

BOIL EASE- benzocaine ointment
Insight Pharmaceuticals LLC

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.

Boil Ease Ointment

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 NOT FINAL	part348	06/21/2010	

Labeler - Insight Pharmaceuticals LLC (055665422)

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Insight Pharmaceuticals LLC