

**DERMAREST PSORIASIS MEDICATED SCALP TREATMENT- salicylic acid gel**  
**Insight Pharmaceuticals 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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**Dermarest Psoriasis Medicated Skin Treatment Gel**

***Drug Facts***

**Active ingredient**

Salicylic acid 3%

**Purposes**

Psoriasis and seborrheic dermatitis treatment

**Uses**

relieves and helps prevent recurrence of scalp:

- itching
- irritation
- redness
- flaking
- scaling

due to psoriasis and seborrheic dermatitis

**Warnings**

***For external use only***

***Ask a doctor before use if you have***

psoriasis that covers a large area of the body

***When using this product***

avoid contact with the eyes. If contact occurs rinse eyes thoroughly with water.

***Stop use and ask a doctor if***

condition worsens or does not improve after regular use as directed

***Keep out of reach of children.***

If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

**Directions**

apply to affected areas one to four times daily or as directed by a doctor

**Other information**

Store at room temperature 15°-25°C (59°-77°F).

**Inactive ingredients**

propylene glycol, purified water, SD alcohol 40, PPG-2 myristyl ether propionate, corn starch modified, zinc PCA, panthenol, aleurites moluccana seed oil, rheum palmatum extract, carthamus tinctorius (safflower) lower extract, camellia sinensis leaf extract, PPG-5-ceteth-20, edetate disodium, xanthan gum, triethanolamine, methylparaben, propylparaben, fragrance

**Questions?**

**1-800-344-7239 [dermarest.com](http://dermarest.com)**

**PRINCIPAL DISPLAY PANEL**

**DERMAREST®**

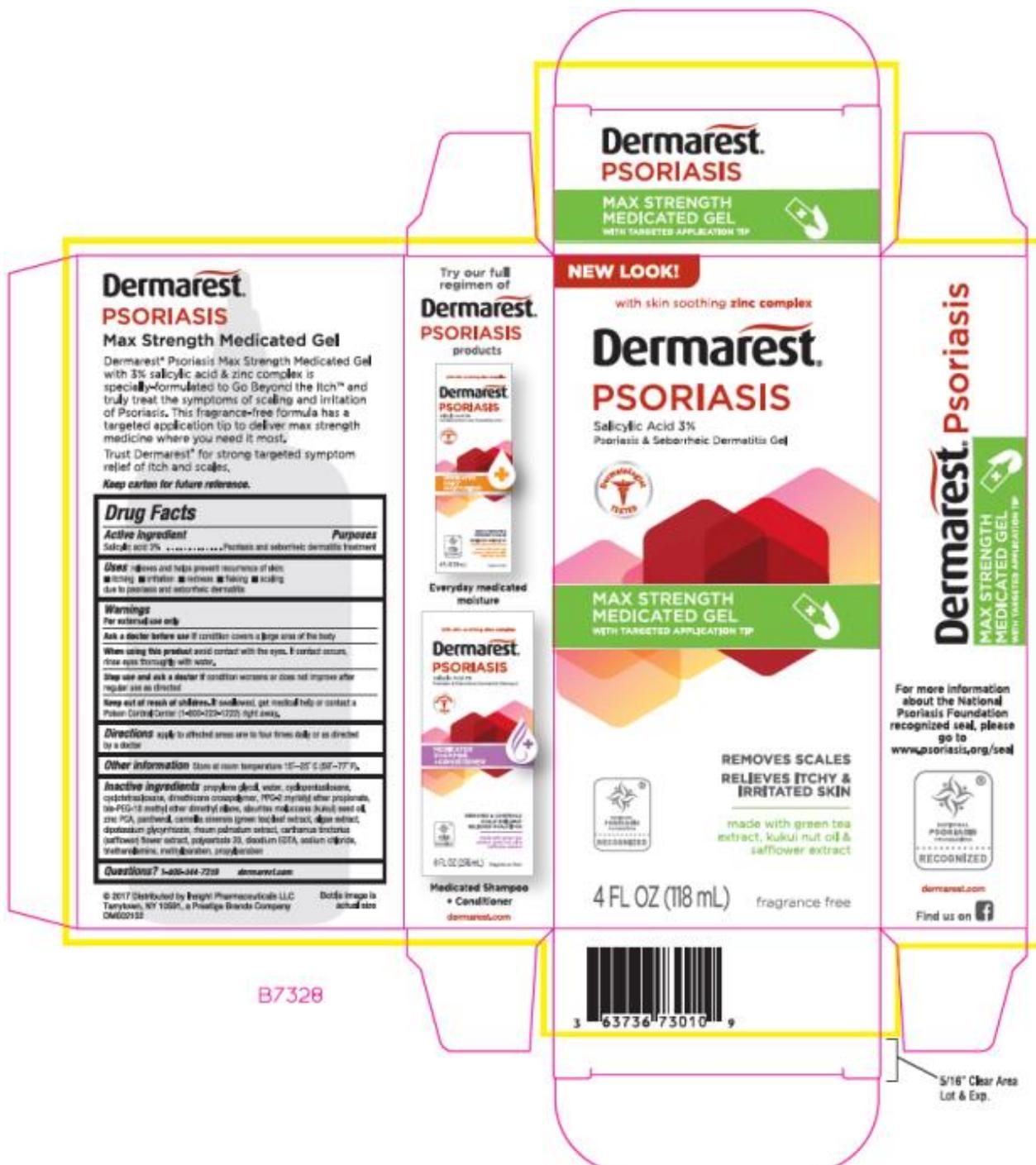
**Psoriasis**

**SALICYLIC ACID 3%**

Psoriasis & Seborrheic Dermatitis Gel

**MAX STRENGTH MEDICATED GEL**

4 FL OZ (118 mL)



## DERMAREST PSORIASIS MEDICATED SCALP TREATMENT

salicylic acid gel

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63736-312
Route of Administration	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	3 g in 100 mL

## Inactive Ingredients

Ingredient Name	Strength
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
CYCLOMETHICONE 5 (UNII: 0THT5PC10R)	
CYCLOMETHICONE 4 (UNII: CZ227117JE)	
DIMETHICONE CROSSPOLYMER (450000 MPAS AT 12% IN CYCLOPENTASILOXANE) (UNII: UF7620L1W6)	
PPG-2 MYRISTYL ETHER PROPIONATE (UNII: 88R97D8U8A)	
BIS-PEG-18 METHYL ETHER DIMETHYL SILANE (UNII: OEB4R3WW9C)	
KUKUI NUT OIL (UNII: TP11QR7B8R)	
ZINC PIDOLATE (UNII: C32PQ86DH4)	
PANTHENOL (UNII: WV9CM0O67Z)	
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
PHYMATOLITHON CALCAREUM (UNII: 6J1M3WA0ZK)	
GLYCYRRHIZINATE DIPO TASSIUM (UNII: CA2Y0FE3FX)	
RHEUM PALMATUM ROOT (UNII: G025DAL7CE)	
SAFFLOWER (UNII: 4VBL71TY4Y)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
EDETATE DISODIUM ANHYDRO US (UNII: 8NLQ36F6MM)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
TROLAMINE (UNII: 9O3K93S3TK)	
METHYL PARABEN (UNII: A2I8C7HI9T)	
PROPYL PARABEN (UNII: Z8IX2SC1OH)	

## Product Characteristics

Color	WHITE	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63736-312-24	24 in 1 CASE	12/20/2010	
1		1 in 1 CARTON		
1		118 mL in 1 BOTTLE; Type 0: Not a Combination Product		

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH FINAL	part358H	12/20/2010	

