

ACNE TREATMENT MEIJER- salicylic acid gel

MEIJER DISTRIBUTION, INC.

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.

MEIJER OIL FREE ACNE SPOT TREATMENT

Active ingredient

Salicylic acid 2.0%

Purpose

Acne Medication

Uses

- For the treatment of acne

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 right away. Avoid contact with eyes. If contact occurs, flush thoroughly with water.

Directions

- Cleanse skin thoroughly before applying product.
- Cover the entire affected area 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
- If bothersome dryness or peeling occurs, reduce application to once a day or every other day.

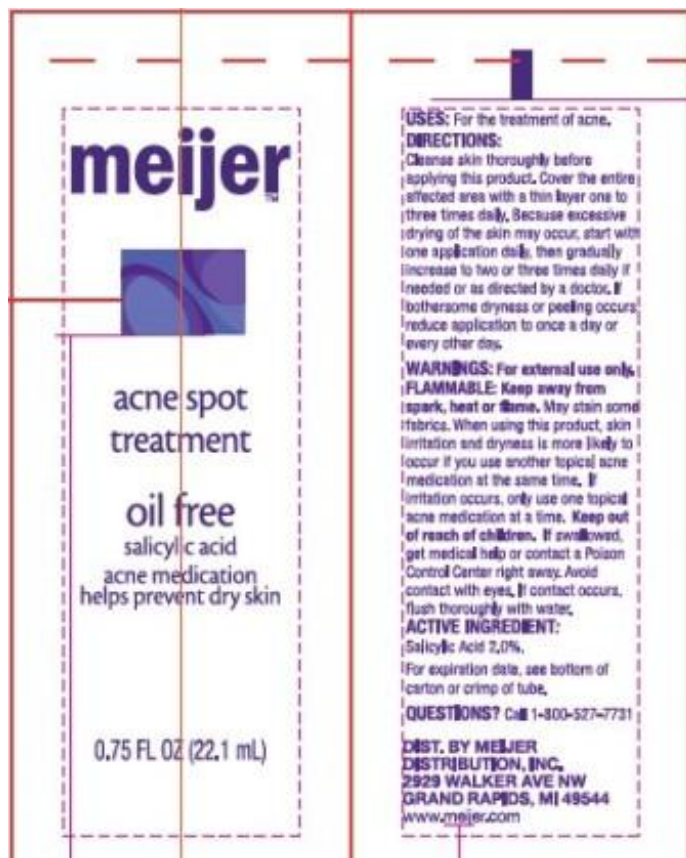
Other information

May stain some fabrics

Inactive ingredients

alcohol denat., ammonium hydroxide, benzalkonium chloride, butylene glycol, C12-15 alkyl lactate, C13-14 isoparaffin, capryloyl glycine, cedrus atlantica bark extract, cetyl lactate, cinnamomum zeylanicum bark extract, cocamidopropyl PG-dimonium chloride phosphate, cyclopentasiloxane, dehydroxanthan gum, ethylparaben, fragrance, glycerin, hamamelis virginiana (witch hazel) extract, hexylene glycol, laureth-7, methylparaben, phenethyl dimethicone, phenoxyethanol, polyacrylamide, polysorbate 20, portulaca oleracea extract, PPG-2 isocetheth-20 acetate, propylparaben, sarcosine, tetrasodium EDTA, water





ACNE TREATMENT MEIJER

salicylic acid gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:41250-560
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	20 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
C12-15 ALKYL LACTATE (UNII: GC844VRD7E)	
C13-14 ISOPARAFFIN (UNII: E4F12ROE70)	
CAPRYLOYL GLYCINE (UNII: 8TY5YO42NJ)	
CEDRUS ATLANTICA BARK (UNII: ITP1Q41UPF)	
COCAMIDOPROPYL PG-DIMONIUM CHLORIDE PHOSPHATE (UNII: H2KVQ74JM4)	
DEHYDROXANTHAN GUM (UNII: 63ZP7I1BQO)	
ETHYLPARABEN (UNII: 14255EXE39)	
DIMETHICONE (UNII: 92RU3N3Y1O)	

AMMONIA (UNII: 5138Q19F1X)	
CETYL LACTATE (UNII: A7EVH2RK4O)	
GLYCERIN (UNII: PDC6A3C0OX)	
HEXYLENE GLYCOL (UNII: KEH0A3F75J)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
SARCOSINE (UNII: Z711V88R5F)	
EDETATE SODIUM (UNII: MP1J8420LU)	
CYCLOMETHICONE 5 (UNII: 0THT5PCI0R)	
HAMAMELIS VIRGINIANA LEAF (UNII: T07U1161SV)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POLYACRYLAMIDE (1500 MW) (UNII: 5D6TC4BRWW)	
LAURETH-7 (UNII: Z95S6G8201)	
ALCOHOL (UNII: 3K9958V90M)	
CINNAMON BARK OIL (UNII: XE54U569EC)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PURSLANE (UNII: M6S840WVG5)	
PPG-2 ISOCETETH-20 ACETATE (UNII: BI6C7YO419)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41250-560-52	1 in 1 CARTON	09/23/2013	
1	NDC:41250-560-51	22.1 mL 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	part333D	09/23/2013	

Labeler - MEIJER DISTRIBUTION, INC. (006959555)

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