

SHOCK-X AMMONIA INHALANT- ammonia inhalant inhalant Platinum Trident Enterprise

Active Ingredients

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 the reach of children.

If swallowed, seek medical help or contact a Poison Control Center right away.

If pregnant or nursing, seek the advice of a doctor before using this product.

Directions

Hold inhalant away from face and slowly approach inhalant to nostrils and inhale. Avoid contact with eyes and skin.

Other Information

Store at room temperature away from light.

Inactive Ingredients

Alcohol, Cotton, Peppermint Oil, Water.

Questions

Contact us at support@shock-x.com

Packaging

**WARNING**

For External Use Only

This product has not been approved by the FDA. Use at your own risk



SHOCK-X™

Questions?
support@shock-x.com



1 98715 81682 2

Drug Facts

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Ammonia 15%

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**SHOCK-X**
FEEL THE SHOCK
SMELLING SALTS


PEPPERMINT
SCENTED

NET WT 1.41 OZ [40 G]

Drug Facts (continued)

Directions Hold inhalant away from face and slowly approach inhalant to nostrils and inhale.

Avoid contact with eyes and skin.

Other Information Store at room temperature away from light.

Inactive Ingredients Ammonium Carbonate, Peppermint Oil, Sea Salt.

SHOCK-X AMMONIA INHALANT

ammonia inhalant inhalant

Product Information				
Product Type		HUMAN OTC DRUG	Item Code (Source)	
Route of Administration		NASAL	NDC:85570-1111	
Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strength	Strength
AMMONIA (UNII: 5138Q19F1X) (AMMONIA - UNII:5138Q19F1X)			AMMONIA	15 g in 100 g
Inactive Ingredients				
Ingredient Name				Strength
SEA SALT (UNII: 87GE52P74G)				
PEPPERMINT OIL (UNII: AV092KU4JH)				
AMMONIUM CARBONATE (UNII: NJ5VT0FKLJ)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:85570-1111-2	1 in 1 BOX	01/01/2025	
1	NDC:85570-1111-1	40 g in 1 BOTTLE; Type 0: Not a Combination Product		
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date

OTC Monograph Drug	505G(a)(3)	01/01/2025	
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Labeler - Platinum Trident Enterprise (119491265)

Revised: 5/2025

Platinum Trident Enterprise