FIRST AID ONLY AMMONIA INHALANTS- ammonia inhalants inhalant
Acme United Corporation

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Ammonia Inhalants

Active Ingredient

Ammonia(15%)

Purpose

Aromatic Stimulant

Uses To prevent or treat fainting

Warnings For external use only.

Do not use •If you have breathing problems such as asthma or emphysema

Stop use and ask a doctor if condition persists

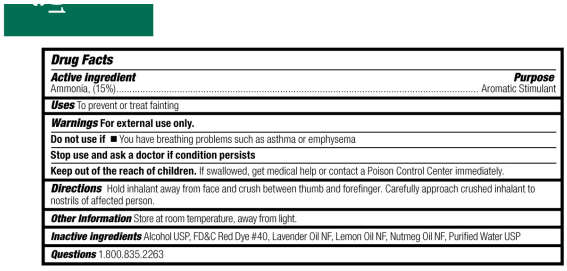
Keep out of the reach of children. If swallowed, get medical help or contact a Poison Control Center immediately

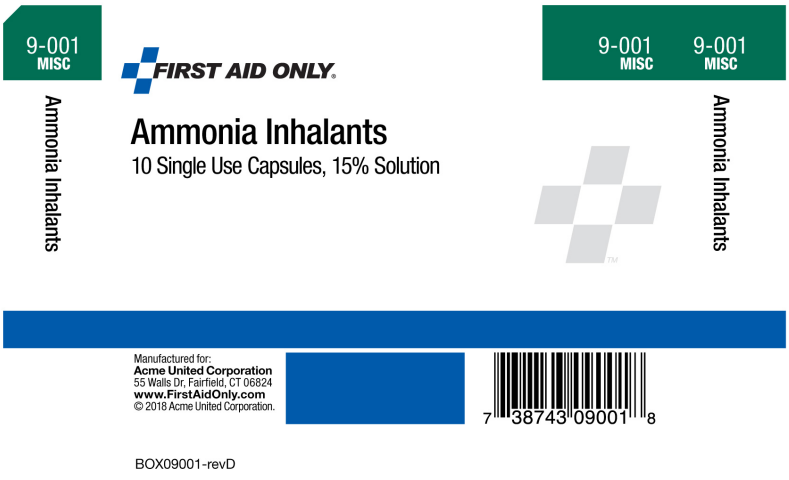
Directions Hold inhalant away from face and crush between thumb and forefinger. Carefully approach crushed inhalant to nostrils of affected person.

Other Information Store at room temperarure, away from light

Inactive Ingredients Alcohol USP, FD&C Red Dye #40, Lavender Oil NF, Lemon Oil NF, Nutmeg Oil NF, Purified Water USP

Questions 1.800.835.2263





FIRST AID ONLY AMMONIA INHALANTS

ammonia inhalants inhalant

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0924-5401(NDC:46414-3333)
Route of Administration	RESPIRATORY (INHALATION)		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
AMMONIA (UNII: 5138Q19F1X) (AMMONIA - UNII:5138Q19F1X)	AMMONIA	0.045 g in 0.3 mL

Inactive Ingredients

Ingredient Name	Strength
LEMON OIL (UNII: I9GRO824LL)	
WATER (UNII: 059QF0KO0R)	
LAVENDER OIL (UNII: ZBP1YXW0H8)	
ALCOHOL (UNII: 3K9958V90M)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
NUTMEG OIL (UNII: Z1CLM48948)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0924-5401-01	10 in 1 CARTON	01/01/2006	
1		0.3 mL in 1 AMPULE; Type 0: Not a Combination Product		
2	NDC:0924-5401-02	100 in 1 CARTON	01/01/2006	
2		0.3 mL in 1 AMPULE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		01/01/2006	

Labeler - Acme United Corporation (001180207)

Registrant - Acme United Corporation (001180207)

Establishment

Name	Address	ID/FEI	Business Operations
Acme United Corporation		045924339	relabel(0924-5401) , repack(0924-5401)

Establishment

Name	Address	ID/FEI	Business Operations
Acme United Corporation		080119599	relabel(0924-5401) , repack(0924-5401)

Revised: 2/2018 Acme United Corporation