CVS HEALTH FIRST AID OUTDOOR PREP-PACK- ammonia
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 Health First Aid Outdoor Prep-Pack

Active Ingredient

Active Ingredient:

Ammonia 3.5%

Purpose

Counterirritant

Use

Temporarily protects and helps relieve minor skin irriatation and itching due to

- insect bites and stings
- poison ivy, oak or sumac

Warnings

Warning: For external use only.

Keep out of reach of children: If swallowed, get medical help or contact a Poison Control Center right away.

Stop use and ask a doctor if

- condition worsens
- symptoms last more than 7 days or clear up and occur again within a few days

When using this product

Do not get into eyes

Directions

Adults and children under 2 years and older dab directly on bite or sting, rub gently and re-apply as needed

Children under 2 years ask a doctor

Inactive Ingredients

Alcohol Ethoxylate, Dimethicone, Mineral Oil and Purified Water

After Bite

The Itch Eraser

Fast Relief from Insect Bites.

Net Contents: 0.037fl. oz.

Contains: One (1) Wipe

Contains Ammonia

Tender Corporation

Littleton, NH 03561

PMS 348

AfterBite®
The Itch Eraser.com

Fast Relief from Insect Bites.

Soulagement rapide des piqûres d'insectes.

Antipruritic / Antiprurigineux

NET CONTENTS: 0.037 fl. oz.
CONTAINS/CONTENU: one (1) wipe/serviette
HPN 02229667 NDC 044224-0001-2
Contains Ammonia/Contient Ammoniaque

Tender CORPORATION
Littleton, NH 03561 USA

DIRECTIONS: Wipe moist towelette on bitten area immediately upon opening. Apply with a wiping motion, do not hold on bitten area. Do not bandage or cover tightly until dry. KEEP OUT OF REACH OF CHILDREN. CAUTION: for external use only. Avoid mouth, eyes, or mucous membranes. If swallowed, do not induce vomiting. Drink milk and citrus juices and consult a physician. If rash, redness, irritation, swelling or pain increases, discontinue use and consult a physician. Do not apply to wounds or damaged skin. INGREDIENTS: Ammonia 3.5% w/v medicinal; Mineral Oil (prevents drying), Alcohol Ethoxylate, Dimeticone, non medicinal.

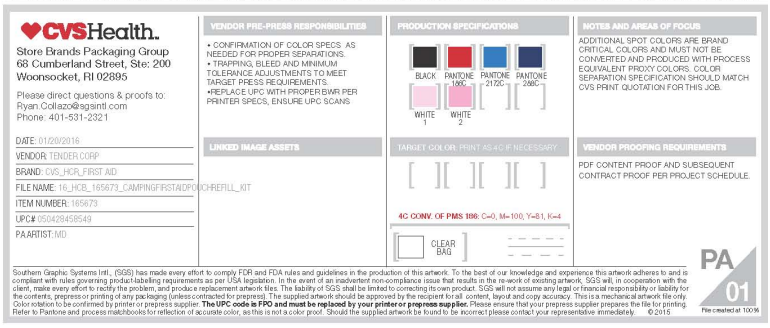
MODE D'EMPLOI: retirer la serviette humide de l'emballage et l'utiliser immédiatement pour nettoyer la plaie d'un mouvement fluide, sans presser sur la piqûre. Ne pas panser ou couvrir la plaie encore humide. HORS DE LA PORTEE DES ENFANTS. ATTENTION: Pour emploi externes enlement. Evitez le contact avec la bouche, les yeux ou membranes muqueuses. Si le produit est avale, ne provoquez pas le vomissement. Buvez du lait et des jus d'agrumes et consultez un médecin. Si l'éruption, la rougeur, l'irritation, l'enflure ou la douleur augmente, cessez l'emploi du produit et consultez un médecin. N'appliquez pas sur une blessure ouverte ou peau endommagée. INGREDIENTS: Ammoniaque 3.5% w/v (Actif), Médicinal; Huile Minéral (prévient le dessèchement), l'Alcool Ethoxylate, Dimeticone, non-medicinal.

0005-3620-1

www.tendercorp.com

TEAR HERE DÉCHIRURE ICI

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



CVS HEALTH FIRST AID OUTDOOR PREP-PACK

ammonia kit

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69842-407

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:69842-407-00	1 in 1 KIT; Type 0: Not a Combination Product	12/01/2016	

Quantity of Parts		
Part #	**Package Quantity**	**Total Product Quantity**
Part 1	4 PACKAGE	3 mL

Part 1 of 1

AFTER BITE WIPE

ammonia swab

Product Information	
Item Code (Source)	NDC:44224-0001
Route of Administration	TOPICAL

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
AMMONIA (UNII: 5138Q19F1X) (AMMONIA - UNII:5138Q19F1X)	AMMONIA	30 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
LIGHT MINERAL OIL (UNII: N6K5787QVP)	
C12-13 ALCOHOLS (UNII: T7ZJT3I9X2)	
DIMETHICONE 1000 (UNII: MCU2324216)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:44224-0001-2	0.7 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	part348	04/15/2011	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	12/01/2014	

Labeler - CVS (062312574)

Establishment

Name	Address	ID/FEI	Business Operations
Tender Corporation		064437304	manufacture(69842-407)

Revised: 1/2017 CVS