GO TIME - ammonia inhalant
Mountain Top Labs, LLC

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.

Drug Facts

Active ingredient (each inhalant)

Ammonia 4.5%

Purpose

Reflex Stimulant

Uses

To arouse consciousness and restore mental alertness.

Warnings

For external use only

Do not use if you have breathing problems such as asthma or emphysema.

When using this product avoid contact with the eyes.

Stop use and ask a doctor if you experience any adverse effects.

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 nostril until desired effect is achieved.

Other information

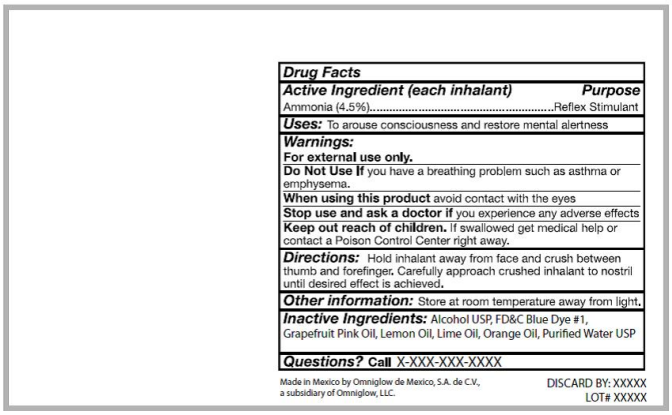
Store at room temperature away from light.

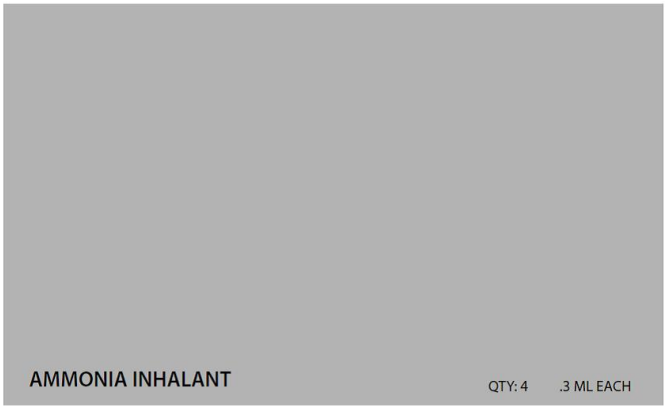
Inactive ingredients

Alcohol USP, FD and C Blue Dye #1, Grapefruit Pink Oil, Lemon Oil, Lime Oil, Orange Oil, Purified Water USP

Questions?

Call 1-801-448-6809





NDC 53063-1114-1

GO TIME

ammonia inhalant

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:53063-1114
Route of Administration	RESPIRATORY (INHALATION)		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
AMMONIA (UNII: 5138Q19F1X) (AMMONIA - UNII:5138Q19F1X)	AMMONIA	0.013 mL in 0.3 mL

Inactive Ingredients	
Ingredient Name	**Strength**
ALCOHOL (UNII: 3K9958V90M)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
GRAPEFRUIT OIL (UNII: YR377U58W9)	
LEMON OIL (UNII: I9GRO824LL)	
LIME OIL (UNII: UZH29XGA8G)	
ORANGE OIL (UNII: AKN3KSD11B)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:53063-1114-2	4 in 1 BOX		
1	NDC:53063-1114-1	0.3 mL in 1 AMPULE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		08/13/2012	

Labeler - Mountain Top Labs, LLC (078408468)

Revised: 9/2012 Mountain Top Labs, LLC