,2-Hexanediol, Cetyl Alcohol, Glyceryl Stearate, Limnanthes Alba (Meadowfoam) Seed Oil, Polyacrylamide/C13-14 Isoparaffin/ Laureth-7, Polysorbate 60, Ethoxydiglycol, Trimethylpentanediol/Adipic Acid Copolymer, CeramideNP/Hydrogenated Polydecene/Stearic Acid/ButyrospermumParkii(Shea) Butter/Ceteareth-20/Glyceryl Citrate/Lactate/Linoleate/Oleate Butylene glycol, Phenoxyethanol, Beta-Glucan, Dipotassium Glycyrrhizate, Fragrance, Sodium Stearoyl Glutamate, Hydrogenated Lecithin, Carbomer, Triethanolamine, Sodium Polyacrylate, Tocopheryl Acetate, Xanthan Gum, Adenosine, Disodium EDTA, Sodium Hyaluronate, Copper Tripeptide-1, Palmitoyl Pentapeptide-4, Acetyl Hexapeptide-8

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

KEEP OUT OF REACH OF CHILDREN

Suggested use

Suggested use: At the final stage of basic makeup, apply an appropriate amount of the cream to the whole face and gently massage the face upwards to help the skin absorb the nutrients.

DOSAGE & ADMINISTRATION

Dosage & administration: Apply 3~4g of cream to the whole face evenly.

PACKAGE LABEL, PRINCIPAL DISPLAY PANEL



AN ADC AP

Intensive Wrinkle Cream Functional anti-wrinkle cosmetics

Suggested use: At the final stage of basic makeup, apply an appropriate amount of the cream to the whole face and gently massage the face upwards to help the skin absorb the nutrients,

*Apply 3~4g of cream to the whole face

NET WT. 50a

(주)에이앤 www.allnationss.com 본사: 경기도 수원시 장안구 필달로 259번길35 AN빌딩 고객센터: 1899—8859 공장: 충청북도 영동군 영동환간로 1452—248

MADE IN KOREA

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







INTENSIVE WRINKLE CREAM

All Nations A.D.C Special Program

Chara

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Drug Facts
Active Ingredient: Dimethicone1.3%

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Inactive Ingredients: Water, Bulylene Glycol,
Cyclopertasiakoane/PEG-10 Dimehicone/Ninyl Dimechicone Cross
Polymer, Dipropylene Glycol, Olive Oli Glycerellt-B Esites, Parthenol,
Cely/Elthylensone, Phytosqualen, 12-Hesanecko, Celyl Alcohol,
Glycery Steerate, Limnarthes Alba (Meadowbam) Seed Oli,
Polyacrystamér CG-3-4 lacparathic Juareth-7, Polyacrystamér CG-20 (Polyacrystamer)

Etmosylajokol, Timnethylepentanedio/Adaptic Acid Copolymer,
Cerandele/Phylogensied Alcoholin, Carbomer, Timethanokamine,
Sodum Polyacryste, Ticopheryl Acetale, Xanthan Gum, Adenosine,
Sodum Billa, Sodum Hysiumonak, Copper Tirpeptde-1, Palmitoyl
Pertapaptide-1, Acetyl Hexapoptide-8

AN ADC SP INTENSIVE WRINKLE

dimethicone cream

Product Information

HUMAN OTC DRUG NDC:69153-100 Product Type Item Code (Source)

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Basis of Strength Strength **Ingredient Name** Dimethicone (UNII: 92RU3N3Y1O) (DIMETHICONE - UNII:92RU3N3Y1O) Dimethicone $0.65\,g\ in\,50\,g$

Inactive Ingredients

Ingredient Name Strength Water (UNII: 059QF0KO0R) Butylene Glycol (UNII: 3XUS85K0RA)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:69153-100- 02	1 in 1 CARTON	05/01/2016			
1	NDC:69153-100- 01	50 g in 1 BOTTLE, GLASS; Type 0: Not a Combination Product				

Marketing Information							
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date				
OTC monograph final	part347	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(69153-100)				

Revised: 6/2016 AN Co Ltd.