REDICARE BLOOD CLOTTING- blood clotting spray aerosol, spray
Redicare LLC

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.

Blood Clotting Spray

Active Ingredients

Benzethonium Chloride 0.2% w/w

Lidocaine 4% w/w

Purpose

Topical Antiseptic

Topical Antiseptic

Uses

- for help in the control of superficial bleeding
- for temporary relief of pain and to help protect against skin infection in minor cuts, minor scrapes, and minor burns

WARNINGS

For external use only.

Flammable

- Keep away from flame. Contents under pressure.
- Do not puncture or incinerate.
- Do not expose to temperatures above 120°(F)

DO NOT USE

- in eyes or other mucous membranes
- on large portions of the body
- in case of deep or puncture wounds
- on raw surfaces
- on blistered areas
- on animal bites
- on serious burns

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Stop use and ask doctor if:
- Redness, swelling, or irritation occurs
- Infection occurs
- The condition persists or worsens
- Symptoms persist for more than 7 days

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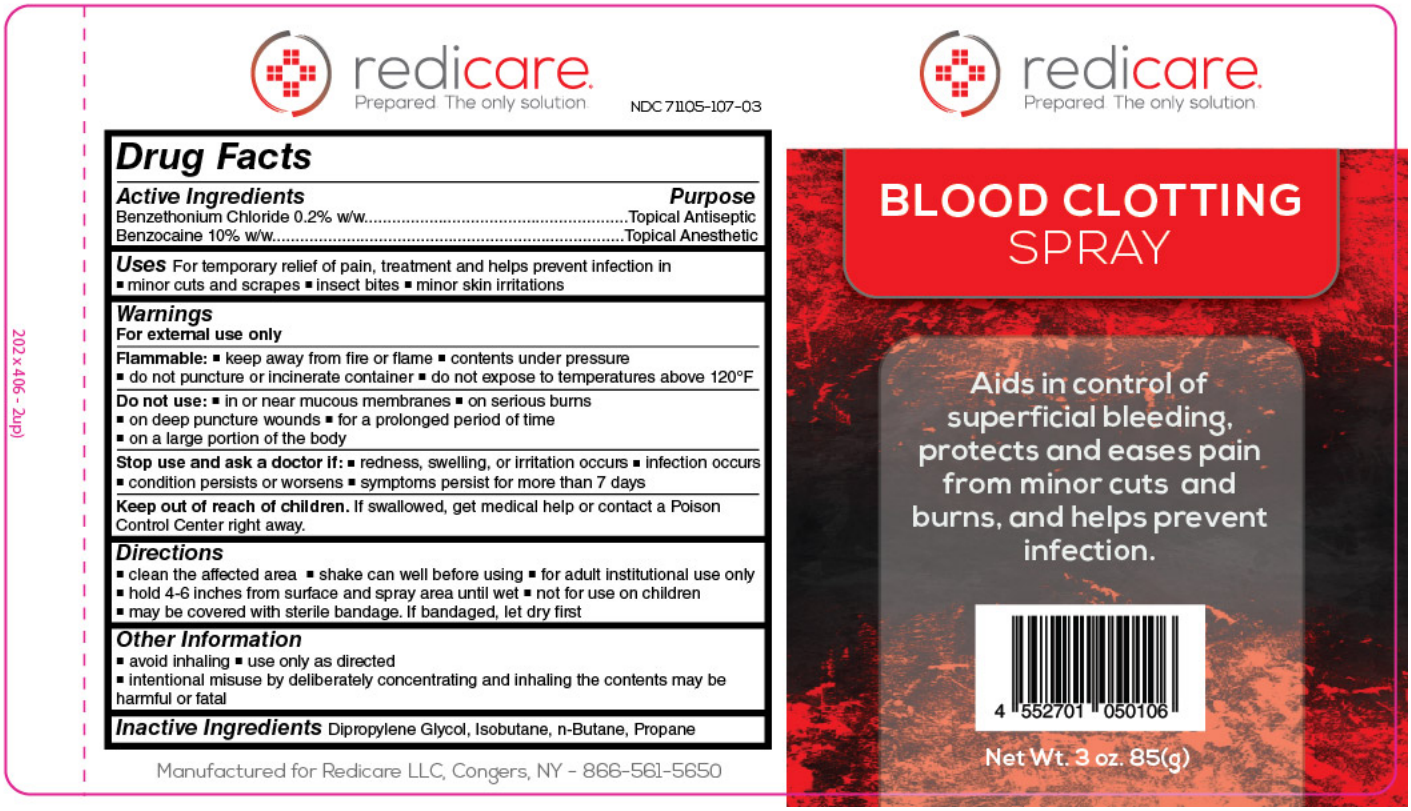
Keep out of reach of children.

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

Directions
- shake well before using
- hold 6-8 inches from moist injured area
- spray until the area is covered
- Scab-like cover will gradually disappear as healing takes place or may be removed with soap and water
- for adult institutional use only
- not intended for use on children

Inactive Ingredients

Isobutane, Isopropyl Myristate, Karaya, N-Butane, Propane, Silica, Silicone, Sorbitan Monooleate, Talc, Tragacanth



REDICARE BLOOD CLOTTING

blood clotting spray aerosol, spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71105-107
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZETHONIUM CHLORIDE (UNII: PH41D05744) (BENZETHONIUM - UNII:1VU15B70BP)	BENZETHONIUM CHLORIDE	170 mg in 85 g
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	3400 mg in 85 g

Inactive Ingredients	
Ingredient Name	**Strength**
PROPANE (UNII: T75W9911L6)	
3-(TRIETHOXYSILYL)PROPYLAMINE (UNII: L8S6UBW552)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
TALC (UNII: 7SEV7J4R1U)	
ISOBUTANE (UNII: BXR49TP611)	
BUTANE (UNII: 6LV4FOR43R)	
SORBITAN MONOOLEATE (UNII: 06XEA2VD56)	
KARAYA GUM (UNII: 73W9IQY50Q)	

SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
TRAGACANTH (UNII: 2944357O2O)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71105-107-03	85 g in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	10/01/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	10/01/2023	

Labeler - Redicare LLC (800149346)

Registrant - Redicare LLC (800149346)

Revised: 9/2023 Redicare LLC