WALGREENS PAIN BURN ITCH RELIEF- benzocaine 20%, menthol 0.5% spray
Walgreens

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.

Walgreens Pain Burn Itch Relief Spray

20% Benzocaine

0.5% Menthol

Topical Analgesic

For temporary relief of pain and itching associated with :

- sunburn
- insect bites
- minor cuts
- scrapes
- minor burns
- minor skin irritations

For External use only

Flamable:

- Do not use near heat, flame, or fire or while smoking

Allergy Alert:

- Do not use this product if you have a history of allergy to local anesthetics such as procaine, benzocaine or other "caine" anesthetics.

Avoid contact with eyes. Do not spray in the face or mouth

Use only as ditected

Intentional misuse by delibrately concentrating or inhailing the contents can be harmful or fatal

Do not puncture or incinerate.Contents under pressure. Do not store at temperatures above 120°F.

Stop use and ask a doctor if

- condition worsens or symptoms persist for more than 7 days
- symptoms clear up and occur again within a few days
- itiching, rash or irritation develops

If swallowed, get medical help or contact a Poision Control Center right away. (1-800-222-1222)

Adults and children 2 years of age and older : Apply to the affected area not more than 3 to 4 times daily

Children under 2 years of age: Cinsult a doctor

- To use this product, hold the can 6 to 12 inches away from the affected area. Direct spray nozzle towards skin and press button to activate spray.
- To apply to face, spray in palm of hand and gently apply.

- Avoid contact with leather, fabric and upholstery to prevent possible staining or discoloration
- Store at 20-25°C (68-77°F)

Aloe Vera, Phenoxyethanol, Polyethylene Glycol 400, Tetrafluoroethane

1-800-925-4733

WALGREENS PAIN BURN ITCH RELIEF

benzocaine 20%, menthol 0.5% spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0363-2053
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	5 mg in 1 g
BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	200 mg in 1 g

Inactive Ingredients	
Ingredient Name	**Strength**

PHENOXYETHANOL (UNII: HIE492ZZ3T)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
NORFLURANE (UNII: DH9E53K1Y8)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-2053-02	78 g in 1 CAN; Type 0: Not a Combination Product	05/05/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	05/05/2022	

Labeler - Walgreens (008965063)

Revised: 6/2022 Walgreens