

**ALKA-SELTZER PLUS MAXIMUM STRENGTH SINUS CONGESTION AND PAIN  
POWERFAST FIZZ- chlorpheniramine maleate, dextromethorphan  
hydrobromide, acetaminophen, phenylephrine hydrochloride tablet,  
effervescent  
Bayer HealthCare LLC.**

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**ASP Maximum Strength Sinus Congestion and Pain PowerFast Fizz UI1614734**

***Drug Facts***

***Active ingredients (in each tablet) Purposes***

Acetaminophen 325 mg.....Pain reliever/fever reducer  
Chlorpheniramine maleate 2 mg.....Antihistamine  
Dextromethorphan hydrobromide 10 mg.....Cough suppressant  
Phenylephrine hydrochloride 5 mg.....Nasal decongestant

**Uses**

temporarily relieves these symptoms due to a cold or flu:

- minor aches and pains
- headache
- cough
- sore throat
- runny nose
- sneezing
- nasal congestion · sinus

temporarily reduces fever

**Liver warning:** This product contains acetaminophen. Severe liver damage may occur if you take

- more than 4,000 mg of acetaminophen in 24 hours
- with other drugs containing acetaminophen
- 3 or more alcoholic drinks every day while using this product

**Allergy alert:** Acetaminophen may cause severe skin or severe allergic reactions. Symptoms may include:

- skin reddening · blisters · rash · hives
- facial swelling · asthma (wheezing) · shock

If a skin or general allergic reaction occurs, stop use and seek medical help right away.

**Sore throat warning:** If sore throat is severe, persists for more than 2 days, is accompanied or followed by fever, headache, rash, nausea,

or vomiting, consult a doctor promptly.

**Do not use to sedate children.**

**Do not use**

- with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.
- if you are now taking a prescription monoamine oxidase inhibitor (MAOI) (certain drugs for depression, psychiatric, or emotional

conditions, or Parkinson's disease), or for 2 weeks after stopping the MAOI drug. If you do not know if your prescription drug

contains an MAOI, ask a doctor or pharmacist before taking this product

- if you have ever had an allergic reaction to this product or any of its ingredients
- in children under 12 years of age

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

- taking the blood thinning drug warfarin
- taking sedatives or tranquilizers

**When using this product**

- **do not exceed recommended dosage**
- may cause marked drowsiness
- avoid alcoholic drinks
- alcohol, sedatives, and tranquilizers may increase drowsiness
- be careful when driving a motor vehicle or operating machinery
- excitability may occur, especially in children

**Stop use and ask a doctor if**

- pain, cough, or nasal congestion gets worse or lasts more than 7 days
- fever gets worse or lasts more than 3 days
- redness or swelling is present
- new symptoms occur
- cough comes back or occurs with rash or headache that lasts. These
- could be signs of a serious condition.
- nervousness, dizziness, or sleeplessness occurs

**If pregnant or breast-feeding,** 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. Quick medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

**Directions**

- do not take more than the recommended dose
- adults and children 12 years and over: take 2 tablets fully dissolved in 4 oz of water every 4 hours. Do not exceed 10 tablets in 24 hours or as directed by a doctor.
- children under 12 years: do not use

**Other information**

**each tablet contains:** potassium 80 mg; sodium 356 mg

store at room temperature. Avoid excessive heat.

**Inactive ingredients** anhydrous citric acid, calcium silicate, dimethicone, flavors, magnesium stearate, maltodextrin, mannitol, potassium bicarbonate, povidone, sodium bicarbonate, sucralose

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



**Alka-Seltzer Plus®**

**MAXIMUM STRENGTH**

**Sinus**

**Congestion**

**& Pain**

**PowerFast Fizz™**

**Berry**

**New!**

Actaminophen/Pain Reliever-Fever Reducer

Chlorpheniramine Maleate/Antihistamine

Dextromethorphan HBr/Cough Suppressant

Phenylephrine Hydrochloride/Nasal Decongestant

- Sinus Congestion & Pressure
- Nasal Congestion
- Headache, Body Ache, Sore Throat
- Runny Nose, Sneezing
- Cough

24 EFFERVESCENT TABLETS

**ALKA-SELTZER PLUS MAXIMUM STRENGTH SINUS CONGESTION AND PAIN POWERFAST FIZZ**

chlorpheniramine maleate, dextromethorphan hydrobromide, acetaminophen, phenylephrine hydrochloride tablet, effervescent

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0280-0055
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
CHLORPHENIRAMINE MALEATE (UNII: V1Q0O9OJ9Z) (CHLORPHENIRAMINE - UNII:3U6IO1965U)	CHLORPHENIRAMINE MALEATE	2 mg
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg

Inactive Ingredients	
Ingredient Name	Strength
CALCIUM SILICATE (UNII: S4255P4G5M)	
MANNITOL (UNII: 3OWL53L36A)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POTASSIUM BICARBONATE (UNII: HM5Z15LEBN)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
DIMETHICONE (UNII: 92RU3N3Y1O)	

Product Characteristics			
Color	white	Score	no score
Shape	ROUND	Size	25mm
Flavor	BERRY (Raspberry and Mixed Berry)	Imprint Code	SINUS
Contains			

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0280-0055-01	10 in 1 CARTON	06/15/2021	
1		2 in 1 POUCH; Type 0: Not a Combination Product		
2	NDC:0280-0055-02	12 in 1 CARTON	03/08/2023	
2		2 in 1 POUCH; Type 0: Not a Combination Product		

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	06/15/2021	

**Labeler** - Bayer HealthCare LLC. (112117283)

Revised: 12/2024

Bayer HealthCare LLC.