BLUE ICE ANALGESIC- menthol gel
Delon Laboratories (1990) Ltd

Blue Ice Gel

Active ingredient

Menthol 2.0%

Purpose

Topical analgesic

Uses

- for the temporary relief of minor aches and pains of muscles and joints

Warnings

For external use only

When using this product
- avoid contact with the eyes
- do not apply to wounds or damaged skin
- do not bandage tightly

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

If pregnant or breast-feeding, ask a health professional before use.

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

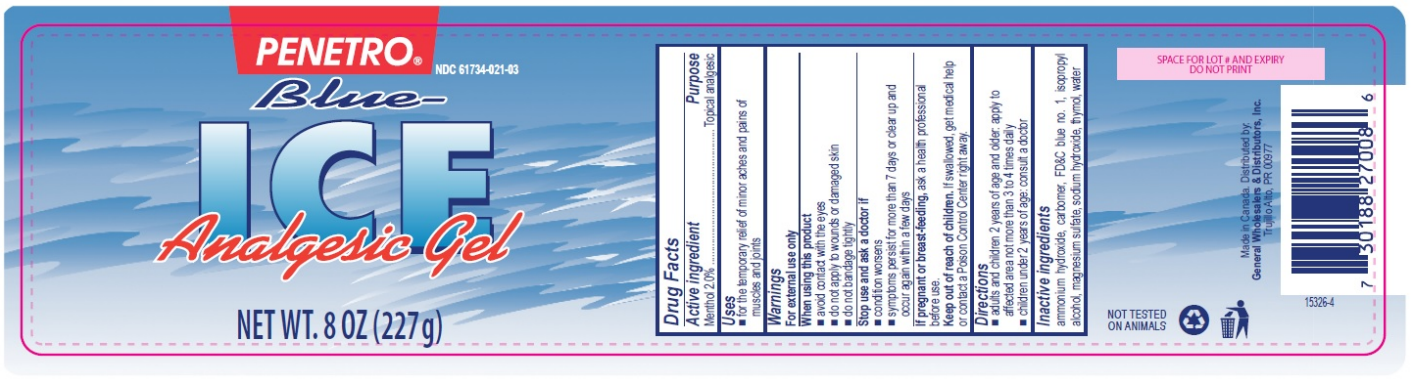
Directions
- adults and children 2 years of age and older: apply to affected area not more than 3 to 4 times daily
- children under 2 years of age: consult a doctor

Inactive ingredients

ammonium hydroxide, carbomer, FD&C blue no. 1, isopropyl alcohol, magnesium sulfate, sodium hydroxide, thymol, water

Delon 8oz (227g)

Penetro 227g



BLUE ICE ANALGESIC

menthol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:61734-021
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**
THYMOL (UNII: 3J50XA376E)	
AMMONIA (UNII: 5138Q19F1X)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
MAGNESIUM SULFATE HEPTAHYDRATE (UNII: SK47B8698T)	
CARBOMER 934 (UNII: Z135WT9208)	

FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
WATER (UNII: 059QF0KO0R)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	

Product Characteristics

Color	blue	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:61734-021-01	100 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/07/2010	11/06/2017
2	NDC:61734-021-02	113.56 g in 1 CONTAINER; Type 0: Not a Combination Product	05/07/2010	12/04/2014
3	NDC:61734-021-03	227 g in 1 CONTAINER; Type 0: Not a Combination Product	05/07/2010	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	05/07/2010	

Labeler - Delon Laboratories (1990) Ltd (248364184)

Establishment

Name	Address	ID/FEI	Business Operations
Laboratoires Delon		208896216	manufacture(61734-021) , pack(61734-021) , label(61734-021)

Revised: 10/2025 Delon Laboratories (1990) Ltd