**AMMONIA INHALANTS- ammonia inhalants inhalant**
**A-S Medication Solutions**

*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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**Ammonia Inhalants**

**Active ingredient(s)**

Ammonia (15%)

**Purpose**

Inhalant

**Use(s)**

To prevent or treat fainting

**Warnings**

Keep away from the Eyes.

**For external use only**

**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**

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.

**Storage**

Store at 20ºC to 25ºC (68ºF to 77ºF)

## Inactive ingredients

Alcohol USP, FDC red dye 40, lavender oil fcc, lemon oil fcc, nutmeg oil fcc, purified water usp

## Questions

Questions? Call 1-866-390-4411 Mon - Fri 9:00 AM - 5:00 PM

## Ammonia Inhalants


# AMMONIA INHALANTS

ammonia inhalants inhalant

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN PRESCRIPTION DRUG | **Item Code (Source)** | NDC:50090-0342(NDC:39822-9900) |
| **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) | |
| **LAVENDER OIL** (UNII: ZBP1YXW0H8) | |

| | |
|---|---|
| **WATER** (UNII: 059QF0KO0R) | |
| **NUTMEG OIL** (UNII: Z1CLM48948) | |
| **LEMON OIL** (UNII: I9GRO824LL) | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:50090-0342-0 | 12 in 1 CARTON | 11/28/2014 | |
| **1** | | .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 - A-S Medication Solutions (830016429)

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| A-S Medication Solutions | | 830016429 | RELABEL(50090-0342) |

Revised: 4/2021 A-S Medication Solutions