

EB301AP PAIN 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.

EB301AP PAIN RELIEF

Drug Facts

Active ingredient

Menthol 1%

Purpose

Topical Analgesic

Uses

For the temporary relief of pain

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 to 4 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, Animal Fat, Euphorbia Hirta, Euphorbia Thymifolia Whole, Gaultheria Procumbens (Wintergreen) leaf Extract, Glycerin, 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 - 100 mL Bottle Carton

**EB
Eminence
Biotech**

EB301AP

PAIN RELIEVING SPRAY

3.38FL OZ (100mL)

NDC 53304-0301-1



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

Made in Taiwan




EB301AP

PAIN RELIEVING SPRAY



Exp: 2 years from manufacture date.
Use within 45 days of opening.

Manufacture date: month/date/year
Please recycle



3.38FL OZ (100mL)
NDC 53304-0301-1

EB301AP PAIN RELIEF

menthol spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:53304-0301
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	1000 mg in 100 mL	
Inactive Ingredients				
Ingredient Name			Strength	
ALCOHOL (UNII: 3K9958V90M)				
EUPHORBIA HIRTA (UNII: L13YF113GN)				
PORK SIDE BACON, CURED FAT (UNII: 6E710H379S)				
GAULTHERIA PROCUMBENS LEAF (UNII: 2125M16OWN)				
GLYCERIN (UNII: PDC6A3C0OX)				
EUPHORBIA THYMIFOLIA WHOLE (UNII: D56Y3D2OTL)				
WATER (UNII: 059QF0K00R)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53304-0301-1	1 in 1 BOX	03/01/2013	
1		100 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product		
2	NDC:53304-0301-3	1 in 1 BOX	03/01/2013	
2		15 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	03/01/2013		

Labeler - EMINENCE BIOTECH CORPORATION LIMITED (078531141)

Revised: 3/2018

EMINENCE BIOTECH CORPORATION LIMITED