

CLARIFYING PEEL- salicylic acid gel
Skin Authority LLC

Skin Authority - CLARIFYING LIQUID PEEL (46007-102)

ACTIVE INGREDIENT

SALICYLIC ACID 2%

PURPOSE

ACNE TREATMENT

USES

HELPS ELIMINATE ACNE-CAUSING BACTERIA, CALMS INFLAMMATION, AND MINIMIZES EXCESS OIL.

WARNINGS

FOR EXTERNAL USE ONLY.

AVOID CONTACT WITH EYES AND EYELIDS. IF CONTACT OCCURS, RINSE EYES THOROUGHLY WITH WATER.

IF IRRITATION DEVELOPS, DISCONTINUE USE.

KEEP OUT OF REACH OF CHILDREN.

DIRECTIONS

APPLY WITH FINGERTIPS OR COTTON PAD IMMEDIATELY AFTER CLEANSING, COVERING FACE AND INFLAMED AREAS. CONTINUE WITH DAILY CARE ROUTINE. REAPPLY THROUGHOUT THE DAY TO REMOVE EXCESS OIL AND DIRT.

OTHER INGREDIENTS

WATER, GLYCOLIC ACID, HAMAMELIS VIRGINIANA (WITCH HAZEL) EXTRACT, SD ALCOHOL, SODIUM PCA, PPG-5-CETETH-20, SODIUM HYDROXIDE, ALOE BARBADENSIS LEAF JUICE, POLYSORBATE 80, PHENOXYETHANOL, CAPRYLYL GLYCOL, ETHYLHEXYLGLYCERIN, HEXYLENE GLYCOL, METHYLGLUCETH-20.

QUESTIONS?

+1-760-268-0047



Drug Facts	
Active Ingredients	Purposes
Salicylic Acid 2%	Acne treatment
Uses: Helps eliminate acne-causing bacteria, calms inflammation, and minimizes excess oil. <i>Aide à éliminer les bactéries qui causent l'acné, à calmer l'inflammation et à réduire l'excès de sébum.</i> • <i>Ayuda a eliminar las bacterias que causan el acné, calma la inflamación y minimiza la grasa excesiva.</i>	
Warnings: For external use only. Keep out of reach of children. Avoid contact with eyes and eyelids. If contact occurs, rinse eyes thoroughly with water. If irritation develops, discontinue use. <i>Pour usage externe seulement. Garder hors de la portée des enfants. Éviter tout contact avec les yeux et les paupières. En cas de contact, rincer abondamment les yeux à l'eau. En cas d'irritation, cesser l'utilisation.</i> • <i>Sólo para uso externo. Manténgalo fuera del alcance de los niños. Evite el contacto con los ojos y los párpados. Si hay contacto, enjuague abundantemente los ojos con agua. Si se presenta irritación, descontinúe el uso.</i>	
Directions: Apply with fingertips or cotton pad immediately after cleansing, covering face and inflamed areas. Continue with daily care routine. Resapply throughout the day to remove excess oil and dirt. <i>Immédiatement après le nettoyage, appliquez du bout des doigts ou à l'aide d'un tampon de coton sur tout le visage et les zones enflammées. Effectuez les autres étapes de la routine de soins quotidienne. Effectuez les autres étapes de la routine de soins quotidienne. Réappliquez au cours de la journée pour éliminer l'excès de sébum et de saleté.</i> • <i>Aplicarlo con las puntas de los dedos o una toalla de algodón inmediatamente después de limpiar, cubriendo el rostro y las áreas inflamadas. Continúa con su rutina diaria de cuidados. Cúbralo con su rutina diaria de cuidados. Aplique nuevamente durante el día para eliminar el exceso de aceite y suciedad.</i>	
Other Ingredients: Water, Glycolic Acid, Hamamelis Virginiana (Witch Hazel) Extract, SD Alcohol, Sodium PCA, PPG-5-Ceteth-20, Sodium Hydroxide, Aloe Barbadensis Leaf Juice, Polysorbate 80, Phenoxethanol, Capryl Glycol, Ethylhexylglycerin, Hexylene Glycol, Methylgluceth 20.	
Questions? +1-760-268-0047	

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 MADE IN USA #51101 (revE)



CLARIFYING PEEL

salicylic acid gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:46007-102
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SALICYLIC ACID (UNII: 0414PZ4LPZ) (SALICYLIC ACID - UNII:0414PZ4LPZ)	SALICYLIC ACID	2 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCOLIC ACID (UNII: 0WT12SX38S)	
HAMAMELIS VIRGINIANA TOP (UNII: UDA30A2JYY)	
ALCOHOL (UNII: 3K9958V90M)	
SODIUM PYRROLIDONE CARBOXYLATE (UNII: 469OTG57A2)	

PPG-5-CETETH-20 (UNII: 4AAN25P8P4)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
HEXYLENE GLYCOL (UNII: KEH0A3F75J)	
METHYL GLUCETH-20 (UNII: J3QD0LD11P)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:46007-102-12	50 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/19/2012	
2	NDC:46007-102-11	20 mL in 1 TUBE; Type 0: Not a Combination Product	04/19/2012	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M006	04/19/2012	

Labeler - Skin Authority LLC (136549396)

Revised: 12/2025

Skin Authority LLC