ALKA-SELTZER PLUS COLD AND FLU- chlorpheniramine maleate, acetaminophen, phenylephrine hydrochloride, dextromethorphan hydrobromide tablet tablet, effervescent tablet, effervescent
Lil' Drug Store Products, Inc.

ALKA-SELTZER PLUS COLD AND FLU

Drug Facts

Active ingredients (in each tablet)

Acetaminophen 250 mg
Chlorpheniramine maleate 2 mg
Dextromethorphan hydrobromide 10 mg
Phenylephrine hydrochloride 5 mg

Purpose

Pain reliever/fever reducer
Antihistamine
Cough suppressant
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 and sinus congestion
- temporarily reduces fever

Warnings

Liver warning

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

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

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

Ask a doctor before use if you have

- liver disease
- heart disease
- high blood pressure
- thyroid disease
- diabetes
- glaucoma
- cough 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 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

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 8 tablets in 24 hours or as directed by a doctor.
- children under 12 years: do not use

Other Information

Other information

- each tablet contains: potassium 80 mg; sodium 356 mg
- store at room temperature. Avoid excessive heat.

Inactive Ingredients

Inactive ingredients

anhydrous citric acid, calcium silicate, dimethicone, FD&C red #40, FD&C yellow #6, flavors, magnesium stearate, maltodextrin,
mannitol, potassium bicarbonate, povidone, sodium bicarbonate, sucralose

Questions or Comments?

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

DO NOT USE IF INNER POUCH OR CARTON IS OPEN OR DAMAGED!
PACKAGE NOT CHILD RESISTANT!

25 Count Box

Alka-

Seltzer

Plus ®

SEVERE

Cold

& Flu

CITRUS

POWERFAST FIZZ™

ACETAMINOPHEN/Pain Reliever-Fever Reducer

Chlorpheniramine Maleate/Antihistamine

Dextromethorphan HBr/Cough Suppressant

Phenylephrine Hydrochloride/Nasal Decongestan

- Fever &
- Body Ache
- Cough
- Nasal Congestion
- Runny Nose
- Sore Throat

[Bayer cross logo]

2 EFFERVESCENT TABLETS PER POUCH

30 Count Box

Alka-

Seltzer

Plus ®

SEVERE

Cold

& Flu

CITRUS

POWERFAST FIZZ™

ACETAMINOPHEN/Pain Reliever-Fever Reducer

Chlorpheniramine Maleate/Antihistamine

Dextromethorphan HBr/Cough Suppressant

Phenylephrine Hydrochloride/Nasal Decongestant

- Fever &

- Body Ache
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- Sore Throat

[Bayer cross logo]

2 EFFERVESCENT TABLETS PER POUCH

ALKA-SELTZER PLUS COLD AND FLU

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

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:29485-7318
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	250 mg
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg
CHLORPHENIRAMINE MALEATE (UNII: V1Q0O9OJ9Z) (CHLORPHENIRAMINE - UNII:3U6IO1965U)	CHLORPHENIRAMINE MALEATE	2 mg

Inactive Ingredients

Ingredient Name	Strength
SUCRALOSE (UNII: 96K6UQ3ZD4)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
POTASSIUM BICARBONATE (UNII: HM5Z15LEBN)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
MANNITOL (UNII: 3OWL53L36A)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
CALCIUM SILICATE (UNII: S4255P4G5M)	

Product Characteristics

Color	white	**Score**	no score
Shape	ROUND	**Size**	25mm
Flavor	CITRUS	**Imprint Code**	ASP;FLU
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:29485-7318-2	25 in 1 BOX	12/23/2021	12/31/2025
1		2 in 1 PACKET; Type 0: Not a Combination Product		
2	NDC:29485-7318-3	30 in 1 BOX	12/23/2021	12/31/2025
2		2 in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M013	12/23/2021	12/31/2025

Labeler - Lil' Drug Store Products, Inc. (093103646)

Revised: 12/2023 Lil' Drug Store Products, Inc.