BURN GEL- lidocaine hci gel
Trifecta Pharmaceuticals USA LLC

Globe First Aid Burn Gel

Active Ingredient

Lidocaine HCL 2%

Purpose

Topical Pain Relief

Uses

Temporary pain relief for minor burns. For professional use only.

Warnings

For external Use Only

Do not use

- in large quantities, particularly over raw or blistered areas
- near eyes, if this happens rinse thoroughly with water

Directions

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

Children under 2 years: do not use, consult a doctor

Inactive Ingredients

Aloe barbadensis leaf juice, Carbomer, Ethylhexylglycerin, Maltodextrin, Menthol, Phenoxyethanol, Propylene glycol, purified water, Triethanolamine, Vitamin E acetate

Questions?

Call 1-888-296-9067

Stop Use and Ask a doctor if

The condition worsens or symptoms persist for more than 7 days or clear up and occur again within a few days.

Storage Information

- Store at room temperature (do not freeze)
- do not use any opened or torn packets

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center immediately.

Other Information

Distributed By:

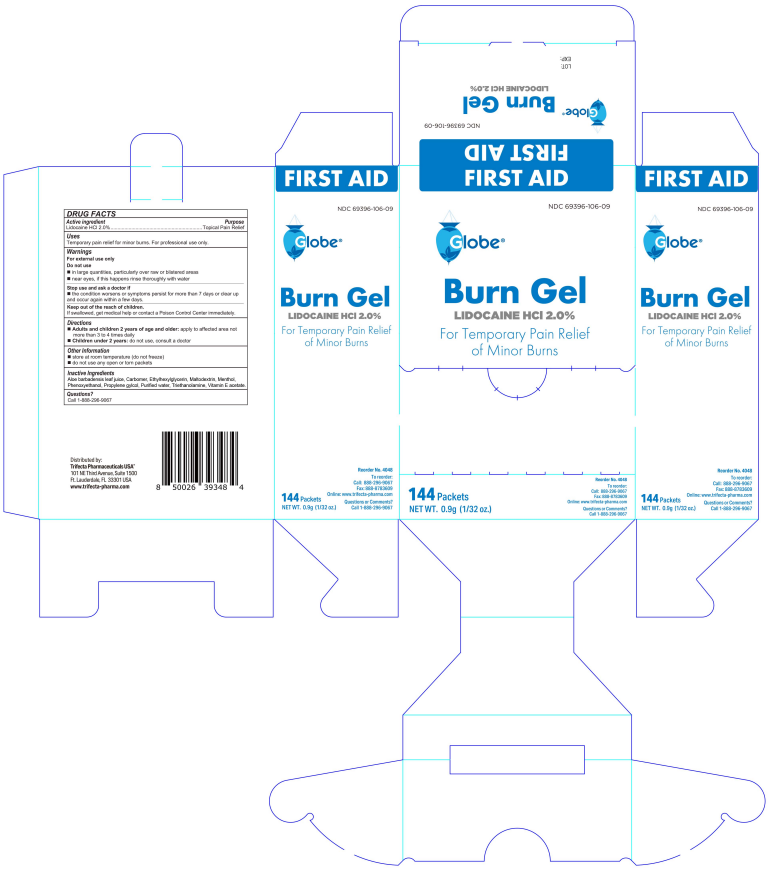
Trifecta Pharmaceuticals USA®

101 NE Third Ave. Ste 1500

Ft. Lauderdale, FL. 33301 USA

www.trifecta-pharma.com

OUTSIDE BOX



INNER PACKET

BURN GEL

lidocaine hci gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69396-106

Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	0.02 g in 1 g

Inactive Ingredients

Ingredient Name	Strength
.ALPHA.-TOCOPHEROL (UNII: H4N855PNZ1)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
MENTHOL (UNII: L7T10EIP3A)	
WATER (UNII: 059QF0KO0R)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
TROLAMINE (UNII: 9O3K93S3TK)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69396-106-09	144 in 1 CARTON	09/27/2022	
1		0.9 g 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	M017	09/27/2022	

Labeler - Trifecta Pharmaceuticals USA LLC (079424163)

Revised: 1/2024 Trifecta Pharmaceuticals USA LLC