OXY OVERNIGHT ACNE REDUCING PATCHES- salicylic acid patch
The Mentholatum Company

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.

Drug Facts

Active ingredient

Salicylic acid 0.5%

Purpose

Salicylic acid - Acne treatment

Uses

treats and helps prevent acne

Warnings

For external use only

When using this product

- keep away from eyes, lips and mouth. If contact occurs, flush thoroughly with water.
- 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.

If pregnant or breast-feeding

ask a health professional before use.

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 this product
- apply patch(es) to affected area(s)
- to remove, gently peel back patch and discard
- because excessive drying of the skin may occur, start with 1 application daily, then gradually increase to 2 or 3 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

Inactive ingredients

acrylates/ethylhexyl acrylate copolymer, water, nonoxynol-30, alcohol denatured, butylene glycol, C13-14 isoparaffin, epilobium angustifolium flower/leaf/stem extract, kaolin, laureth-7, melaleuca alternifolia (tea tree) leaf oil, phenoxyethanol, phytosphingosine, polysorbate 80, PVP, sodium hyaluronate, sodium hydroxide, sodium metabisulfite, sodium polyacrylate, vitis vinifera (grape) seed extract, volcanic ash

Questions?

1-877-636-2677 MON-FRI 9 AM - 5 PM (EST)

Principal Display Panel

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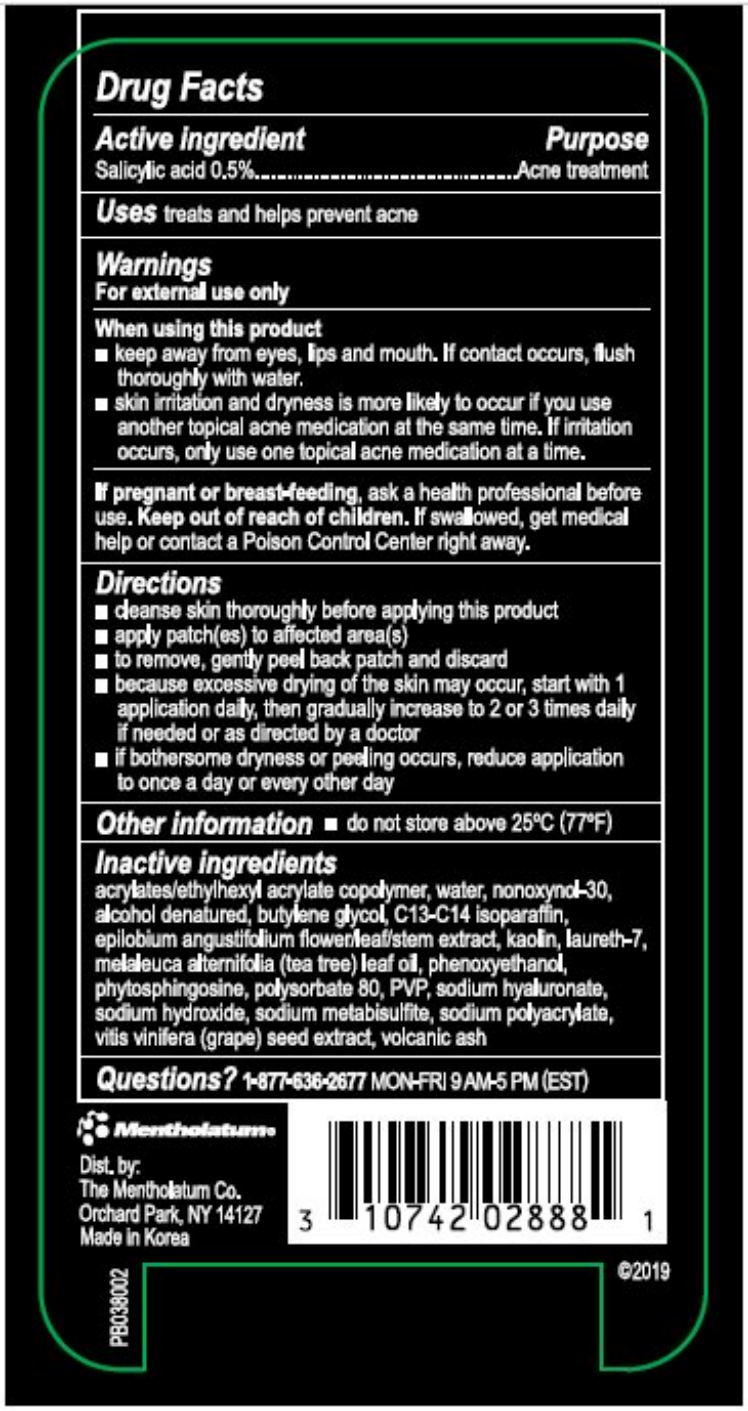

OXY OVERNIGHT ACNE REDUCING PATCHES

salicylic acid patch

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:10742-8177
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	5 mg

Inactive Ingredients

Ingredient Name	Strength
2-ETHYLHEXYL ACRYLATE (UNII: HR49R9S6XG)	
WATER (UNII: 059QF0KO0R)	
NONOXYNOL-30 (UNII: JJX07DG188)	
ALCOHOL (UNII: 3K9958V90M)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
C13-14 ISOPARAFFIN (UNII: E4F12ROE70)	
EPILOBIUM ANGUSTIFOLIUM WHOLE (UNII: C278QS9YBT)	
KAOLIN (UNII: 24H4NWX5CO)	
LAURETH-7 (UNII: Z95S6G8201)	
TEA TREE OIL (UNII: VIF565UC2G)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
PHYTOSPHINGOSINE (UNII: GIN46U9Q2Q)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
SODIUM METABISULFITE (UNII: 4VON5FNS3C)	
SODIUM POLYACRYLATE (8000 MW) (UNII: 285CYO341L)	
VITIS VINIFERA SEED (UNII: C34U15ICXA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10742-8177-1	30 in 1 PACKAGE; Type 0: Not a Combination Product	01/01/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333D	01/01/2018	

Labeler - The Mentholatum Company (002105757)

Registrant - The Mentholatum Company (002105757)

Establishment

Name	Address	ID/FEI	Business Operations

The Mentholatum Company		002105757	manufacture(10742-8177)