NUMB SKIN - lidocaine and benzethonium chloride cream cream
Seenext Venture Ltd

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Drug Facts

Active Ingredient

Lidocaine 4%

benzethonium Chloride 0.25%

Purpose

External Analgesic and First Aid Antiseptic

Uses

- Can be used instead of soap and water to help clean minor cuts, scrapes, and burns
- For the temporary relief of discomfort and pain associated with dermal procedures such as tattoo removal, dermarolling, electrolysis, microblading, and piercing
- Temporarily relieves pain and itch while helping to prevent infection.

Warnings

- **For external use only**
- avoid contact with eyes

Do not use

- do not use in large quantities particularly over raw surfaces or blistered area
- do not exceed the recommended dosage unless directed by a doctor
- in the eyes or apply over large areas of body
- longer than one week unless directed by a doctor

Ask a doctor before use if you have

- deep or puncture wounds, animal bites or serious burns

Stop the use and consult doctor if:

- condition worsenss or symptoms persist for more than 7 days or clear up and occur again within few days.

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away

Directions

- Use NumbSkin® Topical Anesthetic Foam Soap to cleanse the targeted area.
- Shake well before each use.
- Apply 2 to 3 pumps of foam soap andgently rub into the skin. Leave it onfor 5 to 10 minutes. Rinse off gentlywith running water or wipe with cleanpaper towel.
- Use before, during andafter the procedure.
- Make sure tocover the bottle tightly when not inuse, otherwise, the anesthetic effect will be lessened.

Adults and children two years old and older: Use to clean minor cuts, scrapes, and burns by thoroughly washing with water. Rinse and air dry. Use no more than three times daily.

Children under two years of age, ask a doctor.

Other information

- Store at controlled room temperature 59 ^{0}-86 ^{0}F (15 ^{0}-30 ^{0}C)
- Do not expose to temperature above 120 ^{0}F(49 ^{0}C)
- Protect from freezing

Inactive Ingredients

Allantoin, Aloe Barbadensis Leaf Juice, Arnica Montana Flower Extract, Benzyl Alcohol, Glycerin , Lauramidopropyl Betaine, Leuconostoc/Radish Root Ferment Filtrate, Menthoxypropanediol, Phenoxyethanol, Selaginella Lepidophylla Extract, Sodium Hydroxide, Water.

Question or comments?

Call weekdays 9 AM to 6 PM PST at 1-844-700- NUMB(6862) or email us at support@numbskin.com

