LUCKY BLUE ICE- menthol camphor gel
Anicare Pharmaceuticals Pvt. Ltd.

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.

Ice Gel

Active ingredients

Menthol 1.0%
Camphor 0.5%

Purpose

Topical Analgesic

Use

Lucky Super Soft Blue Ice Gel to provide temporary relief of minor pains and aches in your body's joints and muscles associated with:

- sports injuries
- sprains
- arthritis
- bruises

Warnings

for external use only
Do not apply to wound or damaged skin. Do not bandage the applied surface.
If condition worsens, discontinue use of this product.
Consult a physician if symptoms persist for more than 7 days.

Avoid contact with mucus membranes and eyes.

Do not use with heating devices or pads.

Keep out of reach of children

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

Directions

Apply liberally to painful zones and massage until absorbed into the skin. Repeat daily 3 or 4 times.
Keep out of reach of children. Do not apply to children below two years of age.

Inactive Ingredients

Deionized Water, Isopropyl Alcohol, Propylene Glycol,Carbopol, PEG-40 Hydrogenated Castor Oil, Sodium Hydroxide,
Disodium EDTA, Benzyl Alcohol, BHT, FDC Blue No. 1

Package Label

Drug Facts

Active ingredients:	Purpose
Menthol 1.0 %	Topical Analgesic
Camphor 0.5 %	Topical Analgesic

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Hydrogenated Castor Oil, Sodium Hydroxide, Disodium EDTA, Benzyl Alcohol, BHT, FD&C Blue No.1

LUCKY BLUE ICE

menthol camphor gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:47046-167
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	1 g in 100 g
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET) (CAMPHOR (SYNTHETIC) - UNII:5TJD82A1ET)	CAMPHOR (SYNTHETIC)	0.5 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
ALCOHOL (UNII: 3K9958V90M)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
CARBOMER 934 (UNII: Z135WT9208)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
HYDROGENATED CASTOR OIL (UNII: ZF94AP8MEY)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
POLYETHYLENE GLYCOL 200 (UNII: R95B8J264J)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:47046-167-01	227 g in 1 JAR; Type 0: Not a Combination Product	11/15/2009	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	11/15/2009	

Labeler - Anicare Pharmaceuticals Pvt. Ltd. (916837425)