ARTISTRY STUDIO DONE WITH ZIT ACNE TREATMENT CLEARING GEL- salicylic acid gel Access Business Group LLC

Artistry Studio Done With Zit! Acne Treatment + Clearing Gel

Drug Facts

Active ingredient

Salicylic Acid 2.0% (W/W)

Purpose

Acne Treatment

Use

• For the treatment of acne.

Warnings

For external use only

Keep out of reach of children.

If product is swallowed, get medical help or contact a Poison Control Center right away.

Using other topical acne medications at the same time or immediately following use of this product may increase dryness or irritation of the skin. If this occurs, only one product should be used unless directed by a doctor/health care practitioner.

Apply to affected areas only. Do not use on broken skin or apply tolarge areas of the body

Avoid contact with eyes. If contact occurs, rinse thoroughly with water. Discontinue use if excessive skin irritation develops or increases. If irritation persists, consult a doctor/health care practitioner.

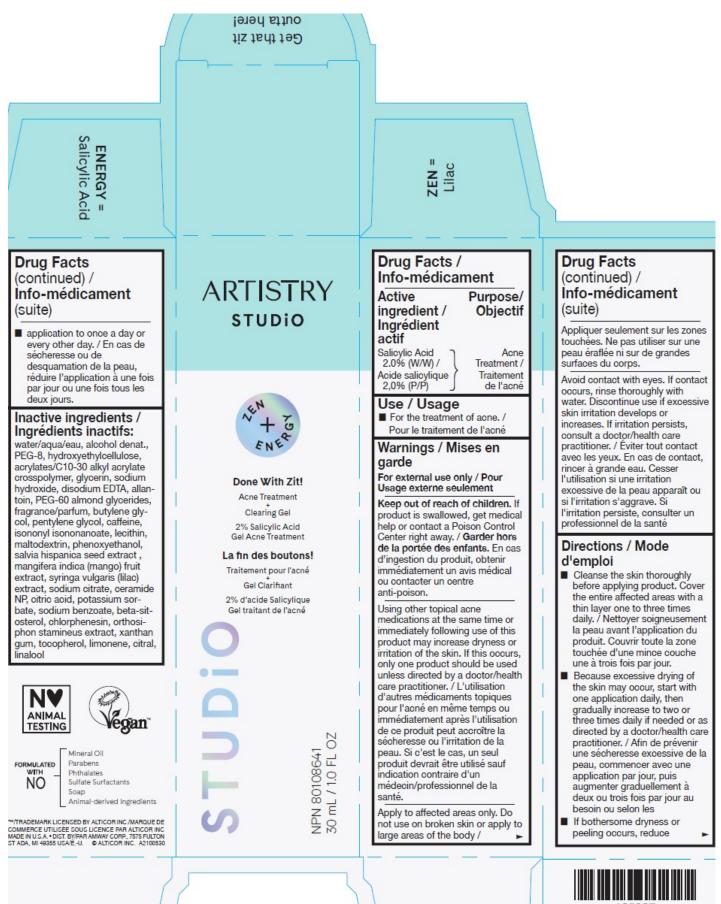
Directions

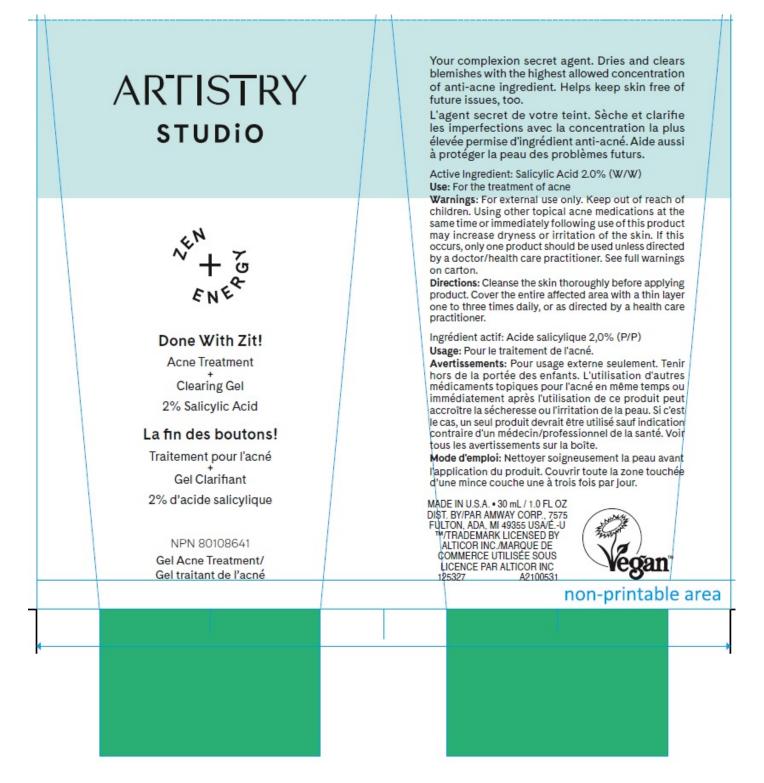
- Cleanse the skin thoroughly before applying product. Cover the entire affected areas with a thin layer one to three times daily.
- Because excessive drying of the skin may occur, start with one application daily, then gradually increase to two or three times daily if needed or as directed by a doctor/health care practitioner.
- If bothersome dryness or peeling occurs, reduce application to once a day or every other day.

