BODY- menthol powder
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 379.001/379AB

Active ingredient

Menthol 0.15%

Purpose

External analgesic

Uses

for temporary relief of pan and itching due to:

- minor burns
- sunburn
- minor cuts
- scrapes
- insect bites
- minor skin irritations
- rashes due to poison ivy, poison oak, poison sumac

warnings

For external use only

When using this product

avoid contact with the eyes

Stop use and ask a doctor

If condition worsens, or if symptoms persist for more than 7 days or clear up and occur again within a few days

keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

adults and children 2 year of age and older - apply to affected area not more than 3 to 4 times daily

children under 2 years of age - do not use, consult a doctor

- for best results dry skin thoroughly before applying

Inactive ingredients

Zea mays (corn) starch. sodium bicarbonate, tricalcium phosphate, zinc oxide, acacia seyal gum, eucalyptol, methyl salicylate, salicylic acid, thymol, zinc stearate

disclaimers

This product is sold by weight, not by volume. Some settling may occur during handling and shipping

Adverse reactions

DISTRIBUTED BY

MEIJER DISTRIBUTION, INC

GRAND RAPIDS, MI 49544

www.meijer.com

principal display panel

Meijer

MEDICATED

BODY POWDER

External Analgesic

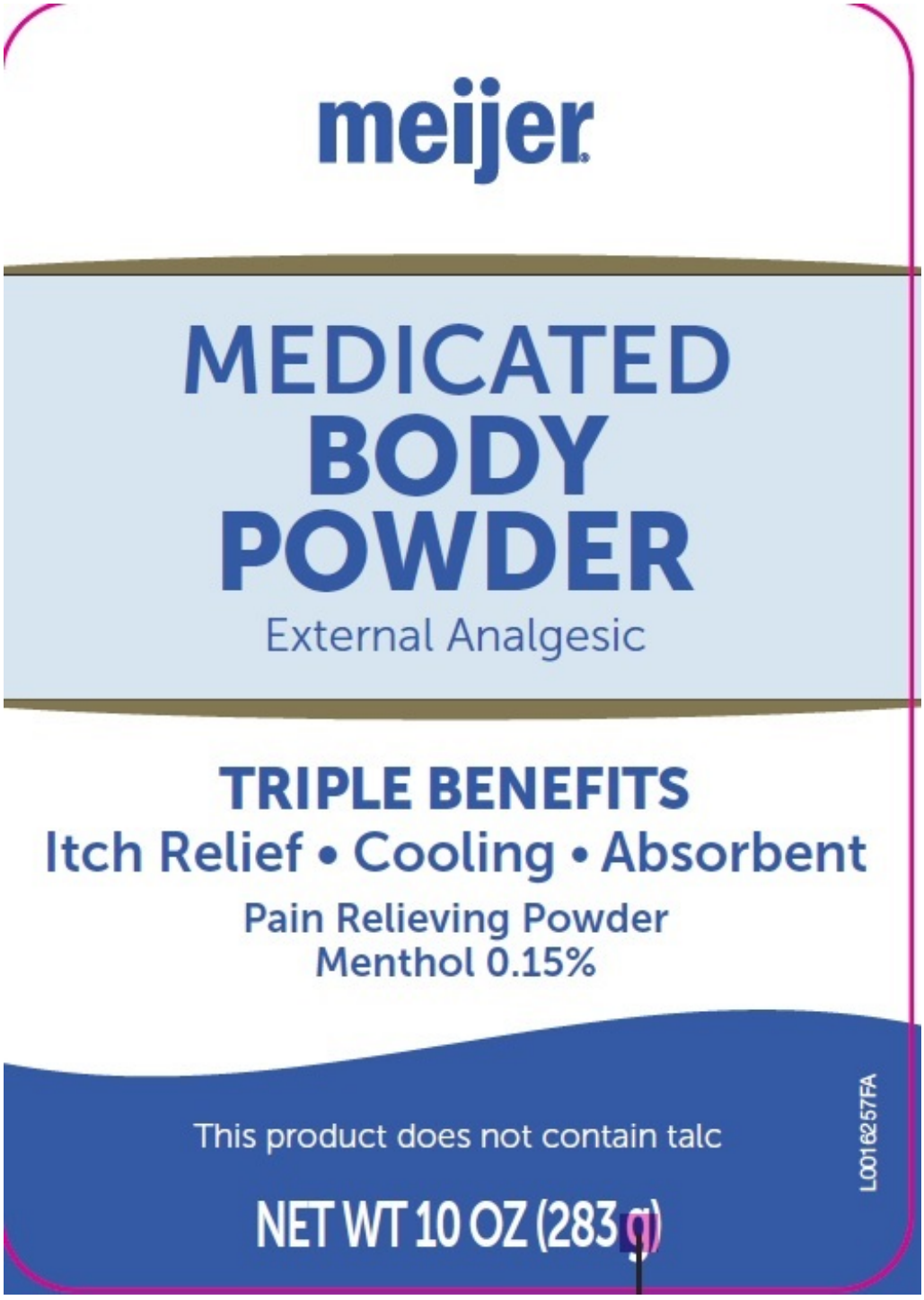
TRIPLE BENEFITS
- Itch Relief
- Cooling
- Absorbent

Pain Relieving Powder

Menthol 0.15%

This product does not contain talc

NET WT 10 OZ (283 g)



BODY

menthol powder

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:41250-043
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**

MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	1.5 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
STARCH, CORN (UNII: O8232NY3SJ)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
TRICALCIUM PHOSPHATE (UNII: K4C08XP666)	
ZINC OXIDE (UNII: SOI2LOH54Z)	
GUM TALHA (UNII: H18F76G097)	
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:41250-043-38	283 g in 1 BOTTLE; Type 0: Not a Combination Product	11/15/1987	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	11/15/1987	

Labeler - Meijer Distribution, Inc (006959555)

Registrant - Vi-Jon, LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
MK Packaging		790752542	manufacture(41250-043)

Revised: 4/2022 Meijer Distribution, Inc