CA-REZZ - NORISC - INCONTINENT WASH- benzethonium chloride solution FNC MEDICAL CORPORATION

FNC - CA-REZZ - NORISC - INCONTINENT WASH (60762-113)

ACTIVE INGREDIENT

BENZETHONIUM CHLORIDE 0.12%

PURPOSE

ANTIBACTERIAL

USES

CA-REZZ NORSIC WASH HAS BEEN FORMULATED AS AN INCONTINENT WASH TO QUICKLY EMULSIFY BODY WASTE EVEN IN THE MOST SENSITIVE PERINEAL AREAS.

CA-REZZ NORSIC WASH IS PH BALANCED AND ENRICHED WITH ALOE VERA, TO SOOTHE AND HELP PROMOTE HEALING OF FRAGILE, IRRITATED SKIN. STOPS ODORS.

WARNINGS

FOR EXTERNAL USE ONLY. IF RASH OR IRRITATION DEVELOP, DISCONTINUE USE. CONSULT A PHYSICIAN IF IRRITATION PERSISTS.

KEEP OUT OF REACH OF CHILDREN.

DIRECTIONS

- 1. SPRAY CA-REZZ NORSIC WASH WHERE URINE, FECES OR EMESIS HAS SOILED SKIN PAD LINENS AND CLOTHING.
- 2. REMOVE BULK OF FECES OR EMESIS.
- 3. SPRAY ADDITIONAL CA-REZZ NORSIC WASH ON WARM WET CLOTH. CLEANSE SKIN OF ALL REMAINING RESIDUE. PAT DRY. NOTE: IF AREA NEEDING CARE IS SIZEABLE AND FECAL OR EMESIS MATERIAL MASSIVE, RINSING IS ADVISED.
- 4. ADD ONE CAPFUL OF CA-REZZ NORSIC WASH TO THE REGULAR LAUNDRY TO ELIMINATE RESIDUAL ODORS.

OTHER INFORMATION

CA-REZZ NORSIC WASH IS ENRICHED WITH ALOE.

INGREDIENTS

DEIONIZED WATER, SODIUM LAURETH SULFATE, PROPYLENE GLYCOL, DISODIUM COCOAMPHODIACETATE, POLYSORBATE 80 (AND) CETYL ALCOHOL (AND)

QUESTIONS OR COMMENTS?

PLEASE CALL 1-800-440-2888



CA-REZZ - NORISC - INCONTINENT WASH

benzethonium chloride solution

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:60762-113
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
BENZETHONIUM CHLORIDE (UNII: PH41D05744) (BENZETHONIUM - UNII:1VU15B70BP)	BENZETHONIUM CHLORIDE	0.12 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
DISODIUM COCOAMPHODIACETATE (UNII: 18L9G3U51M)	
POLYSORBATE 80 (UNII: 60ZP39ZG8H)	
CETYL ACETATE (UNII: 4Q43814HXS)	
ACETYLATED LANOLIN ALCOHOLS (UNII: SNN716810P)	
ALOE VERA WHOLE (UNII: KIZ 4X2EHYX)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	

ı	Packaging				
	# Item Code	Package Description	Marketing Start Date	Marketing End Date	
		237 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	02/05/2018		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M003	02/05/2018		

Labeler - FNC MEDICAL CORPORATION (849207519)

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