

KILLA EXTRA STRENGTH 4 PACK- salicylic acid patch
ZitSticka, Inc.

Drug Facts	
Active Ingredients	Purpose
Salicylic Acid 2%	Acne Treatment
Uses:	
- For the management of acne, - Penetrates pores to eliminate most blemishes.	
Warnings: For external use only	
When using this product:	
Skin irritation and dryness may occur if you use another topical acne medication at the same time or immediately following. If irritation occurs, only use one topical acne medication at a time unless directed by a doctor.	
Stop use and ask a doctor if irritation becomes severe.	
Keep out of reach of children.	
If swallowed, get medical help or contact a Poison Control Center right away.	
Directions:	
- Cleanse skin thoroughly before applying the product. - Apply product directly to the affected area. - Leave on for 2 hours or overnight then remove. - If bothersome dryness or peeling occurs, reduce application to once a day or every other day.	
Other information:	
Store at room temperature. Avoid use if you have very sensitive skin or are allergic to Salicylic Acid.	
Sunburn Alert	
This product contains an Alpha Hydroxy Acid (AHA) that may increase your skin's sensitivity to the sun and particularly the possibility of sunburn. Use a sunscreen, wear protective clothing, and limit sun exposure while using this product and for a week afterwards.	
Inactive Ingredients:	
Sodium Hyaluronate, Phytosphingosine HCL, Niacinamide, Oligopeptide-10, Glycolic Acid, Madecassoside, 4-Butylresorcinol.	
Patch: Hydrocolloid	

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Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:81746-417
Route of Administration	TOPICAL		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)		SALICYLIC ACID	0.0125 mg in 0.6216 mg
Inactive Ingredients			
Ingredient Name			Strength
OLIGOPEPTIDE-10 (UNII: Q46328TRNK)			
GLYCOLIC ACID (UNII: 0WT12SX38S)			
4-BUTYLRESORCINOL (UNII: 2IK4UQ3Z GA)			
HYALURONATE SODIUM (UNII: YSE9PPT4TH)			
NIACINAMIDE (UNII: 25X51I8RD4)			
PHYTOSPHINGOSINE HYDROCHLORIDE (UNII: TT871XV7TU)			
MADECASSOSIDE (UNII: CQ2F5O6YIY)			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81746-417-01	3660 mg in 1 BOX; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)	01/01/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M006	01/01/2023	

Labeler - ZitSticka, Inc. (117778611)

Registrant - ZitSticka, Inc. (117778611)

Establishment

Name	Address	ID/FEI	Business Operations
Raphas Co., Ltd		695914964	manufacture(81746-417)

Revised: 9/2025

ZitSticka, Inc.