

SCANTILY PAD- dermala acne treatment liquid I Shay Cosmetics 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.

Dermala Acne Treatment

Use

For the treatment of Acne.

Active Ingredient

Salicylic acid 1%Acne Treatment

Warning

WARNINGS For external use only. Skin irritation and dryness is more likely to occur when using this product if you use another topical acne medication at the same time. If irritation occurs, only use one topical acne medication at a time.

Inactive ingredients

INACTIVE INGREDIENTS Organic Aloe vera, organic grain alcohol, cocamidopropyl dimethylamine, SE Microbiome Complex™, sodium hydroxide, organic lemon essential oil.

DIRECTIONS Cleanse the skin. Use one pad to apply a thin layer of solution to affected areas. Start with one application daily, then increase to two times per day (morning and evening). If dryness or peeling occurs, reduce application to once a day or every other day. Test on a small area of skin before applying to face. If eye contact occurs, flush thoroughly with water.

children

keep out of reach of children

indications and usage

For the treatment of Acne

Label

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Dermala Inc., 4040 Sorrento Valley Blvd., Suite C, San Diego, CA 92121

DERMALA®

SCANTILY Pad™

AM/PM Acne Treatment

With SE Microbiome Complex™

50 pads

Drug Facts

ACTIVE INGREDIENTS	Purpose
Salicylic acid 1%	Acne medication

USE For the treatment of acne.

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SCANTILY PAD

dermala acne treatment liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:65112-268
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	0.01 mg in 1 mg

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	0.8 mg in 1 mg

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65112-268-01	50 mg in 1 JAR; Type 0: Not a Combination Product	05/04/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333D	05/04/2021	

Labeler - I Shay Cosmetics Inc (151582384)

Revised: 12/2021

I Shay Cosmetics Inc