BIKINI ZONE MEDICATED GEL- lidocaine gel
Kingsway Pharmaceuticals dba NFI, LLC

Bikini Zone® Medicated Gel

Active Ingredients

Lidocaine HCL 2.0% w/w

Purpose

Topical Analgesic

Uses

For temporary relief of pain and itching.

Warnings

For external use only

Avoid contact with eyes

Do not use

in large quantities, particularly over raw surfaces or blistered areas

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. Discontinue use.

Keep out of reach of children

- If product is swallowed, get medical help or contact a Poison Control Center right away.

Directions

For adults and children two-years or older: Apply to affected area not more then 3 to 4 times daily. Children under 2 years of age: consult a physician.

Other information

Protect this product from excessive heat and direct sun.
Store at room temperature 15°-25°C (59°-77°F)

Inactive Ingredients

Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aloe Barbadensis Leaf Extract, Camphor, Ethylhexylglycerin, Glycerin, Menthol, Phenoxyethanol, Propylene Glycol, Salicylic Acid, SD Alcohol 40-B, Sodium Phytate, Triethanolamine, Water, Yarrow Extract, Yucca Extract

Questions or Comments?

(888) 990-2454

Package/Label Principal Display Panel – Carton Label

Bikini Zone®

SOOTHING
MEDICATED
GEL

For use after any type
of hair removal

LIDOCAINE
Topical Analgesic

Instantly
relieves pain
associated with:

- ✓ Waxing
- ✓ Shaving
- ✓ Electrolysis
- ✓ Depilation

HELPS RELIEVE
BIKINI BUMP IRRITATION

NET WT. 1 OZ (28 G)

Distributed By:
Kingsway Pharmaceuticals
dba NFI Consumer Products
636 Shelby Street, Suite 300
Bristol, TN 37620

Made in the U.S.A

BIKINI ZONE MEDICATED GEL

lidocaine gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69993-044
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	2 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CAMPHOR (NATURAL) (UNII: N20HL7Q941)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
GLYCERIN (UNII: PDC6A3C0OX)	
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SALICYLIC ACID (UNII: O414PZ4LPZ)	
ALCOHOL (UNII: 3K9958V90M)	
PHYTATE SODIUM (UNII: 88496G1ERL)	
TROLAMINE (UNII: 9O3K93S3TK)	
WATER (UNII: 059QF0KO0R)	
ACHILLEA MILLEFOLIUM WHOLE (UNII: 2FXJ6SW4PK)	
YUCCA SCHIDIGERA WHOLE (UNII: 08A0YG3VIC)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69993-044-01	1 in 1 CARTON	09/01/2025	
1		28 g in 1 TUBE; 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	09/01/2025	

Labeler - Kingsway Pharmaceuticals dba NFI, LLC (121681919)

Revised: 8/2025 Kingsway Pharmaceuticals dba NFI, LLC