DR. GROOT ANTI-DANDRUFF SCALP CARE- salicylic acid shampoo TAI GUK PHARM. 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.

Dr. Groot Anti-dandruff Scalp Care Shampoo

Drug Facts

Active ingredient

Salicylic Acid 3%

Purposes

Anti-dandruff

Uses

controls flaking, scaling and itcing associated with dandruff

Warnings

For external use only

When using this product

- avoid contact with the eyes
- if contact occurs, rinse eyes thoroughly with water

Stop use and ask a doctor if

■ condition worsens or does not improve after regular use of this product as directed.

Keep out of reach of children if swallowed, get medical help or contact a Poison Control Center right away

Directions

■ shake well before use ■ wet hair, massage onto scalp, rinse, repeat if desired ■ for best results use at least twice a week or as directed by a doctor ■ for maximum dandruff control, use every time you shampoo

Other information

■ Store at room temperature

Inactive Ingredients

water, sodium C14-16 olefin sulfonate, cocamidopropyl betaine, disodium cocoyl glutamate, sodium chloride, polyquaternium-7, fragrance, sodium cocoyl glutamate, sodium benzoate, PEG-40 hydrogenated castor oil, menthol, tetrasodium EDTA, citric acid, biotin, caffeine, hexylene glycol, honey, niacinamide, panthenol, sodium citrate, polysorbate 20, alcohol, butylene glycol, propolis extract, houttuynia cordata extract, rosmarinus officinalis (rosemary) leaf extract, zingiber officinale (ginger) root extract

Questions?

U.S.A.: 1-800-FOR-AVON

www.avon.com

MADE IN KOREA

DISTR. LG H&H Co., Ltd. 58, SAEMUNAN-RO, JONGNO-GU, SEOUL, KOREA

PRINCIPAL DISPLAY PANEL - 300 mL Bottle Carton

Dr. Groot

ANTI DANDRUFF SCALP CARE SHAMPOO

SALICYLIC ACID 3%

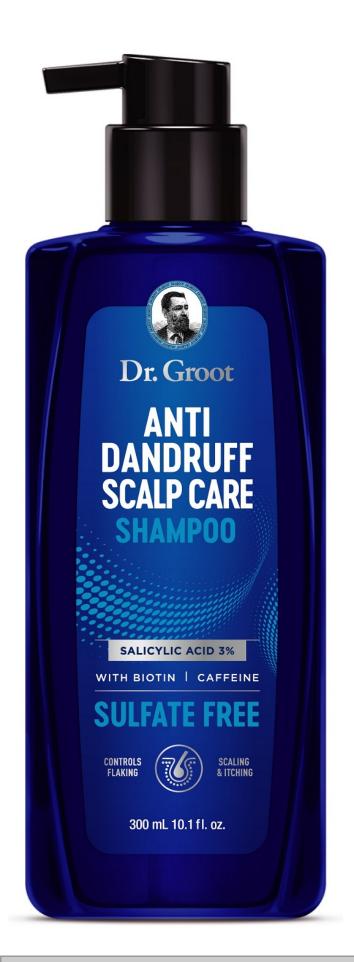
WITH BIOTIN | CAFFEINE

SULFATE FREE

CONTROLS FLAKING

SCALING & ITCHING

300 mL 10.1 fl. oz.





DR. GROOT ANTI-DANDRUFF SCALP CARE

Product Information

Product Type HUMAN OTC DRUG **Item Code (Source)** NDC:43136-701

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength	
0.4.455.4.553.4.6.4.6.6.5.4.6.5.4.6.5.4.6.5.4.5.5.4.5.5.4.5.5.4.6.5	6 44 46 44 46 4 64 5		

SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ) SALICYLIC ACID 30 mg in 1 mL

Inactive Ingredients

mactive mgreatenes		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
SODIUM C14-16 OLEFIN SULFONATE (UNII: O9W3D3YF5U)		
COCAMIDOPROPYL BETAINE (UNII: 50CF3011KX)		
SODIUM CHLORIDE (UNII: 451W47IQ8X)		

Packaging

ı	Packaging				
	# Item Code	Package Description	Marketing Start Date	Marketing End Date	
	1 NDC:43136-701-01	300 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/15/2022		

Marketing Information

	ia keeing in ormation			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph final	part358H	09/15/2002		

Labeler - TAI GUK PHARM. CO., LTD (689060246)

DISODIUM COCOYL GLUTAMATE (UNII: MBK0CP8F5A)

Establishment Address ID/FEI **Business Operations** TAI GUK PHARM. CO., LTD 689060246 manufacture(43136-701)

Revised: 9/2022 TAI GUK PHARM. CO., LTD