EB302 ARTHRITIS RELIEF- menthol spray EMINENCE BIOTECH CORPORATION LIMITED

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.

EB302 ARTHRITIS RELIEF

Drug Facts

Active ingredient

Menthol 1.7307%

Purpose

Topical Analgesic

Uses

For the temporary relief of minor aches & pains of muscles & joints associated with arthritis

Warnings

For external use only

Do not use

- on ears or nasal cavity
- if pregnant
- on wounds or damaged skin
- otherwise than as directed

When using this product avoid contact with the eyes. If contact occurs, rinse eyes thoroughly with water.

Stop use and ask a doctor if

- Conditions worsen
- Symptoms persist for more than 7 days, or symptoms clear up and occur again within a few days

Keep out of reach of children.

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

Directions

Adults and children 12 years of age and over

Spray sufficient amount onto affected area to cover it completely, and rub until dry. Repeat no more than 3 times daily.

Children under 12 years of age

Consult a doctor.

Other information

Store at cool place (room temperature) away from heat and light sources.

Inactive ingredients

Alcohol, Amaranthus Viridis Leaf, Animal Fat, Gaultheria Procumbens (Wintergreen) Leaf Extract, Glycerin, Mesona Chinensis Whole, Water

Questions?

Call 1-562-207-8624

Distributed by:

EMINENCE BIOTECH CORPORATION LIMITED

3411 SILVERSIDE ROAD, RODNEY BUILDING # 104 WILMINGTON DE19810 NEW CASTLE COUNTY, USA

PRINCIPAL DISPLAY PANEL - 50mL Bottle Carton

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Eminence

Biotech

EB302

ARTHRITIS RELIEF SPRAY

1.69FL OZ (50mL)

NDC 53304-03021



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Made in Taiwan



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ARTHRITIS RELIEF SPRAY



Exp: 2 years from manufacture date.

Use within 45 days of opening.

Manufacture date: month/date/year Please recycle

4 7 1 10 2 1 4 2 3 2 16



EB302 ARTHRITIS RELIEF

menthol spray

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:53304-0302	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A) (MENTHOL, UNSPECIFIED FORM - UNII:L7T10EIP3A)	MENTHOL, UNSPECIFIED FORM	865.4 mg in 50 mL		

Inactive Ingredients		
Ingredient Name	Strength	
ALCOHOL (UNII: 3K9958V90M)		
AMARANTHUS VIRIDIS LEAF (UNII: 7T0 WZ76 K1T)		
PORK SIDE BACON, CURED FAT (UNII: 6E710H379S)		
GAULTHERIA PRO CUMBENS LEAF (UNII: 2125M16 OWN)		
GLYCERIN (UNII: PDC6A3C0OX)		
PLATOSTOMA PALUSTRE WHOLE (UNII: FE70 D0 62O9)		
WATER (UNII: 059QF0KO0R)		

]	Packaging				
7	t Item Code	Package Description	Marketing Start Date	Marketing End Date	
:	NDC:53304-0302-	1 in 1 BOX	11/0 1/20 12		
:		50 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product			

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC MONOGRAPH NOT FINAL	part348	11/0 1/20 12			

Labeler - EMINENCE BIOTECH CORPORATION LIMITED (078531141)

Revised: 3/2018 EMINENCE BIOTECH CORPORATION LIMITED