EQUALINE BLUE ICE PAIN RELIEVING - menthol gel
SUPERVALU 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.

Drug Facts

Active Ingredient Purpose

Menthol 2.0% .....Topical Analgesic

Uses: Temporary relief of minor aches and pains in: Muscles and joints.

Warnings:

For external use only. Avoid contact with eyes and mucus membranes.

When using this product do not:

- Use with heating pads or heating devices
- Use, pour, spill, or store near open flame
- Use with other creams, sprays or liniments
- Apply to damaged skin or wounds
- Bandage area tightly

To do so may result in excessive skin irritation or skin burn.

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

and consult a physician. If you have sensitive skin, consult a physician. If skin irritation develops, discontinue use and seek the advice of a physician before

using this product.

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

Directions:

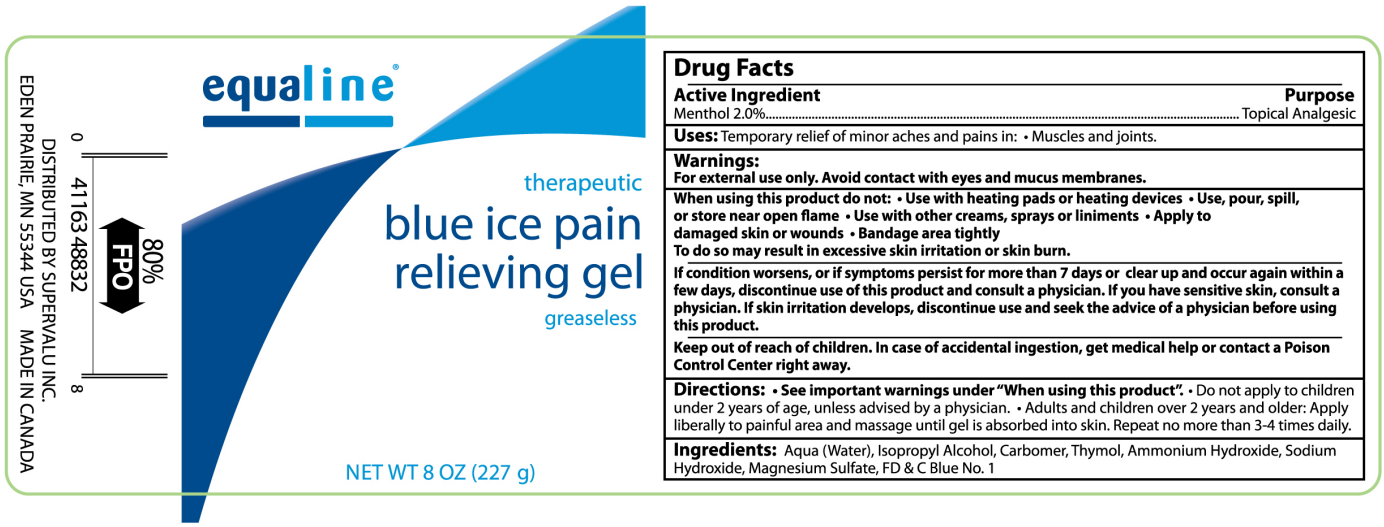
- See important warnings under "When using this product."
- Do not apply to children under 2 years of age, unless advised by a physician
- Adults and children over 2 years and older: Apply liberally to painful area and massage until gel is absorbed into skin. Repeat no more than 3-4 times daily

Inactive Ingredients: Aqua (water), Isopropyl Alcohol, Carbomer, Thymol, Ammonium Hydroxide, Sodium Hydroxide, Magnesium Sulfate, FD and C Blue No.1

DISTRIBUTED BY SUPERVALU INC.

EDEN PRAIRIE, MN 55344 USA

MADE IN CANADA



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EQUALINE BLUE ICE PAIN RELIEVING

menthol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:41163-488
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	2 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
THYMOL (UNII: 3J50XA376E)	
AMMONIA (UNII: 5138Q19F1X)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
MAGNESIUM SULFATE, UNSPECIFIED (UNII: DE08037SAB)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41163-488-32	227 g in 1 CONTAINER		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part348	10/03/2011	

Labeler - SUPERVALU INC (006961411)

Revised: 10/2011 SUPERVALU INC