SYBORG SALTS SMELLING SALT- ammonium carbonate and sodium carbonate granule Direct Outlet Supplier 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.

SYBORG SALTS SMELLING SALTS

Active Ingredients

Ammonium Carbonate	15%
Sodium Carbonate	.10%

Purpose

The purpose of Syborg Smelling Salts is a boost of arousal in your respiratory system which causes the muscles that control breathing to be fast acting and put you into a fight or flight state whether its for training, working or when it matters the most The purpose of Syborg Smelling Salts is a boost of arousal in your respiratory system which causes the muscles that control breathing to be fast-acting and put you into a fight or flight state whether its for training, working or when it matters the most Game Time.

Usage

The combination of Ammonium Carbonate + Sodium Carbonate added with some water and well shaken will release ammonia gas that irritates the nasal pathways that provides instant fight or flight with boosts in strength, endurance and focus.

Warning and Precautions

Keep away from fire or flame Always keep bottle 6 inches from nose.

Keep away from children.

Do not use if allergic to any ingredients or pregnant.

Directions

Product is already pre-activated. Just shake the bottle and sniff!

Inactive Ingredient

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SYBORG SALTS SMELLING SALT

ammonium carbonate and sodium carbonate granule

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:85813-001
Route of Administration	RESPIRATORY (INHALATION)		

Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
AMMONIUM CARBONATE (UNII: NJ5VT0FKLJ) (AMMOUNII:54S68520I4)	NIUM CATION -	AMMONIUM CATION	15 g in 100 g
SODIUM CARBONATE (UNII: 45P3261C7T) (CARBON	ATE ION - UNII:7UJQ5OPE7D)	SODIUM CARBONATE	10 g in 100 g

Inactive Ingredients			
Ingredient Name	Strength		
ALCOHOL (UNII: 3K9958V90M)			

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:85813-001- 01	47 g in 1 BOTTLE; Type 0: Not a Combination Product	09/03/2025	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		09/03/2025	

Labeler - Direct Outlet Supplier LLC (133619845)

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