

EQUATE BURN RELIEF- lidocaine hydrochloride gel

Walmart Inc

Walmart 79903-237-08

Active ingredient

Lidocaine HCl 0.5%

Purpose

External analgesic

Uses

for the temporary relief of pain and itching associated with

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

Warnings

For external use only.

Do not use in large quantities, particularly over raw surfaces or blistered areas

When using this product avoid contact with eyes

Stop use and ask a doctor if condition worsens, or if symptoms persist for more than 7 days or clear up and occur again within a few days

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 no more than 3 to 4 times daily
- children under 2 years of age: ask a doctor

Inactive ingredients

Aloe Barbadensis Leaf Juice, Blue 1, Carbomer, Disodium EDTA, Ethylhexylglycerin, Glycerin, Maltodextrin, Menthol, Phenoxyethanol, Polysorbate 20, Propanediol, SD Alcohol, Sodium Hydroxide, Water, Yellow 5



label

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Questions? 1-888-287-1915

DISTRIBUTED BY: Walmart Inc., Bentonville, AR 72716

PRODUCT OF CHINA

Information: 1-888-287-1915 or Walmart.com/help

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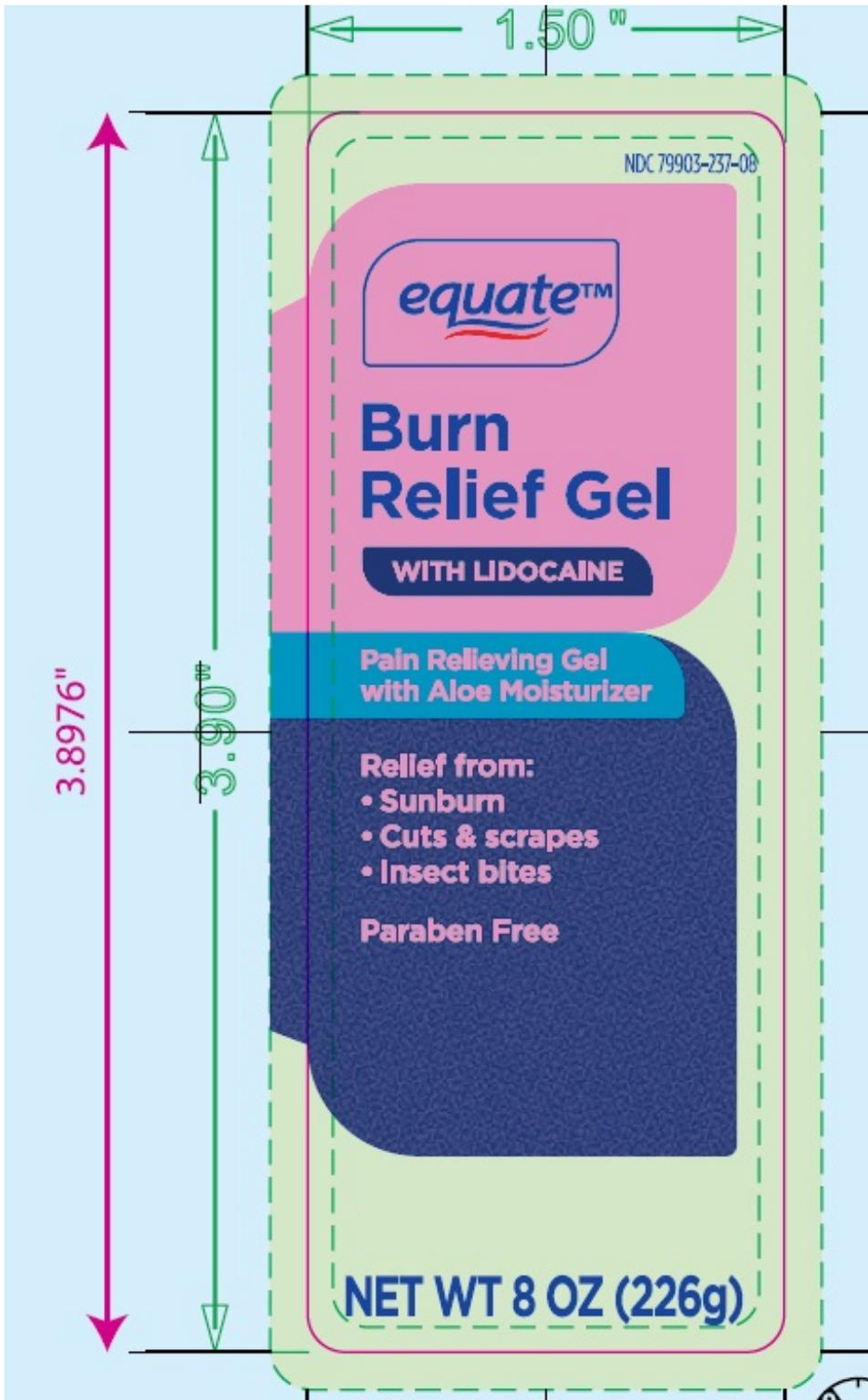
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EQUATE BURN RELIEF

lidocaine hydrochloride gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79903-237
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	0.5 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALOE (UNII: V5VD430YW9)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
CARBOMER 934 (UNII: Z135WT9208)	
DISODIUM HEDTA (UNII: KME849MC7A)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
MENTHOL (UNII: L7T10EIP3A)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
PROPANEDIOL (UNII: 5965N8W85T)	
ALCOHOL (UNII: 3K9958V90M)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79903-237-08	226 g in 1 BOTTLE; Type 0: Not a Combination Product	11/15/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	11/15/2023	

Labeler - Walmart Inc (051957769)**Establishment**

Name	Address	ID/FEI	Business Operations
Nantong Health & Beyond Hygienic Products Inc.		421280161	manufacture(79903-237)

Revised: 10/2025

Walmart Inc