BIGELOIL- menthol gel W. F. Young, Inc.

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

BIGELOIL® LIQUID GEL

Can be used before, during and after competition

Can also be used on dogs

Indications

For temporary relief of minor pain and swelling from sprains, strains, bruises, arthritis, overworked muscles and minor wounds.

Directions for Use

Rub onto sore area 3 to 4 times a day. For faster relief, rub with hair and wrap or blanket. For minor cuts and scrapes apply without rubbing.

Caution: Avoid contact with eyes and mucous membranes.

Stop using if skin irritation develops or symptoms persist for more than 10 days.

For severe injuries consult a veterinarian.

For external use only. Keep out of reach of children.

Warning

Flammable. Keep away from fire, sparks and heated surfaces.

Store at room temperature.

For animal use only.

Active Ingredient

Menthol 2%. Also contains Alcohol: 60%.

Distributed by: W. F. Young, Inc. 302 Benton Drive E. Longmeadow, MA 01028 USA © WFY

www.absorbine.com

RM 342360-5

PRINCIPAL DISPLAY PANEL - 414 mL Bottle Label

BIGELOIL® LIQUID GEL

- Reduces Pain & Swelling of Sore Muscles, Joints, Legs & Arthritis
- Excellent Under Wraps

TOPICAL PAIN RELIEF GEL

HERBAL FORMULA

14 fl oz. (414 mL)



BIGELOIL

menthol gel

Product	Inform	ation
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Product Type OTC ANIMAL DRUG Item Code (Source) NDC:11444-136

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name

Basis of Strength

FORM (UNII: L7T10EIP3A) (MENTHOL, UNSPECIFIED

MENTHOL,

2 q

MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A) (MENTHOL, UNSPECIFIED MENTHOL, UNSPECIFIED FORM - UNII:L7T10EIP3A)

2 g in 112 mL

Packaging

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l	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	-	NDC 11444 126 01	ATA I I DOTTLE DI ACTIC		

1 NDC:11444-136-01 414 mL in 1 BOTTLE, PLASTIC

Marketing Information

Marketing	Application Number or Monograph Citation	Marketing Start	Marketing End
Category		Date	Date
UNAPPROVED DRUG OTHER		01/01/1997	

Labeler - W. F. Young, Inc. (001114669)

Revised: 11/2024 W. F. Young, Inc.