ALKA SELTZER PLUS MAXIMUM STRENGTH COUGH CHEST CONGESTION POWERFAST FIZZ- dextromethorphan hydrobromide, guaifenesin tablet, effervescent
Bayer Healthcare LLC.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Alka-Seltzer Plus Severe Cold & Cough PowerFast Fizz Effervescent Tablets

Drug Facts

Active ingredients (in each tablet) Purposes

Dextromethorphan hydrobromide 10 mg.....Cough suppressant

Guaifenesin 200 mg.....Expectorant

Uses

Uses

- temporarily relieves:
- cough due to minor throat and bronchial irritation as may occur with a cold
- the intensity of coughing
- helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus and make coughs more productive

Warnings

Do not use

● 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 you have

● cough that occurs with excessive phlegm (mucus) ● persistent or chronic cough such as occurs with smoking, asthma, chronic bronchitis, or emphysema ● a sodium-restricted diet

Stop use and ask a doctor if cough lasts more than 7 days, comes back, or is accompanied by a fever, rash, or persistent headache. These could be signs of a serious condition.

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

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 12 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 237 mg

● store at room temperature. Avoid excessive heat.

anhydrous citric acid, flavors, magnesium stearate, mannitol, potassium bicarbonate, sodium bicarbonate, sucralose

Questions or comments

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

Carton 24 count

Alka Seltzer Plus Maximum Strength Cough & Chest Congestion

POWERFAST FIZZ™

Dextromethorphan HBr / Cough Suppressant

Guaifenesin/Expectorant

24 EFFERVESCENT TABLETS

ALKA SELTZER PLUS MAXIMUM STRENGTH COUGH CHEST CONGESTION POWERFAST FIZZ

dextromethorphan hydrobromide, guaifenesin tablet, effervescent

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0280-0089
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg

GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	200 mg

Inactive Ingredients

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

Product Characteristics

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0280-0089-01	20 in 1 CARTON	04/05/2022	
1		2 in 1 POUCH; Type 0: Not a Combination Product		
2	NDC:0280-0089-02	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 final	part341	04/05/2022	

Labeler - Bayer Healthcare LLC. (112117283)

Revised: 8/2023 Bayer Healthcare LLC.