# **EVEN KEEL RESURFACING PEEL-** salicylic acid gel Personal Day Inc.

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#### **EVEN KEEL RESURFACING PEEL**

## **Drug Facts**

## **Active Ingredients**

Salicylic Acid 1.05%

### **Purpose**

Acne treatment

#### Uses

- Treats acne
- Helps prevent the development of new acne blemishes

## Warnings

## For external use only

# When using this product

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

# Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center immediately.

#### **Directions**

- After cleansing, cover the affected area with a thin layer one to two times per week.
- Because excessive drying of the skin may occur, start with one application (weekly), then gradually increase to two times weekly, if needed or as directed by a doctor.
- If bothersome dryness or peeling occurs, reduce application to every other week.

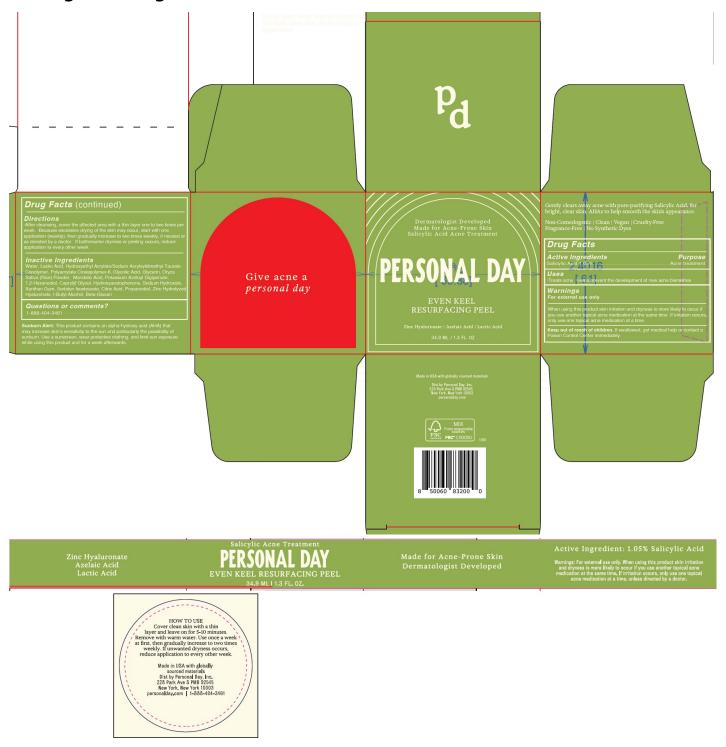
# Inactive Ingredients

Water, Lactic Acid, Hydroxyethyl Acrylate/Sodium Acryloyldimethyl Taurate Copolymer, Polyacrylate Crosspolymer-6, Glycolic Acid, Glycerin, Oryza Sativa (Rice) Powder, Mandelic Acid, Potassium Azeloyl Diglycinate, 1,2-Hexanediol, Caprylyl Glycol, Hydroxyacetophenone, Sodium Hydroxide, Xanthan Gum, Sorbitan Isostearate, Citric Acid, Propanediol, Zinc Hydrolyzed Hyaluronate, t-Butyl Alcohol, Beta-Glucan

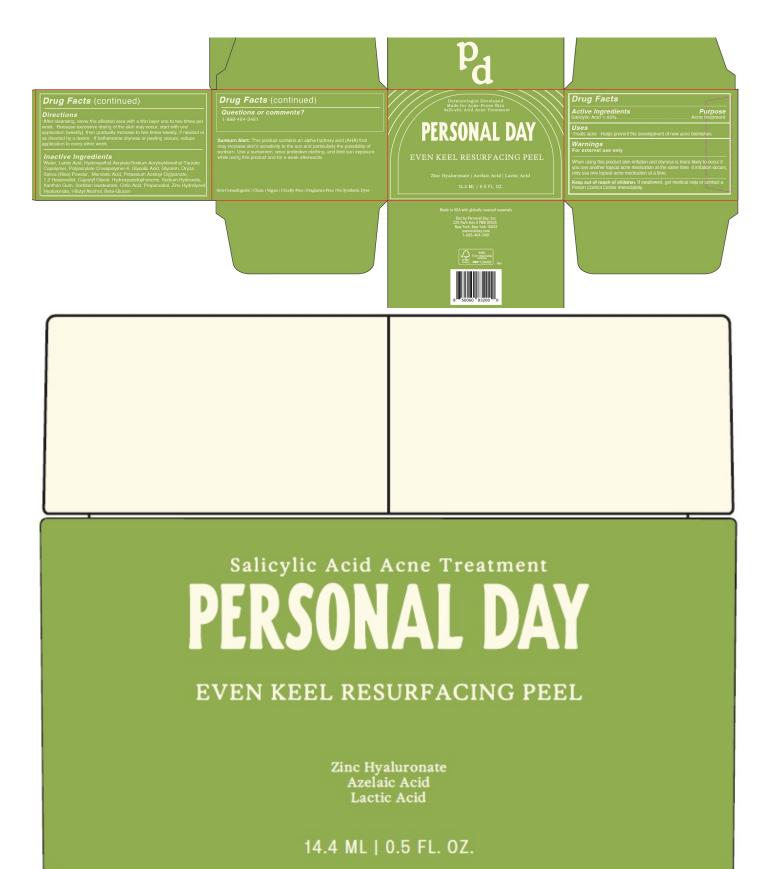
#### Questions or comments?

1-888-404-3461

## **Package Labeling:**



# Package Labeling:



# **EVEN KEEL RESURFACING PEEL** salicylic acid gel

#### **Product Information**

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:84234-000
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	10.5 mg in 1 mL	

Inactive Ingredients			
Ingredient Name	Strength		
WATER (UNII: 059QF0KO0R)			
LACTIC ACID (UNII: 33X04XA5AT)			
AMMONIUM ACRYLOYLDIMETHYLTAURATE, DIMETHYLACRYLAMIDE, LAURYL METHACRYLATE AND LAURETH-4 METHACRYLATE COPOLYMER, TRIMETHYLOLPROPANE TRIACRYLATE CROSSLINKED (45000 MPA.S) (UNII: Q7UI015FF9)			
GLYCOLIC ACID (UNII: 0WT12SX38S)			
GLYCERIN (UNII: PDC6A3C0OX)			
WHITE RICE (UNII: A195V20H7A)			
MANDELIC ACID (UNII: NH496X0UJX)			
POTASSIUM AZELOYL DIGLYCINATE (UNII: N02RVN6NYP)			
1,2-HEXANEDIOL (UNII: TR046Y3K1G)			
CAPRYLYL GLYCOL (UNII: 00YIU5438U)			
HYDROXYACETOPHENONE (UNII: G1L3HT4CMH)			
SODIUM HYDROXIDE (UNII: 55X04QC32I)			
XANTHAN GUM (UNII: TTV12P4NEE)			
SORBITAN ISOSTEARATE (UNII: 01S2G2C1E4)			
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)			
PROPANEDIOL (UNII: 5965N8W85T)			
TERT-BUTYL ALCOHOL (UNII: MD83SFE959)			

Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:84234-000- 00	1 in 1 CARTON	11/22/2024		
1		34.9 mL in 1 JAR; Type 0: Not a Combination Product			
2	NDC:84234-000- 01	1 in 1 CARTON	11/22/2024		
2		14.4 mL in 1 JAR; Type 0: Not a Combination Product			

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M006	11/22/2024	

# Labeler - Personal Day Inc. (119231300)

Revised: 12/2024 Personal Day Inc.