#### ALKA-SELTZER GOLD- alka-seltzer gold tablet, effervescent Bayer HealthCare LLC.

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Alka-Seltzer Gold

#### **Drug Facts**

Drug Facts

# Active Ingredients

Anhydrous citric acid 1000 mg	Antacid
Potassium bicarbonate 344 mg	Antacid.
Sodium bicarbonate (heat-treated) 1050 mg	Antacid

## Purpose

Antacid

**Uses** for the relief of

 $\bullet$  heartburn  $\bullet$  acid indigestion  $\bullet$  sour stomach

# Do Not Use

**Do not use** if you have ever had an allergic reaction to this product or any of its ingredients

## Ask a doctor before use if you have

- kidney disease
- a potassium or sodium-restricted diet

## Ask a doctor or pharmacist before use if you are

Ask a doctor or pharmacist before use if you are presently taking a prescription drug. Antacids may interact with certain prescription drugs.

## When using this product

When using this product do not exceed recommended dosage

## Stop use and ask a doctor if

Stop use and ask a doctor if you have taken the maximum dose for 2 weeks

If pregnant or breast-feeding, ask a health professional before use.

# Keep out of reach of children

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#### Directions

• fully dissolve tablets in 4 ounces of water before taking

	2 tablets every 4 hours as needed, or asdirected by a doctor	do not exceed8 tablets in 24 hours
adults 60 years and over	2 tablets every 4 hours as needed, or as directed by a doctor	do not exceed 6 tablets in 24 hours
children under 12 years	1 tablet every 4 hours as needed, or as directed by a doctor	do not exceed 4 tablets in 24 hours

# Other information

- each tablet contains: potassium 135 mg
- each tablet contains: sodium 309 mg
- store at room temperature. Avoid excessive heat.
- this product does not contain aspirin
- Alka-Seltzer Gold in water contains principally the

antacids sodium citrate and potassium citrate

## Inactive ingredients

Inactive ingredients magnesium stearate, mannitol

## **Questions or comments**

#### Questions or comments?1-800-986-0369 (Mon - Fri 9AM - 5PM EST)

**Reye's syndrome:** Children and teenagers who have or are recovering from chicken pox or flu-like symptoms should not use this product. When using this product, if changes in behavior with nausea and vomiting occur, consult a doctor because these symptoms could be an early sign of Reye's syndrome, a rare but serious illness.

Allergy alert: Aspirin may cause a severe allergic reaction which may include:

 $\cdot$  hives  $\cdot$  facial swelling  $\cdot$  asthma (wheezing)  $\cdot$  shock

**Stomach bleeding warning:** This product contains an NSAID, which may cause severe stomach bleeding. The chance is

higher if you

- are age 60 or older
- have had stomach ulcers or bleeding problems
- take a blood thinning (anticoagulant) or steroid drug
- take other drugs containing prescription or nonprescription

NSAIDs (aspirin, ibuprofen, naproxen, or others)

have 3 or more alcoholic drinks every day while using this product

• take more or for a longer time than directed





ALKA-SELTZER GOLD					
alka-seltzer gold tablet, efferv	escent				
Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source)		NDC:0280-4100	
Route of Administration	ORAL				
Active Ingredient/Active	Μοιετγ				
Ingredient Name			Basis of Strength		Strength
<b>CITRIC ACID MONOHYDRATE</b> (UNII: 2968PHW8QP) (ANHYDROUS CITRIC ACID - UNII:XF417D3PSL)			ANHYDROUS CITRIC ACID		1000 mg
POTASSIUM BICARBONATE (UNII UNII:295053K152)	: HM5Z15LEBN) (POTASSIU	M CATION -	POTASSIL	JM CATION	344 mg
			SODIUM BICARBON	NATE	1050 mg
Inactive Ingredients					
Ingredient Name				Strength	

MAGNESIUM STEARATE (UNII: 70097M6I30)

MANNITOL (UNII: 30WL53L36A)						
<b>Product Chara</b>	Product Characteristics					
Color		white	Score		no score	
Shape		ROUND	Size		25mm	
Flavor			Imprint Code			
Contains						
Packaging						
# Item Code		Package Description		ľ	Aarketing Start Date	Marketing End Date
<b>1</b> NDC:0280-4100- 63	18 in 1 C	CARTON		11/19/2014		
1	2 in 1 PO Product	n 1 POUCH; Type 0: Not a Combination				
Marketing Information						
Marketing Category	Арр	Application Number or Monograpl Citation		h Marketing Star Date		Marketing End Date
OTC Monograph Drug	g M001				11/19/2014	

Labeler - Bayer HealthCare LLC. (112117283)

Revised: 12/2024

Bayer HealthCare LLC.