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.

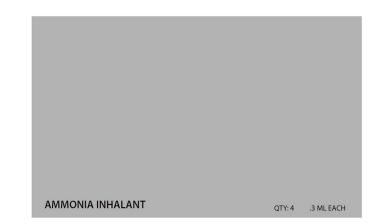
Inactive ingredients

Alcohol USP, Cinnamon Cassia Oil, Eucalyptus Oil, FD and C Blue Dye #1, Purified Water USP

Questions?

Call 1-801-448-6809

Dru	rug F	Fact	ts									
Act	ctive	e Ing	gred	die	nt (eac	:h ii	nhala	nt)		Purp	ose
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Do Not Use If you have a breathing problem such as asthma or												
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NDC 53063-1113-1

GO TIME

ammonia inhalant

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:53063-1113
Route of Administration	RESPIRATORY (INHALATION)		

Active Ingredient/Active Moiety					
Ingredient Name	Basis of Strength	Strength			
AMMO NIA (UNII: 5138 Q 19 F1X) (AMMO NIA - UNII: 5138 Q 19 F1X)	AMMO NIA	$0.013\ mL$ in $0.3\ mL$			

Inactive Ingredients	
Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
CINNAMON OIL (UNII: E5GY4I6 YCZ)	
EUCALYPTUS OIL (UNII: 2R04ONI662)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
WATER (UNII: 059QF0KO0R)	

P	Packaging							
#	Item Code	Package Description	Marketing Start Date	Marketing End Date				
1	NDC:53063-1113-2	4 in 1 BOX						
1	NDC:53063-1113-1	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