

ALKA-SELTZER PLUS SEVERE COLD NIGHT POWERFAST FIZZ- aspirin, chlorpheniramine maleate, dextromethorphan hydrobromide, phenylephrine bitartrate tablet, effervescent
Bayer HealthCare LLC.

Alka-Selzer Plus Severe Cold PowerFast Fizz Night effervescent tablets UI 1614459

Drug Facts

Active ingredients (in each tablet) Purposes

Aspirin 325 mg (NSAID)*.....Pain reliever/fever reducer
Chlorpheniramine maleate 2 mg.....Antihistamine
Dextromethorphan hydrobromide 10 mg.....Cough suppressant
Phenylephrine bitartrate 7.8 mg.....Nasal decongestant

*nonsteroidal anti-inflammatory drug

Uses

- temporarily relieves these symptoms due to a cold:
 - minor aches and pains
 - headache
 - runny nose
 - sinus congestion and pressure
 - cough
 - sneezing
 - sore throat
 - nasal congestion
- temporarily reduces fever

Warnings

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:

- hives
- facial swelling
- asthma (wheezing)
- 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

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

- if you are allergic to aspirin or any other pain reliever/fever reducer
- 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 before use if

- stomach bleeding warning applies to you
 - you have a history of stomach problems, such as heartburn
 - you have high blood pressure, heart disease, liver cirrhosis, or kidney disease
 - you are taking a diuretic
 - you have
 - asthma
 - diabetes
 - thyroid disease
 - glaucoma
 - cough that occurs with excessive phlegm (mucus)
 - a breathing problem such as emphysema or chronic bronchitis
 - difficulty in urination due to enlargement of the prostate gland
 - persistent or chronic cough such as occurs with smoking, asthma, or emphysema
 - a sodium-restricted diet

Ask a doctor or pharmacist before use if you are

- taking a prescription drug for
 - gout
 - diabetes
 - arthritis
- 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

- an allergic reaction occurs. Seek medical help right away.
- you experience any of the following signs of stomach bleeding
 - feel faint
 - vomit blood
 - have bloody or black stools
 - have stomach pain that does not get better
- 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
- ringing in the ears or a loss of hearing occurs
- 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.

It is especially important not to use aspirin at 20 weeks or later in pregnancy unless definitely directed to do so by a doctor because it may cause problems in the unborn child or complications during delivery.

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

Directions

- adults and children 12 years and over: take 2 tablets fully dissolved in 4 oz of water every 4 hours. Do not exceed 8 tablets in 24 hours or as directed by a doctor.
- children under 12 years of age: 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, mannitol, potassium bicarbonate, povidone, sodium bicarbonate, sucralose

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

Carton 24 count



RESTRICTED

Alka-Seltzer Plus®

SEVERE

Cold

LEMON

POWERFAST FIZZ™ SEE NEW DOSING DIRECTIONS

NIGHT

Aspirin (**NSAID**)/Pain Reliever-Fever Reducer

Chlorpheniramine Maleate / Antihistmine

Dextromethorphan HBr / Cough Suppressant

Phenylephrine Bitartrate / Nasal Decongestant

- Nasal Congestion
- Headache + Body Ache
- Cough
- Runny Nose
- Sore Throat

24 EFFERVESCENT TABLETS

ALKA-SELTZER PLUS SEVERE COLD NIGHT POWERFAST FIZZ

aspirin, chlorpheniramine maleate, dextromethorphan hydrobromide, phenylephrine bitartrate tablet, effervescent

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0280-0121
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ASPIRIN (UNII: R16CO5Y76E) (ASPIRIN - UNII:R16CO5Y76E)	ASPIRIN	325 mg
CHLORPHENIRAMINE MALEATE (UNII: V1Q0O9OJ9Z) (CHLORPHENIRAMINE - UNII:3U6IO1965U)	CHLORPHENIRAMINE MALEATE	2 mg
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg
PHENYLEPHRINE BITARTRATE (UNII: 27O3Q5ML57) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE BITARTRATE	7.8 mg

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
CALCIUM SILICATE (UNII: S4255P4G5M)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
MANNITOL (UNII: 3OWL53L36A)	
POTASSIUM BICARBONATE (UNII: HM5Z15LEBN)	
POVIDONE (UNII: FZ989GH94E)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	

Product Characteristics

Color	white	Score	no score
Shape	ROUND	Size	25mm
Flavor	LEMON	Imprint Code	ASP;NT
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0280-0121-01	12 in 1 CARTON	05/24/2023	
1		2 in 1 POUCH; Type 0: Not a Combination Product		
2	NDC:0280-0121-	10 in 1 CARTON	05/24/2023	

2	02	10 III 1 CARTON	05/24/2023	
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		05/24/2023	

Labeler - Bayer HealthCare LLC. (112117283)

Revised: 1/2025

Bayer HealthCare LLC.