CVS MAXIMUM STRENGTH BOIL RELIEF- benzocaine ointment
CVS Pharmacy, Inc.

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.

CVS Boil Relief Ointment 1 oz 48551 ZDP

Active ingredient Purpose

Benzocaine 20%..... Pain Reliever

Uses for the temporary relief of pain and discomfort caused by boils

Warnings

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 presist 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 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, eucalyptol, menthol, petrolatum, thymol, yellow wax

DISTRIBUTED BY:

CVS PHARMACY, INC.

ONE CVS DRIVE

WOONSOCKET, RI 02895

CVS MAXIMUM STRENGTH BOIL RELIEF

benzocaine ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:59779-444
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**
YELLOW WAX (UNII: 2ZA36H0S2V)	
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)	
EUCALYPTUS OIL (UNII: 2R04ONI662)	
LANOLIN (UNII: 7EV65EAW6H)	
MENTHOL (UNII: L7T10EIP3A)	
THYMOL (UNII: 3J50XA376E)	
PETROLATUM (UNII: 4T6H12BN9U)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:59779-444-28	1 in 1 TUBE	10/13/2014	
1		28 g in 1 BOX; 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	10/13/2014	

Labeler - CVS Pharmacy, Inc. (062312574)

Revised: 3/2022 CVS Pharmacy, Inc.