GOOD NEIGHBOR PHARMACY THERAPEUTIC BLUE- menthol gel AmerisourceBergen Drug Corporation

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.

Good Neighbor Pharmacy Therapeutic Blue

Drug Facts

Active Ingredients

Active ingredient

Menthol 2%

pain relieving gel

Uses

for the temporary relief of minor aches and pains of muscles and joints associated with

- simple backache arthritis strains
- bruises sprains

warnings

For external use only

when using this product

- avoid contact with the eyes
- do not bandage tightly
- do not apply to wounds or damaged skin
- do not use with heating pads or other heating devices

Stop using and ask a doctor if

• condition worsens, or if symptoms persist for more than 7 days or clear up and occur again within a few days

Keep out of the reach of children

If swallowed get medical help or contact a Poison Control Center.

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: do not use, consult a doctor

Inactive Ingredients

Ammonium Hydroxide, Carbomer, Cupric Sulfate, FD&C Blue No. 1, Isopropyl Alcohol, Magnesium Sulfate, Sodium Hydroxide, Thymol, Water.

This product is not manufactured by Novartis Consumer Health, Inc., owner of the registered trademark Mineral Ice®.



GOOD NEIGHBOR PHARMACY THERAPEUTIC BLUE

menthol gel

Product Information				
l	Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:24385-225

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		
AMMO NIA (UNII: 5138 Q 19 F1X)			
CARBO MER 934 (UNII: Z135WT9208)			
CUPRIC SULFATE (UNII: LRX7AJ16 DT)			
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)			
ISOPROPYL ALCOHOL (UNII: ND2M416302)			
MAGNESIUM SULFATE (UNII: DE08037SAB)			
SO DIUM HYDRO XIDE (UNII: 55X0 4QC32I)			
THYMOL (UNII: 3J50 XA376E)			
WATER (UNII: 059QF0KO0R)			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:24385-225-40	227 g in 1 JAR; Type 0: Not a Combination Product		
Н				

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part348	03/11/1988		

Labeler - AmerisourceBergen Drug Corporation (007914906)

Establishment				
Name	Address	ID/FEI	Business Operations	
Product Quest Mfg, LLC		927768135	manufacture(24385-225), label(24385-225)	

Revised: 7/2015 AmerisourceBergen Drug Corporation