

HURRICaine ONE- topical anesthetic and antiseptic spray
Beutlich Pharmaceuticals LLC

HC1 - Lidocaine

Drug Facts

Active Ingredients

Lidocaine 4%

Benzalkonium Chloride 0.13%

Purpose

Topical Anesthetic/Analgesic

Topical First-Aid Antiseptic

Uses

Uses:

- For the temporary relief of pain
- First aid to help protect against infection in minor cuts, scrapes, and burns

Warnings

- For external use only. Do not use In the eyes or apply over large areas of the body, particularly over raw surfaces or blistered areas. In case of deep or puncture wounds, animal bites, or serious burns, consult a doctor.
- If condition worsens, or If symptoms persist for more than 7 days or clear up and occur again within a few days, discontinue use of this product and consult a doctor. Do not use longer than 1 week unless directed by a doctor.
- Flammable. Keep away from fire or flame.

Do not use

Do not Use

- if foil pouch has been previously opened
- in children under 2 years of age

When using this product

- avoid contact with eyes

Keep out of reach of children.

If more than used for pain is accidentally swallowed, get medical help or contact a Poison Control Center Immediately

Directions

- do not exceed recommended dosage

adults and children 2 years of age and older: Clean the affected area. Spray a small amount of this product on the area 1 to 3 times daily. May be covered with a sterile bandage. If bandaged, let dry first.

children under 2 years of age: do not use, consult a physician

Other information

- store at 15-30 °C (59-86 °F)

Inactive ingredients

Polyethylene glycol, Ethanol

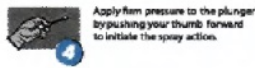
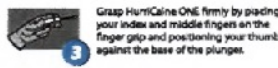
Questions or comments?

1-800-238-4582

M-F: 8:00 a.m. - 4:30 p.m. ET

Principal Display Panel

DIRECTIONS



NDC 0283-1024-11

Beutlich
Pharmaceuticals, LLC

Manufactured for
Beutlich Pharmaceuticals, LLC
Bunnell, FL 32110



HURRICaine ONE®

**4% Lidocaine Topical Analgesic
& Antiseptic Spray**

PRODUCT DIAGRAM



2 Unit-Dose Non-Aerosol Sprays – 0.013 fl. oz. (0.37 mL) each



HurriCaine ONE® is a trademark of
Beutlich Pharmaceuticals, LLC.
7775 S. US Hwy 1, Suite H, Bunnell, FL 32110
www.beutlich.com

LOT

EXP

**No Coating
in this area**



0283-1024-11

Please retain Drug Facts for future reference

Drug Facts

Active ingredient

Lidocaine 4% Topical Analgesic
Benzalkonium chloride 0.13% First Aid Antiseptic

Purpose

Uses

- For the temporary relief of pain.
- First aid to help protect against infection in minor cuts, scrapes, and burns.

Warnings

- For external use only. Do not use in the eyes or apply over large areas of the body, particularly over raw surfaces or blistered areas. In case of deep or puncture wounds, animal bites, or serious burns, consult a doctor.
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Drug Facts (continued)

Warnings (continued)

Do not use

- if foil pouch has been previously opened
- in children under 2 years of age

When using this product avoid contact with eyes

Keep out of reach of children. If accidentally swallowed, get medical help or contact a Poison Control Center immediately.

Directions

do not exceed recommended dosage

age	dose
adults and children 2 years of age and older	Clean the affected area. Spray a small amount of this product on the area 1 to 3 times daily. May be covered with a sterile bandage. If bandaged, let dry first.
children under 2 years of age	do not use, consult a physician

Other information • store at 15-30°C (59-86°F)

Inactive ingredients

Ethanol, Polyethylene glycol

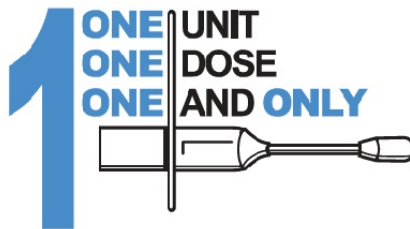
Questions or comments?

