

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)



DELON. TEMPORARY RELIEF OF MINOR ACHEs & PAINs OF MUSCLES & JOINTS
NDC 61734-021-03

Blue-ICE
Analgesic Gel

NET WT. 8 OZ

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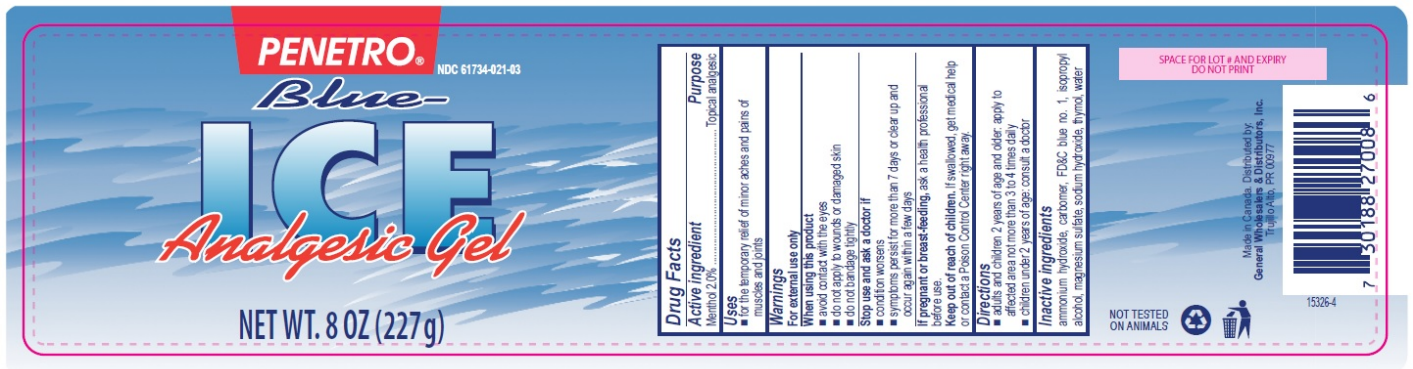
Made in Canada by:
LABORATOIRES DELON (1980) INC.
Pointe-Claire, Quebec, Canada, H8R 1E2
www.labdelon.com @labdelon

14285-4
NOT TESTED ON ANIMALS

SPACE FOR LOT # AND EXPIRY DO NOT PRINT

0 59338 12825 2

Penetro 227g



PENETRO. NDC 61734-021-03

Blue-ICE
Analgesic Gel

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Made in Canada. Distributed by:
General Wholesalers & Distributors, Inc.
Toronto, Ontario, Canada, M5S 1A7
Tel: 416-491-1111 PR 05977

NOT TESTED ON ANIMALS

SPACE FOR LOT # AND EXPIRY DO NOT PRINT

7 30188 27008 6

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: 3/2025

Delon Laboratories (1990) Ltd