BENGAY ULTRA STRENGTH LARGE SIZE- menthol, unspecified form patch
Johnson & Johnson Consumer 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.

BENGAY® ULTRA STRENGTH PAIN RELIEVING PATCH
Large Size

Drug Facts

Active ingredient

Menthol 5%

Purpose

Topical analgesic

Uses

temporarily relieves minor aches and pains of muscles and joints associated with:
- simple backache
- arthritis
- strains
- bruises
- sprains

Warnings

For external use only.

Do not use
- on wounds or damaged skin
- with a heating pad
- on a child under 12 years of age with arthritis-like conditions

Ask a doctor before use if you have redness over the affected area

When using this product
- avoid contact with eyes or mucous membranes
- do not bandage tightly

Stop use and ask a doctor if
- condition worsens or symptoms persist for more than 7 days
- symptoms clear up and occur again within a few days
- excessive skin irritation occurs

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- open pouch and remove patch
- if desired, cut patch to size
- peel off protective backing and apply sticky side to affected area
- adults and children 12 years of age and older: apply to affected area not more than 3 to 4 times daily
- children under 12 years of age: ask a doctor

Other information

- store at 20° to 25°C (68° to 77°F)

Inactive ingredients

calcined kaolin, cellulose gum, glycerin, methyl acrylate/2-ethylhexyl acrylate copolymer, methylparaben, polyacrylic acid, polysorbate 80, propylparaben, silica, sodium polyacrylate, sodium polyacrylate starch, sorbitan oleate, sorbitol, tartaric acid, titanium dioxide, water

Questions?

call **1-800-223-0182** (toll-free) or **215-273-8755** (collect)

Distributed by: **JOHNSON & JOHNSON COMSUMER INC.**
Skillman, NJ 08558

PRINCIPAL DISPLAY PANEL - 1 Patch Pouch Carton

***BENGAY*®**
MENTHOL 5% TOPICAL
ANALGESIC PATCH

ULTRA STRENGTH
PAIN RELIEVING
PATCH
Targeted relief that's
designed to **stay in place**

4
LARGE
FOR BACK TO HIP

4 INDIVIDUALLY SEALED PATCHES
3.9 IN x 7.9 IN (10 cm x 20 cm)

34.394 mm

15.345 mm

LOT:

EXP:

BENGAY®

ULTRA STRENGTH PAIN RELIEVING PATCH

34.459 mm

BENGAY®

MENTHOL 5% TOPICAL ANALGESIC PATCH

ULTRA STRENGTH
PAIN RELIEVING
PATCH

Targeted relief that's designed to **stay in place**

4

LARGE
FOR BACK TO HIP

4 INDIVIDUALLY SEALED PATCHES
3.9 IN x 7.9 IN (10 cm x 20 cm)

36.130 mm

0 74300 08149 6

18.874 mm

6.000 mm

6.000 mm

BENGAY®
ULTRA STRENGTH

Provides deep, penetrating relief from muscle aches & joint pain

Feels cool, then warm

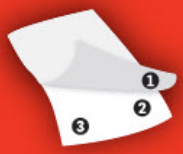
Convenient, no mess

34.459 mm

BENGAY® ULTRA STRENGTH PAIN RELIEVING PATCH

For best results, please ensure skin is clean and dry before applying.

- Apply one Ultra Strength BENGAY® Pain Relieving Patch to the site of pain to deliver the relief you need.
- Large size is best for lower back, upper back, shoulders and hips.



1. Easy to remove backing
2. Soft breathable pad
3. Pain relieving hydrogel layer

Also available in Regular size

Drug Facts

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Menthol 5%	Topical analgesic

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ACTUAL SIZE

4 LARGE

18.034 mm

BENGAY® ULTRA STRENGTH PAIN RELIEVING PATCH

34.460 mm

BENGAY ULTRA STRENGTH LARGE SIZE

menthol, unspecified form patch

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69968-0487
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Menthol, Unspecified Form (UNII: L7T10EIP3A) (Menthol, Unspecified Form - UNII:L7T10EIP3A)	Menthol, Unspecified Form	50 mg

Inactive Ingredients	
Ingredient Name	**Strength**
Kaolin (UNII: 24H4NWX5CO)	
Carboxymethylcellulose Sodium, Unspecified Form (UNII: K679OBS311)	
Glycerin (UNII: PDC6A3C0OX)	
Methylparaben (UNII: A2I8C7HI9T)	
Polyacrylic Acid (8000 MW) (UNII: 73861X4K5F)	
Polysorbate 80 (UNII: 6OZP39ZG8H)	
Propylparaben (UNII: Z8IX2SC1OH)	
Silicon Dioxide (UNII: ETJ7Z6XBU4)	
Sodium Polyacrylate (8000 MW) (UNII: 285CYO341L)	
Sorbitan Monooleate (UNII: 06XEA2VD56)	
Sorbitol (UNII: 506T60A25R)	
Tartaric Acid (UNII: W4888I119H)	
Titanium Dioxide (UNII: 15FIX9V2JP)	
Water (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:69968-0487-1	1 in 1 CARTON	04/03/2019	
1		1 in 1 POUCH; Type 0: Not a Combination Product		
2	NDC:69968-0487-4	4 in 1 CARTON	09/01/2019	
2		1 in 1 POUCH; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part348	04/03/2019	

Labeler - Johnson & Johnson Consumer Inc. (002347102)