Principal Display Panel

NDC 70907-002-50

NUMBSKIN

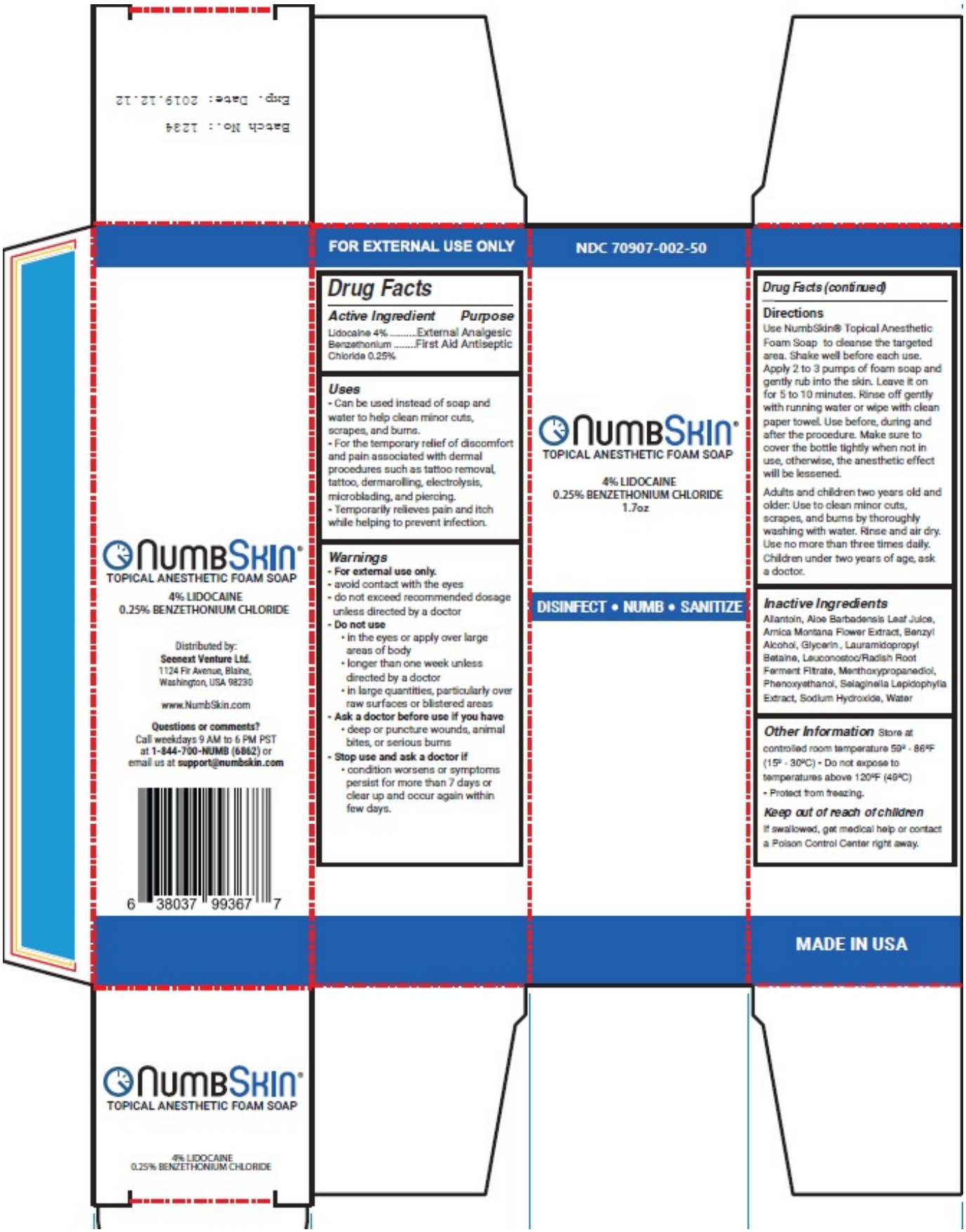
TOPICAL ANESTHETIC FOAM SOAP

4% LIDOCAINE

0.25% BENZETHONIUM CHLORIDE

DISINFECT NUMB SANITIZE

1.7 oz



NUMB SKIN

lidocaine and benzethonium chloride cream cream

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70907-002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	4 mg in 50 g
BENZETHONIUM CHLORIDE (UNII: PH41D05744) (BENZETHONIUM - UNII:1VU15B70BP)	BENZETHONIUM CHLORIDE	0.25 mg in 50 g

Inactive Ingredients

Ingredient Name	Strength
ALLANTOIN (UNII: 344S277G0Z)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
GLYCERIN (UNII: PDC6A3C0OX)	
LAURAMIDOPROPYL BETAINE (UNII: 23D6XVI233)	
LEUCONOSTOC/RADISH ROOT FERMENT FILTRATE (UNII: D2QHA03458)	
3-((L-MENTHYL)OXY)PROPANE-1,2-DIOL (UNII: KD6TZ2QICH)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
SELAGINELLA LEPIDOPHYLLA (UNII: 02JQ564P1G)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70907-002-50	1 in 1 CARTRIDGE	09/11/2017	
1		50 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 not final	part348	09/11/2017	

Labeler - Seenext Venture Ltd (203416862)

Registrant - Seenext Venture Ltd (203416862)