

**NAUZENE- sodium citrate tablet, chewable**  
**Alva-Amco Pharmacal Companies, Inc.**

*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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**Nauzene Chewables**

**Active ingredient (in each chewable tablet)**

Sodium citrate dihydrate 230 mg

**Purpose**

Upset stomach reliever/antacid

**Uses**

For the relief of nausea associated with upset/sour stomach, including that due to overindulgence in food and drink.

**Warnings**

**Do not use** if you have Hereditary Fructose Intolerance (HFI). This product contains fructose.

**Ask a doctor before use if you**

- have diabetes because this product contains sugar
- are on a sodium-restricted diet
- have phenylketonuria because each chewable tablet contains 4.5 mg phenylalanine.

**Ask a doctor or pharmacist before use if you are** taking any other medications. This product may interact with certain prescription drugs.

**When using this product,** do not take more than 24 tablets in a 24-hour period.

**Stop use and ask a doctor if** nausea lasts more than two weeks or recurs frequently.

**If pregnant or breastfeeding,** ask a health professional before use.

**Keep out of reach of children.** In case of overdose, get medical help or contact a Poison Control Center right away.

**Directions**

- Adults: 2 - 4 tablets.
- Children: Consult a doctor for appropriate dosage.
- Chew tablets completely.
- Dosage may be repeated after 15 minutes, not to exceed 24 tablets in a 24-hour period unless advised by a doctor.
- Read all package directions and warning before use and use only as directed.
- Nauzene Chewables are intended for use by normally healthy persons only.
- Persons under 18 years of age should use only as directed by a doctor.

### **Other information**

- Sodium content: 60 mg/tablet
- Store at room temperature.
- **\*\*Contents sealed:** Each round pink Nauzene chewable tablet bears the identifying mark "ALVA" and is sealed in a clear plastic blister with a foil backing. Do not use if seal appears broken or if product contents do not match product description. Slight red speckling may occur over time.
- **Note:** Nauzene is not intended as a substitute for a balanced nutritional diet or as an electrolyte replenishment.
- You may report serious side effects to the phone number provided under *Questions?* below.

### **Inactive ingredients**

Aspartame, bitter masking salt, dextrose, FDC Red No. 40 Lake, flavors, food starch-modified, fructose, hypromellose, magnesium stearate, maltodextrin, povidone, silicon dioxide, stearic acid, sucrose.

**Questions? 1-800-792-2582**



<b>NAUZENE</b>			
sodium citrate tablet, chewable			
<b>Product Information</b>			
<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:52389-242
<b>Route of Administration</b>	ORAL		
<b>Active Ingredient/Active Moiety</b>			
<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>	
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K) (ANHYDROUS CITRIC ACID - UNII:XF417D3PSL)	ANHYDROUS CITRIC ACID	230 mg	

## Inactive Ingredients

Ingredient Name	Strength
<b>ASPARTAME</b> (UNII: Z0H242BBR1)	
<b>FD&amp;C RED NO. 40</b> (UNII: WZB9127XOA)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	
<b>MALTODEXTRIN</b> (UNII: 7CVR7L4A2D)	
<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)	
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	
<b>DEXTROSE</b> (UNII: IY9XDZ35W2)	
<b>FRUCTOSE</b> (UNII: 6YSS42VSEV)	
<b>MODIFIED CORN STARCH (1-OCTENYL SUCCINIC ANHYDRIDE)</b> (UNII: 461P5CJN6T)	
<b>HYPROMELLOSES</b> (UNII: 3NXW29V3WO)	
<b>POVIDONE K30</b> (UNII: U725QWY32X)	
<b>SUCROSE</b> (UNII: C151H8M554)	
<b>CHERRY</b> (UNII: BUC5I9595W)	

## Product Characteristics

<b>Color</b>	pink	<b>Score</b>	no score
<b>Shape</b>	ROUND	<b>Size</b>	16mm
<b>Flavor</b>	CHERRY (Wild Cherry Flavor)	<b>Imprint Code</b>	ALVA
<b>Contains</b>			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52389-242-40	4 in 1 CARTON	07/08/2000	
1		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:52389-242-42	3 in 1 CARTON	07/06/2017	
2		14 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:52389-242-50	5 in 1 CARTON	10/01/2006	
3		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
4	NDC:52389-242-10	1 in 1 CARTON	07/15/2016	
4		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
5	NDC:52389-242-56	4 in 1 CARTON	11/16/2019	
5		14 in 1 BLISTER PACK; Type 0: Not a Combination Product		

## Marketing Information

<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
unapproved drug other		07/08/2000	

**Labeler** - Alva-Amco Pharmacal Companies, Inc. (042074856)

Revised: 10/2021

Alva-Amco Pharmacal Companies, Inc.