PERFECT PURITY MEDICATED BODY- menthol powder
Davion, 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.

Perfect Purity Medicated Body Powder

Active Ingredient

Menthol 0.15%

Purpose

External Analgesic

Uses

temporarily relieves pain and itch associated with:

- Minor Cuts
- Sunburn
- Insect Bites
- Scrapes
- Minor Burns
- Minor Skin Irritations

Warning

- For external use only.
- Avoid contact with eyes

Stop use and ask a doctor if

- condition worsens
- symptoms persists more than 7 days or clear up and occur again within a few days

Keep out of reach of children

In case of accidental ingestion, get medical help or contact a Poison Control Center right away.

Directions

- adults and children 2 years and older apply freely up to 3 or 4 times daily.
- children under 2 years ask a doctor.
- For best results, dry skin thoroughly before applying.

Inactive Ingredients

talc, zinc oxide, eucalyptol, methyl salicylate, salicylic acid, thymol, zinc stearate.

PRINCIPAL DISPLAY PANEL

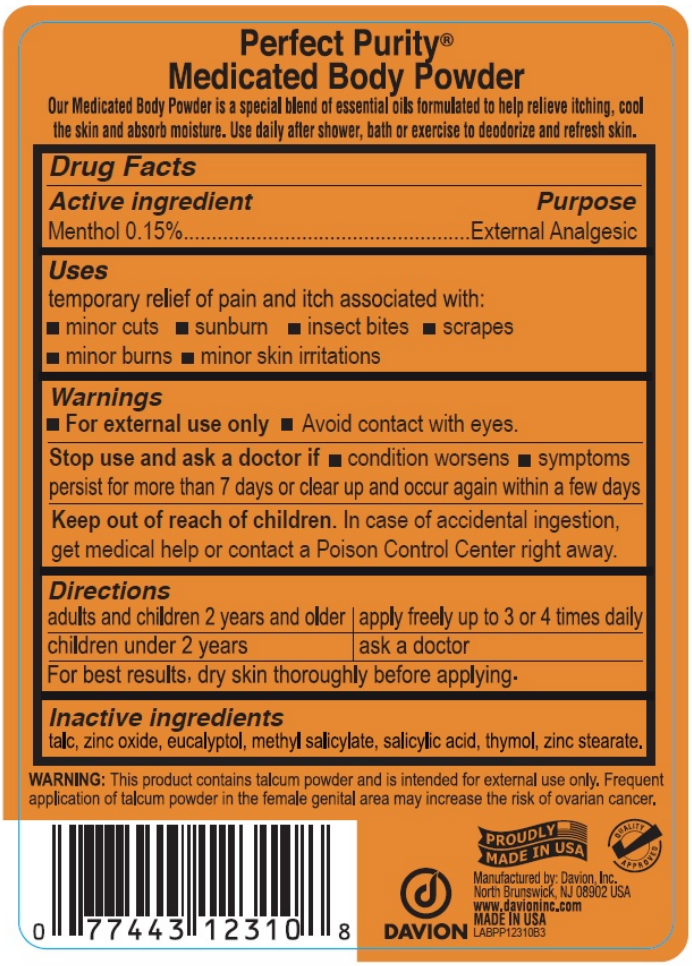
NDC 42669-208-01

Perfect Purity

medicated Body Powder

use daily to help skin feel fresh and healthy

NET WT 10OZ (283 g)



PERFECT PURITY MEDICATED BODY

menthol powder

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:42669-208
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	0.15 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
TALC (UNII: 7SEV7J4R1U)	

ZINC OXIDE (UNII: SOI2LOH54Z)	
EUCALYPTOL (UNII: RV6J6604TK)	
METHYL SALICYLATE (UNII: LAV5U5022Y)	
SALICYLIC ACID (UNII: O414PZ4LPZ)	
THYMOL (UNII: 3J50XA376E)	
ZINC STEARATE (UNII: H92E6QA4FV)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:42669-208-01	283 g in 1 CONTAINER; Type 0: Not a Combination Product	12/19/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part346	12/19/2017	

Labeler - Davion, Inc. (174542928)

Registrant - Davion, Inc. (174542928)

Establishment

Name	Address	ID/FEI	Business Operations
Davion, Inc.		174542928	manufacture(42669-208)

Revised: 12/2017 Davion, Inc.