EEZ-AWAY RELIEF- menthol spray
EEZAWAY RELIEF 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.

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eez-away RELIEF

ACTIVE INGREDIENT:

Menthol 1.25%

Liquid Analgesic

Keep out of reach of children.

FOR THE TEMPORARY RELIEF OF MINOR ACHES & PAINS ASSOCIATED WITH ARTHRITIS, SIMPLE BACKACHE & SORE, TIRED MUSCLES.

CAUTION: Do not apply to wounds or damaged skin. If rash or irritation occurs, discontinue use.

The application of external heat, such as an electric heating pad, may result in excessive skin irritation or burn. Avoid contact with the eyes and mucous membranes. Do not bandage tightly. Do not inhale.

DIRECTIONS FOR USE

Apply EEZ-AWAY® generously to the affected area. Let dry, then re-apply. For maximum pain relief, apply, let dry and re-apply five times daily or as needed.

INGREDIENTS: Isopropyl Alcohol, Deionized Water (Aqua), PEG-75 Lanolin, Iodine, Oleth-10, PPG-20 Methyl Glucose Ether Distearate, Sodium Iodide, Sodium Thiosulfate, Ethyl Alcohol, Fragrance.

To reorder call: 302-339-3030

Packaging

EEZ-AWAY RELIEF

menthol spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69678-101
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	1.25 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	
IODINE (UNII: 9679TC07X4)	

ISOPROPYL ALCOHOL (UNII: ND2M416302)	
POLYOXYL-10 OLEYL ETHER (UNII: JD797EF70J)	
PEG-75 LANOLIN (UNII: 09179OX7TB)	
PPG-20 METHYL GLUCOSE ETHER DISTEARATE (UNII: 0057334FAB)	
SODIUM IODIDE (UNII: F5WR8N145C)	
SODIUM THIOSULFATE (UNII: HX1032V43M)	

Product Characteristics

Color	brown (amber)	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69678-101-08	236.6 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	03/19/2015	
2	NDC:69678-101-06	177.4 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	05/31/2017	
3	NDC:69678-101-04	118 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	05/31/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	03/19/2015	

Labeler - EEZAWAY RELIEF INC (079751465)

Establishment

Name	Address	ID/FEI	Business Operations
Topiderm Inc.		049121643	manufacture(69678-101)

Revised: 6/2017 EEZAWAY RELIEF INC