Inactive ingredients

water/aqua/eau, alcohol denat., PEG-8, hydroxyethylcellulose, acrylates/C10-30 alkyl acrylate crosspolymer, glycerin, sodium hydroxide, disodium EDTA, allantoin, PEG-60 almond glycerides, fragrance/parfum, butylene glycol, pentylene glycol, caffeine, isononyl isononanoate, lecithin, maltodextrin, phenoxyethanol, salvia hispanica seed extract, mangifera indica (mango) fruit extract, syringa vulgaris (lilac) extract, sodium citrate, ceramide NP, citric acid, potassium sorbate, sodium benzoate, betasitosterol, chlorphenesin, orthosiphon stamineus extract, xanthan gum, tocopherol, limonene, citral, linalool

Package Labeling:





Circumference/Unrolling: 94,2 mm TUBE HEAD

ARTISTRY STUDIO DONE WITH ZIT ACNE TREATMENT CLEARING GEL

salicylic acid gel

Product Information

Product Type		HUMAN OTC DRUG	Item Code (S	Source)	NDC:1	10056-045			
Route of Admini	stration	TOPICAL							
Active Ingredient/Active Moiety									
Ingredient Name				Basis of Strength		Strength			
SALICYLIC ACID (UNII: 0414PZ4LPZ) (SALICYLIC ACID - UNII:0414PZ)414P74IP7)	SALICYLIC ACID		20 mg in 1 mL			
			,						
Inactive Ingre	dients								
		Ingredient Nam	e			Strength			
WATER (UNII: 059Q	F0KO0R)	ingreatent itali	C			Juciyu			
ALCOHOL (UNII: 3K9958V90M)									
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)									
HYDROXYETHYL CELLULOSE, UNSPECIFIED (UNII: T4V6TWG28D)									
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)									
GLYCERIN (UNII: PD		- ,			,				
		40C32I)							
	SODIUM HYDROXIDE (UNII: 55X04QC32I) EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)								
ALLANTOIN (UNII: 3445277G0Z)									
		JNII: 4Y0F651N0F)							
PEG-60 ALMOND GLYCERIDES (UNII: 4Y0E651N0F) BUTYLENE GLYCOL (UNII: 3XUS85K0RA)									
PENTYLENE GLYCOL (UNII: 50C1307PZG)									
CAFFEINE (UNII: 3G6A5W338E)									
ISONONYL ISONONANOATE (UNII: S4V5BS6GCX)									
MALTODEXTRIN (UNII: 7CVR7L4A2D)									
PHENOXYETHANOL (UNII: HIE492ZZ3T)									
CHIA SEED (UNII: NUOOLX06F8)									
MANGO (UNII: 1629I3NR86)									
SODIUM CITRATE	(UNII: 1Q73Q2JI	JLR)							
CERAMIDE NP (UNII: 4370DF050B)									
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)									
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)									
SODIUM BENZOAT	E (UNII: 0J2456	E5EU)							
.BETASITOSTEROL (UNII: S347WMO6M4)									
CHLORPHENESIN (UNII: I670DAL4SZ)									
XANTHAN GUM (UNII: TTV12P4NEE)									
TOCOPHEROL (UNII: R0ZB2556P8)									
LIMONENE, (+)- (UNII: GFD7C86Q1W)									
CITRAL (UNII: T7EU009VPP)									
LINALOOL, (+/-)- (UNII: D81QY6I88E)									
Packaging									
			Mark	eting Start	Mar	keting End			
# Item Code	Pa	ckage Description	Mark	Date	mar	Date			
1 NDC:10056-045-	1 in 1 CARTON	1	06/23/20	21					
• 00				2 I					
1	30 mL in 1 TU	BE; Type 0: Not a Combinat	tion						

F	roduct						
Marketing Information							
Marketing	Application Number or Monograph	Marketing Start	Marketing End				
Category	Citation	Date	Date				
OTC Monograph Drug	M006	06/23/2021					

Labeler - Access Business Group LLC (839830713)

EstablishmentNameAddressID/FEIBusiness OperationsAccess Business Group LLC839830713manufacture(10056-045)

Revised: 10/2023

Access Business Group LLC