1-800-238-8542

M-F: 8:00 a.m. – 4:30 p.m. ET

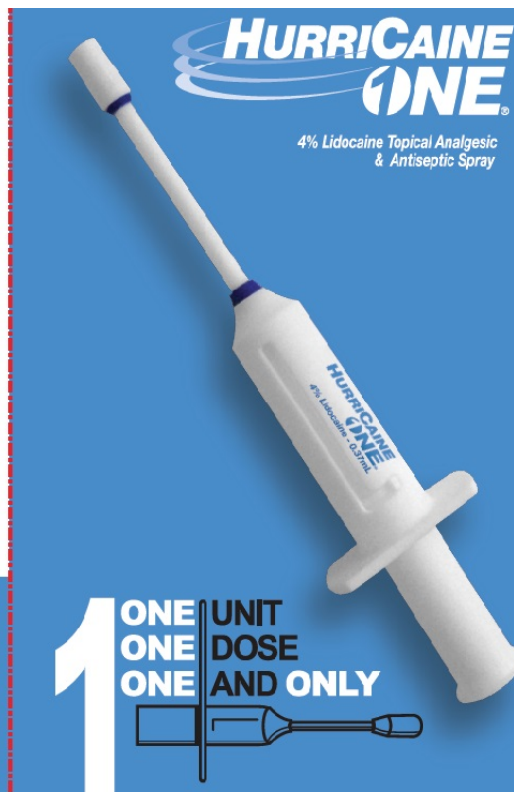
HOLB2 965 1224

Principal Display Panel



Please retain Drug Facts for future reference

Drug Facts	
Active ingredient	Purpose
Lidocaine 4%	Topical Analgesic
Benzalkonium chloride 0.13%	First Aid Antiseptic
Uses	
<ul style="list-style-type: none"> For the temporary relief of pain. First aid to help protect against infection in minor cuts, scrapes, and burns. 	
Warnings	
<ul style="list-style-type: none"> For external use only. Do not use in the eyes or apply over large areas of the body, particularly over raw surfaces or blistered areas. In case of deep or puncture wounds, animal bites, or serious burns, consult a doctor. If condition worsens, or if symptoms persist for more than 7 days or clear up and occur again within a few days, discontinue use of this product and consult a doctor. Do not use longer than 1 week unless directed by a doctor. Flammable. Keep away from fire or flame. 	
Do not use	
if foil pouch has been previously opened in children under 2 years of age	
When using this product avoid contact with eyes	
Keep out of reach of children. If accidentally swallowed, get medical help or contact a Poison Control Center immediately.	
Directions	
do not exceed recommended dosage	dosage
adults and children 2 years of age and older	Clean the affected area. Spray a small amount of this product on the area 1 to 3 times daily. May be covered with a sterile bandage. If bandaged, let dry first.
children under 2 years of age	do not use, consult a physician
Other information - store at 15-30°C (59-86°F)	
Inactive ingredients Ethanol, Polyethylene glycol	
Questions or comments? 1-800-238-8542 M-F 8:00 a.m. - 4:30 p.m. ET	



0283-1024-25

LOT
EXP

No Coating
in this area

Beutlich
PHARMACEUTICALS LLC
Since 1955

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7775 S. US Hwy 1, Suite 10, Bunnell, FL 32110, www.beutlich.com

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HURRICANE ONE

topical anesthetic and antiseptic spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0283-1024
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z 41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	40 mg in 1 g

BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)		BENZALKONIUM CHLORIDE	1.3 mg in 1 g	
Inactive Ingredients				
Ingredient Name			Strength	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)			200 mg in 1 g	
ALCOHOL (UNII: 3K9958V90M)			758.7 mg in 1 g	
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0283-1024-11	2 in 1 BOX	12/10/2025	
1	NDC:0283-1024-43	0.37 g in 1 APPLICATOR; Type 0: Not a Combination Product		
2	NDC:0283-1024-26	25 in 1 BOX	12/10/2025	
2	NDC:0283-1024-43	0.37 g in 1 APPLICATOR; 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	12/10/2025	

Labeler - Beutlich Pharmaceuticals LLC (005209325)

Registrant - Beutlich Pharmaceuticals LLC (005209325)

Establishment

Name	Address	ID/FEI	Business Operations
Beutlich Pharmaceuticals, LLC		005209325	pack(0283-1024) , label(0283-1024)

Revised: 12/2025

Beutlich Pharmaceuticals LLC