BENGAY VANISHING SCENT PAIN RELIEVING- menthol, unspecified form gel
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® VANISHING SCENT MENTHOL PAIN RELIEVING GEL

Drug Facts

Active ingredient

Menthol 2.5%

Purpose

Topical analgesic

Uses

temporarily relieves the 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
- 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

water, isopropyl alcohol, PEG-40 hydrogenated castor oil, carbomer, isoceteth-20, sodium hydroxide, DMDM hydantoin, camphor

Questions?

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

Distributed by:
JOHNSON & JOHNSON CONSUMER INC.
Skillman, NJ 08558

PRINCIPAL DISPLAY PANEL - 57 g Tube Carton

VANISHING SCENT

NON-GREASY GEL

BENGAY®
MENTHOL PAIN RELIEVING GEL

NET WT 2 OZ (57 g)

BENGAY VANISHING SCENT PAIN RELIEVING

menthol, unspecified form gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69968-0240
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	25 mg in 1 g

Inactive Ingredients	
Ingredient Name	**Strength**
Water (UNII: 059QF0KO0R)	
Isopropyl Alcohol (UNII: ND2M416302)	
Polyoxyl 40 Hydrogenated Castor Oil (UNII: 7YC686GQ8F)	
Carboxypolymethylene (UNII: 0A5MM307FC)	
Isoceteth-20 (UNII: O020065R7Z)	
Sodium Hydroxide (UNII: 55X04QC32I)	
DMDM Hydantoin (UNII: BYR0546TOW)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:69968-0240-2	1 in 1 CARTON	11/30/2009	
1		57 g in 1 TUBE; 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	11/30/2009	

Labeler - Johnson & Johnson Consumer Inc. (002347102)

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