

OXY CLEANSING PADS DAILY DEFENSE- salicylic acid swab
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 2%

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.
- do not leave pad on skin

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

- use a pad to cleanse and cover the affected area with a thin layer of medicine 1 to 3 times daily
- because too much drying of the skin may occur, start with 1 application, 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

- *Sensitivity test:* Apply product sparingly to a small affected area for the first 3 days. If no discomfort occurs, follow direction above.

Other information

- KEEP TIGHTLY CLOSED
- avoid storing at temperatures above 100°F (38°C)
- protect from freezing
- keep away from flame, fire and heat

Inactive ingredients

alcohol (46% v/v), fragrance, isoceteth-20, PEG-8/SMDI copolymer, purified water, trisodium EDTA, trolamine

Questions?

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

Package/Label Principal Display Panel



Principal Display Panel

Drug Facts Active ingredient Purpose Salicylic acid 2% Acne treatment		Drug Facts (continued) with a thin layer of medicine 1 to 3 times daily ■ because too much drying of the skin may occur, start with 1 application, 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 ■ Sensitivity Test: Apply product sparingly to a small affected area for the first 3 days. If no discomfort occurs, follow directions above.	 Take the OXY® 28 Day Challenge for clear skin oxy28daychallenge.com  Download on the App Store  GET IT ON Google Play Clinically Proven Active Ingredient
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. ■ do not leave pad on skin 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 ■ use a pad to cleanse and cover the affected area			
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OXY CLEANSING PADS DAILY DEFENSE

salicylic acid swab

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:10742-3204
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SALICYLIC ACID (UNII: 0414PZ4LPZ) (SALICYLIC ACID - UNII:0414PZ4LPZ)	SALICYLIC ACID	20 mg

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
ISOCETETH-20 (UNII: 0020065R7Z)	
PEG-8/SMDI COPOLYMER (UNII: CCX72L6NY6)	
WATER (UNII: 059QF0KO0R)	
EDETATE TRISODIUM (UNII: 420IP921MB)	
TROLAMINE (UNII: 903K93S3TK)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10742-3204-1	90 in 1 JAR; Type 0: Not a Combination Product	08/01/2016	
2	NDC:10742-3204-2	55 in 1 JAR; Type 0: Not a Combination Product	08/01/2016	
3	NDC:10742-3204-3	115 in 1 JAR; Type 0: Not a Combination Product	08/01/2016	

4	NDC:10742-3204-4	70 in 1 JAR; Type 0: Not a Combination Product	08/01/2016
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Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333D	08/01/2016	

Labeler - The Mentholatum Company (002105757)

Registrant - The Mentholatum Company (002105757)

Establishment

Name	Address	ID/FEI	Business Operations
The Mentholatum Company		002105757	manufacture(10742-3204)

Revised: 2/2023

The Mentholatum Company