CVS PET WOUND CARE PREP-PACK- benzalkonium chloride CVS

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.

CVS Pet Wound Care Prep-Pack

Active Ingredient

Benzalkonium Chloride .13%

Purpose

First Aid Antiseptic

Use

For Professional and Hospital use. Helps prevent infection. Antiseptic cleansing of face, hands and body without soap and water.

Warnings

For external use only. Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away. If unusual redness, swelling or other symptoms occur, cunsult a physician immediately.

Do not use

In the eyes, or over large areas of the body

Directions

Tear open packet, unfold towelette and use to cleanse desired skin are. Discard towelette appropriately after single use

Inactive Ingredients

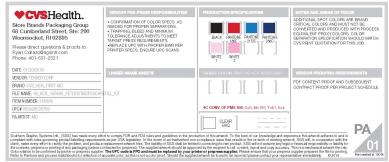
Purified Water

Antiseptic Towelette





PLEASE LEAVE OUR CVS SLUG ON ALL FINAL ART AND CONTRACT PROOFS FOR FINAL APPROVAL



Antiseptic Towelette GFA Production Xiamen Co., Ltd www.gfaproduction.com





CVS PET WOUND CARE PREP-PACK

benzalkonium chloride kit

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:69842-405

Packaging

	# Item Code	Item Code Package Description		Marketing End Date
П	NDC:69842-405-00 1 in 1 BAG; Type 0: Not a Combination Product		12/01/2016	

Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	8 PACKAGE	6 mL in .8

Part 1 of 1

ANTISEPTIC

benzalkonium chloride swab

Product Information			
Item Code (Source)	NDC:52124-0001		
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7) (BENZALKO NIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL	

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

l	Packaging				
l	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
l	1	NDC:52124-0001-1	0.8 mL in 1 PACKAGE; 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	part333E	12/0 1/20 16		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph final	part333A	12/01/2016		

Labeler - CVS (062312574)

Establishment				
Name	Address	ID/FEI	Business Operations	
Tender Corporation		064437304	manufacture(69842-405)	

Revised: 12/2016 CVS