

MEDI-FIRST STING RELIEF- benzocaine 6%, isopropyl alcohol 60% swab
Unifirst First Aid Corporation

Medi-First Sting Relief Wipes

Drug Facts

Active ingredient

Benzocaine 6% w/v

Isopropyl alcohol 60% w/v

Purpose

Topical anesthetic

Antiseptic

Uses

For temporary relief of pain and itching associated with minor burns, scrapes and insect bites.

Warnings

For external use only.

Flammable, keep away from fire or flame.

Do not use

- in the eyes. If contact occurs, rinse thoroughly with water.

Stop use and ask a doctor if

- irritation, redness or other symptoms develop
- the condition persists or gets worse

Keep out of 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 3-4 times daily.

Children under 2 years: Consult physician.

Other information

- store at room temperature 59°-86°F (15°-30°C)
- tamper evident sealed packets
- do not use any opened or torn packets

Inactive ingredients

purified water

Questions or comments?

1-800-634-7680

Medi-First Sting Relief Wipes Label

Medi-First®

Sting Relief Wipes

Benzocaine/ Isopropyl Alcohol

Topical Anesthetic/ Antiseptic

Product # 23112

NET CONTENTS: 10 wipes per box

Sting Relief Wipes

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10 per Box

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Stop use and ask a doctor if irritation, redness or other symptoms develop the condition persists or gets worse
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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:47682-234
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength

BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)		BENZOCAINE	0.6 mg in 1 mL	
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)		ISOPROPYL ALCOHOL	0.6 mL in 1 mL	
Inactive Ingredients				
Ingredient Name		Strength		
WATER (UNII: 059QF0KO0R)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47682-234-99	0.6 mL in 1 PACKET; Type 0: Not a Combination Product	07/02/2018	
2	NDC:47682-234-12	10 in 1 BOX	07/02/2018	
2		0.6 mL in 1 PACKET; Type 0: Not a Combination Product		
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug		M003	07/02/2018	

Labeler - Unifirst First Aid Corporation (832947092)

Revised: 9/2025

Unifirst First Aid Corporation