BOIL EASE- benzocaine ointment
Insight Pharmaceuticals LLC

Boil Ease Ointment - 63736-040

BOIL EASE®
PAIN RELIEVING OINTMENT

Drug Facts

Active ingredient

Benzocaine 20%

Purpose

Pain Reliever

Use

for the temporary relief of pain and discomfort caused by boils

Warning

For external use only

Do not use

for more than 3 days

Ask a doctor before use if you have

boils on the lips, nose, cheeks, or forehead

When using this product

avoid contact with the eyes

Stop use and ask a doctor if

- fever occurs
- redness around the boil develops
- condition worsens or does not improve
- symptoms persist for more than 3 days
- symptoms 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 (1-800-222-1222)

right away.

Directions

Adults and children 2 years of age and older	Apply to affected area no more than 2 times daily
Children under 2 years of age	Consult a doctor

Other information

Keep carton for full drug facts

Inactive ingredients

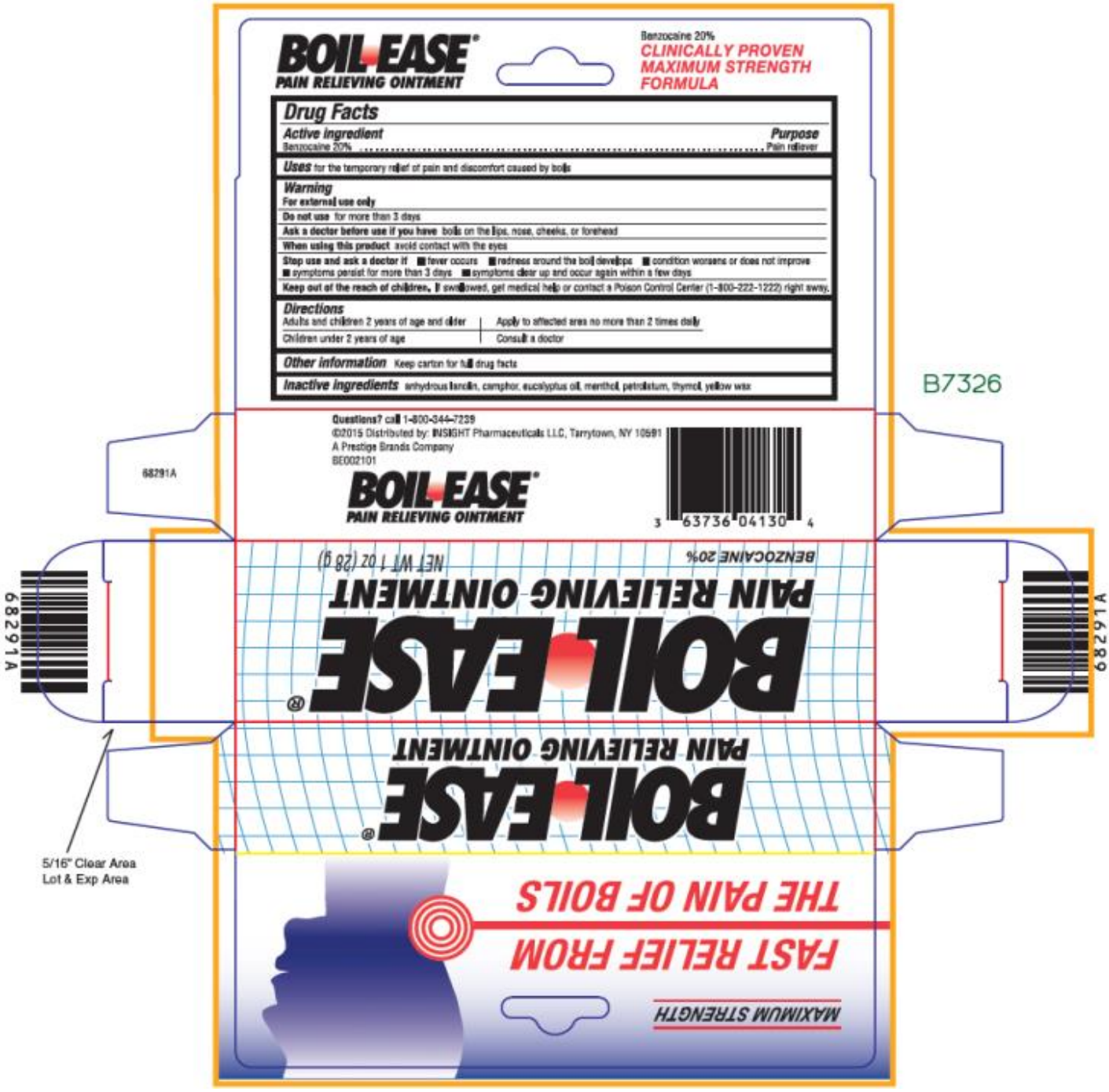
anhydrous lanolin, camphor, eucalyptus oil, menthol, petrolatum, thymol, yellow wax

Questions?

call 1-800-344-7239

PRINCIPAL DISPLAY PANEL

28 g Tube Carton
BOIL EASE®
PAIN RELIEVING OINTMENT
BENZOCAINE 20%
NET WT 1 oz (28 g)



BOIL EASE

benzocaine ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63736-040
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	20 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
LANOLIN (UNII: 7EV65EAW6H)	
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)	
EUCALYPTUS OIL (UNII: 2R04ONI662)	
MENTHOL (UNII: L7T10EIP3A)	
PETROLATUM (UNII: 4T6H12BN9U)	
THYMOL (UNII: 3J50XA376E)	
YELLOW WAX (UNII: 2ZA36H0S2V)	

Product Characteristics

Color	yellow	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63736-040-28	1 in 1 BOX	06/21/2010	
1		28 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	06/21/2010	

Labeler - Insight Pharmaceuticals LLC (055665422)

Revised: 10/2024 Insight Pharmaceuticals LLC