AMMONIA INHALANTS- ammonia inhalants inhalant
James Alexander 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

Drug Facts

Active Ingredients (each inhalant)

Ammonia (15%)

Purpose

inhalant

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

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 temperature away from light.

Inactive Ingredients

Alcohol USP, FD&C Red Dye # 40, Lavender Oil FCC, Lemon Oil FCC, Nutmeg Oil FCC, Purified Water USP.

Questions?

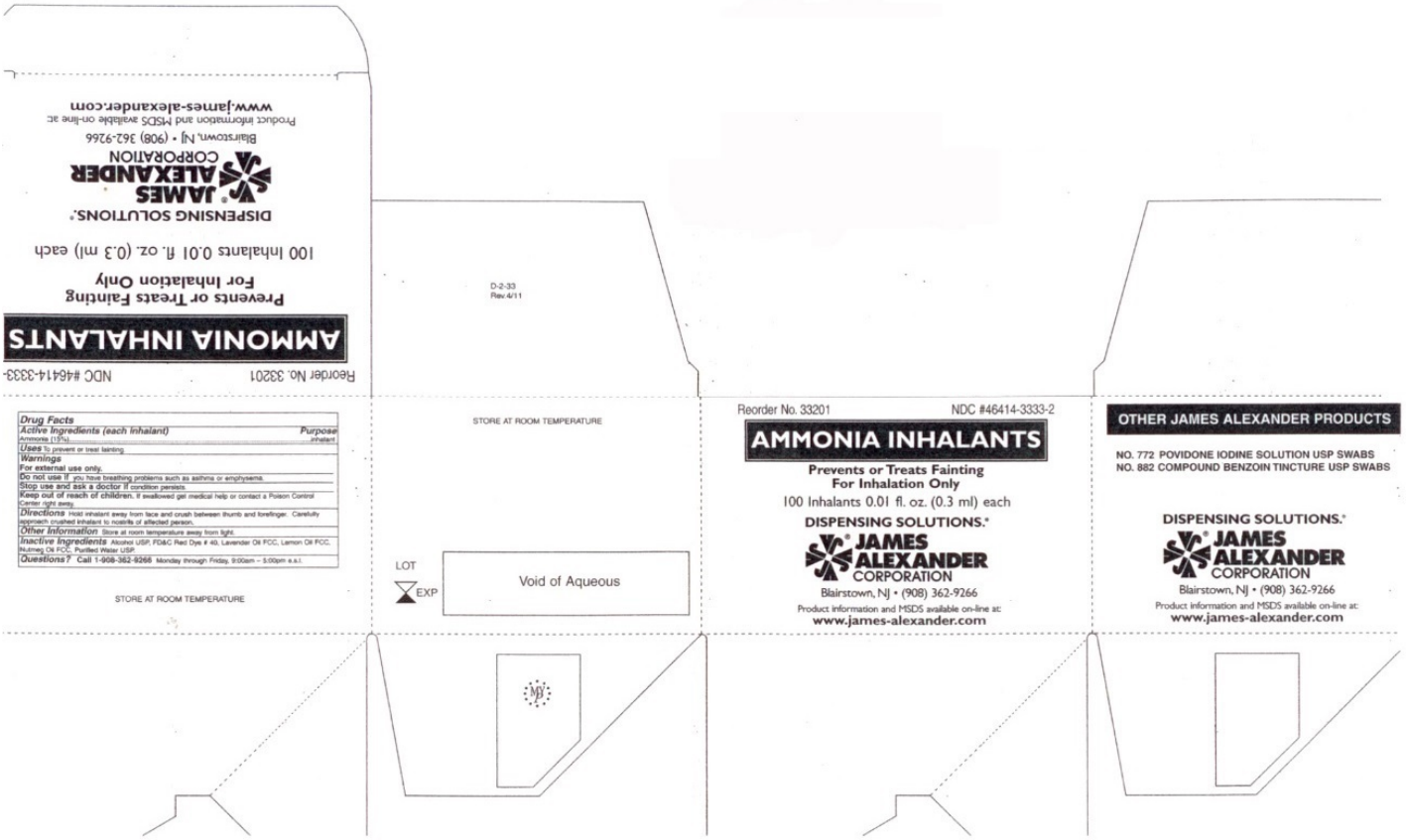
Call 1-908-362-9266 Monday through Friday. 9:00am - 5:00pm e.s.t

