### LIQUID- salicylic acid liquid Oxygen Development LLC

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.

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### BE CLARIFIED Acne Gel-Cream Cleanser Salicylic Acid Acne treatment

#### **Active Ingredient**

Salicylic Acid

#### **Purpose**

Acne Treatment

#### Uses

For the treatment of acne

Helps prevent new acne blemishes from forming

## Warnings

For external use only

# When using this product

Skin irritation and dryness are more likely to occur if you use another topical acne medication at the same time. If irritation occurs, only use one topical acne treatment at a time.

Avoid contact with eyes. If products occurs, rinse eyes thoroughly with water

# Keep out of reach of children

If product is swallowed, get medical help or contact a poison control center immediately

#### **Directions**

Shake well before use

Start with damp skin

Apply a small amount onto your fingertips and emulsify

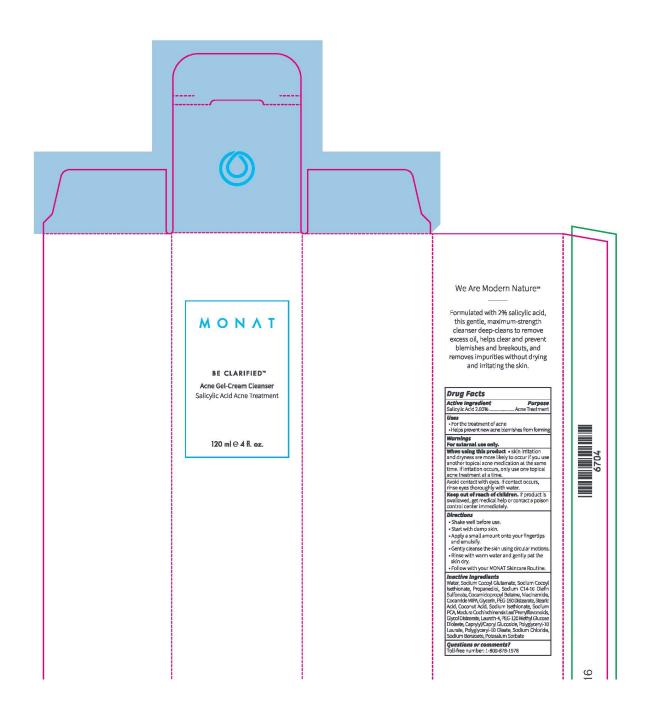
Gently cleanse the skin using circular motions

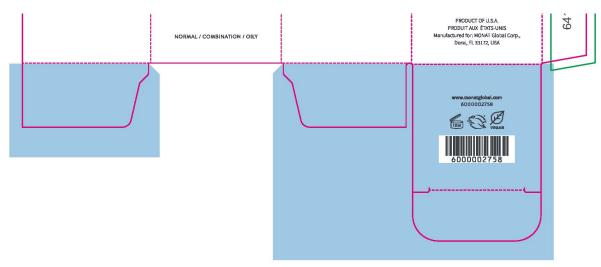
Rinse with warm water and gently pat the skin dry Follow with your MONAT Skincare Routine

# **Inactive Ingredients**

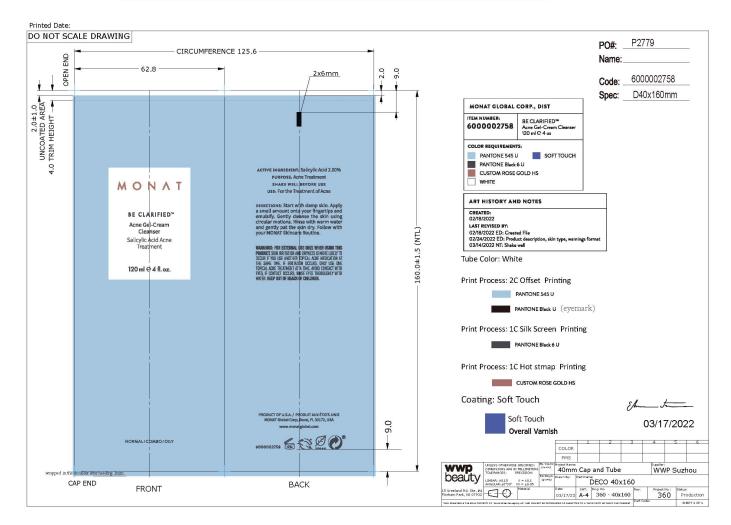
Water/Aqua, Sodium Cocoyl Glutamate, Sodium Cocoyl Isethionate, Propanediol, Sodium C14-16 Olefin Sulfonate, Cocamidopropyl Betaine, Niacinamide, Cocamide MIPA, Glycerin, PEG-150 Distearate, Stearic Acid, Coconut Acid, Sodium PCA, Maclura Cochinchinensis Leaf Prenylflavonoids, Glycol Distearate, Laureth-4, PEG-120 Methyl Glucose Dioleate, Caprylyl/Capryl Glucoside, Polyglyceryl-10 Laurate, Polyglyceryl-10 Oleate, Sodium Chloride, SOdium Benzoate, Potassium Sorbate

#### Package Label









# LIQUID

salicylic acid liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:61354-091
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	<b>Basis of Strength</b>	Strength	
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	2 mg in 100 mg	

Inactive Ingredients			
Ingredient Name	Strength		
SODIUM COCOYL GLUTAMATE (UNII: BMT4RCZ3HG)	4.2 mg in 100 mg		
PROPANEDIOL (UNII: 5965N8W85T)	2.99 mg in 100 mg		
SODIUM C14-16 OLEFIN SULFONATE (UNII: O9W3D3YF5U)	2 mg in 100 mg		
COCO MONOISOPROPANOLAMIDE (UNII: 21X4Y0VTB1)	0.56 mg in 100 mg		
COCAMIDOPROPYL BETAINE (UNII: 50CF3011KX)	1.79 mg in 100 mg		
SODIUM COCOYL ISETHIONATE (UNII: 518XTE8493)	1.68 mg in 100 mg		
WATER (UNII: 059QF0KO0R)	78.25 mg in 100 mg		
PEG-120 METHYL GLUCOSE DIOLEATE (UNII: YM0K64F20V)	1.087 mg in 100 mg		
SODIUM PYRROLIDONE CARBOXYLATE (UNII: 4690TG57A2)	0.5 mg in 100 mg		

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
<b>1</b> NDC:61354-091-	1 in 1 CARTON	03/22/2023	
1 NDC:61354-091-	1 mg in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	M006	03/22/2023	

# Labeler - Oxygen Development LLC (137098492)

Establishment				
Name	Address	ID/FEI	<b>Business Operations</b>	
Oxygen Development LLC		137098492	manufacture(61354-091)	

Revised: 3/2023 Oxygen Development LLC