

**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](#).*

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**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

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<b>Questions? Call</b> X-XXX-XXX-XXXX	

Made in Mexico by Omniciglow de Mexico, S.A. de C.V., a subsidiary of Omniciglow, LLC. DISCARD BY: XXXXX LOT# XXXXX

AMMONIA INHALANT

QTY: 4 .3 ML EACH

NDC 53063-1113-1

## GO TIME

ammonia inhalant

### Product Information

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

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
AMMONIA (UNII: 5138 Q19 F1X) (AMMONIA - UNII:5138 Q19 F1X)	AMMONIA	0.013 mL in 0.3 mL

### Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958 V90M)	
CINNAMON OIL (UNII: E5GY4I6 YCZ)	
EUCALYPTUS OIL (UNII: 2R04ONI662)	
FD&C BLUE NO. 1 (UNII: HBR47K3TBD)	
WATER (UNII: 059 QF0 KO0R)	

### 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	