DISPENSING SOLUTIONS®

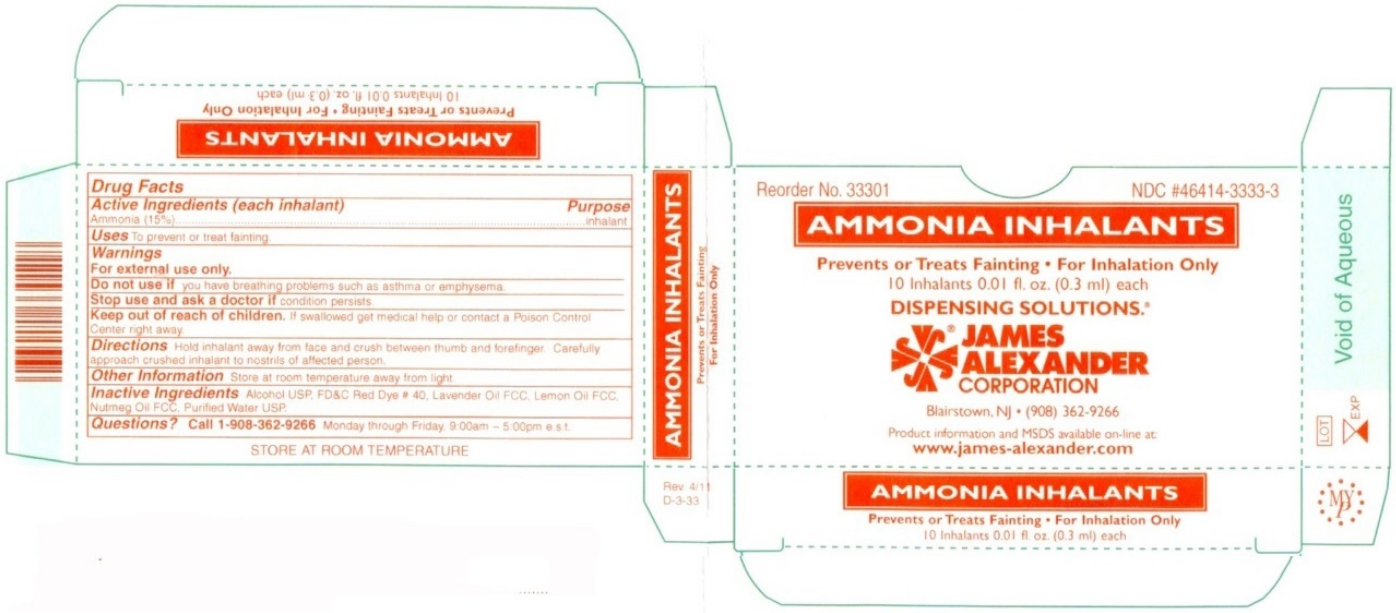
JAMES ALEXANDER CORPORATION
Blairstown, NJ• (908) 362-9266
Product information and MSDS available on-line at:
www.james-alexander.com

Void of Aqueous

Packaging



Packaging



AMMONIA INHALANTS

ammonia inhalants inhalant

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	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**
ALCOHOL (UNII: 3K9958V90M)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:46414-3333-2	100 in 1 CARTON	02/14/1976	
1	NDC:46414-3333-3	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		02/14/1976	

Labeler - James Alexander Corporation (040756421)

Registrant - James Alexander Corporation (040756421)

Establishment

Name	Address	ID/FEI	Business Operations
James Alexander Corporation		040756421	manufacture(46414-3333)

Revised: 1/2019 James Alexander Corporation