

NICK RELIEF - aluminum sulfate stick
220 LABORATORIES INC

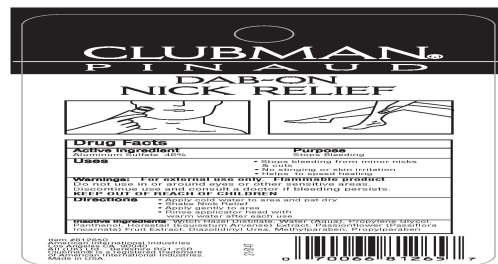
Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

NICK RELIEF PACKAGING

Warnings: For external use only. Flammable product.
Do not use in or around eyes or sensitive areas.
Discontinue use and consult a doctor if bleeding persists.



FRONT



BACK

PMS 349
PMS 485
PMS 877
PMS BLACK



Directions:

Apply cold water to area and pat dry.
Shake Nick Relief.
Apply gently to area.
Rinse applicator head with warm water after each use.

Uses:

Stops bleeding from minor nicks and cuts.
No stinging or skin irritation.
Helps to speed healing.

Active Ingredient: Aluminum Sulfate 46%

Inactive ingredients: Witch Hazel Distillate, Water (Aqua), Propylene Glycol, Panthenol, Horsetail (Equisetum Arvense) Extract, Passionflower (Passiflora Incarnata) Fruit Extract, Diazolidinyl Urea, Methylparaben, Propylparaben

KEEP OUT OF REACH OF CHILDREN

Purpose: Stops bleeding

Clubman

Nick Relief

For razor nicks and cuts

Stops Bleeding Instantly

**CLUBMAN.
NICK RELIEF**
For Razor Nicks and Cuts
Stops Bleeding Instantly
7 ml e .25 fl oz

Directions: Apply cold water to area and pat dry. Gently shake Nick Relief. Dab on affected area. Rinse applicator head with warm water after each use.

CAUTION: For external use only
AVOID EYE AREA
KEEP OUT OF REACH OF CHILDREN
ACTIVE INGREDIENT: Aluminum Sulfate 46%

Item # 812650
American International Industries
Clubman® is a registered trademark of
American International Industries, Los Angeles, CA 90040
All (UK) Ltd. Berkshire RG1 7SR
www.aiibeauty.com Made in USA

23-2421

NICK RELIEF

aluminum sulfate stick

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:57367-007
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALUMINUM SULFATE (UNII: 34S289N54E) (ALUMINUM CATION - UNII:3XHB1D032B)	ALUMINUM SULFATE	46 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
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WATER (UNII: 059QF0KO0R)
WITCH HAZEL (UNII: 101I4J0U34)
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)
DIAZOLIDINYLUREA (UNII: H5RIZ3MPW4)
PANTHENOL (UNII: WV9CM0O67Z)
METHYL PARABEN (UNII: A2I8C7HI9T)
PROPYL PARABEN (UNII: Z8IX2SC1OH)
MELISSA OFFICINALIS (UNII: YF70189L0N)
MATRICARIA RECUTITA (UNII: G0R4UBI2ZZ)
ALOE VERA LEAF (UNII: ZY81Z83H0X)
GERANIUM EXTRACT (UNII: 6H429I0Y3L)
QUERCUS ALBA (UNII: XR6BC2ZUAM)
ROSA CANINA FRUIT (UNII: 3TNW8D08V3)
SAGE (UNII: 065C5D077J)
COMFREY (UNII: D05HXX6R3G)
YARROW FLOWER (UNII: YQR8R0SQEA)
EAST INDIAN LEMONGRASS OIL (UNII: UP0M8M3VZW)
.ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:57367-007-20	1 in 1 BLISTER PACK		
1	NDC:57367-007-07	7 mL in 1 APPLICATOR		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part347	11/27/1997	

Labeler - 220 LABORATORIES INC (783247950)

Registrant - 220 LABORATORIES INC (783247950)

Establishment

Name	Address	ID/FEI	Business Operations
220 LABORATORIES INC		783247950	manufacture