ALKA-SELTZER PLUS MAXIMUM STRENGTH DAY AND NIGHT COLD AND FLU POWERMAX GELS- acetaminophen, dextromethorphan hydrobromide, phenylephrine hydrochloride, doxylamine succinate Bayer HealthCare LLC.

Alka-Seltzer Plus® Maximum Strength Day & Night Cold & Flu PowerMax™ Gels - 36 count

Alka-Seltzer Plus

Maximum Strength Day & Night Cold & Flu PowerMax Gels

Do not take these products at the same time.

Alka-Seltzer Plus® Maximum Strength Day Cold & Flu PowerMax™ Gels

Drug Facts

Active ingredients (in each capsule)

Acetaminophen 325 mg

Dextromethorphan hydrobromide 10 mg

Phenylephrine hydrochloride 5 mg

Purposes

Pain reliever/fever reducer

Cough suppressant

Nasal decongestant

Uses

- · temporarily relieves these symptoms due to a cold or flu:
- \cdot minor aches and pains \cdot headache \cdot cough
- \cdot sore throat \cdot nasal and sinus congestion
- · temporarily reduces fever

Warnings

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

- \cdot more than 4,000 mg of acetaminophen in 24 hours
- \cdot with other drugs containing acetaminophen
- \cdot 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:

 \cdot skin reddening \cdot blisters \cdot rash \cdot 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

- 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 before use if you have

- liver disease heart disease high blood pressure
- thyroid disease diabetes
- cough with excessive phlegm (mucus)
- difficulty in urination due to enlargement of the prostate gland
- persistent or chronic cough such as occurs with smoking, asthma, or emphysema

Ask a doctor or pharmacist before use if you are taking the blood thinning drug warfarin

When using this product do not exceed recommended dosage

Stop use and ask a doctor if 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 capsules with water every 4 hours. Do not exceed 10 capsules in 24 hours or as directed by a doctor.
- · children under 12 years: do not use

Other information

Other information

store at room temperature. Avoid temperatures above 25°C (77°F).

Inactive ingredients FD&C yellow #6, ferric oxide, gelatin, glycerin, polyethylene glycol, potassium aluminum silicate, povidone, propylene glycol, purified water, shellac, sodium hydroxide, sorbitol sorbitan solution, titanium dioxide

Questions or comments?

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

Alka-Seltzer Plus® Maximum Strength Night Cold & Flu PowerMax™ Gels

Drug Facts

Active ingredients (in each capsule)

Acetaminophen 325 mg

Dextromethorphan hydrobromide 10 mg

Doxylamine succinate 6.25 mg

Phenylephrine hydrochloride 5 mg

Purposes

Pain reliever/fever reducer
Cough suppressant
Antihistamine
Nasal decongestant

Uses

Uses

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

Warnings

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
- \cdot 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:

- \cdot skin reddening \cdot blisters \cdot rash \cdot 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 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

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 than7 days
- \cdot fever gets worse or lasts more than 3 days
- · redness or swelling is present
- · new symptoms occur
- \cdot cough comes back or occurs with rash or headache that lasts.

These could be signs of a serious condition.

 \cdot 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 capsules with water every 4 hours. Do not exceed 10 capsules in 24 hours or as directed by a doctor.
- · children under 12 years: do not use

Other information

Other information

store at room temperature. Avoid temperatures above 25°C (77°F).

Inactive ingredients D&C yellow No. 10, FD&C blue No. 1, gelatin, glycerin, polyethylene glycol, potassium aluminum silicate, povidone, propylene glycol, purified water, shellac, sodium hydroxide, sorbitol sorbitan solution, titanium dioxide

Questions or comments?

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

36 count carton





ALKA-SELTZER PLUS MAXIMUM STRENGTH DAY AND NIGHT COLD AND FLU POWERMAX GELS

acetaminophen, dextromethorphan hydrobromide, phenylephrine hydrochloride, doxylamine succinate kit

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:0280-0125

l	P	Packaging					
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
	1	NDC:0280-0125- 01	1 in 1 CARTON; Type 0: Not a Combination Product	07/02/2018			

Quant	Quantity of Parts				
Part #	Package Quantity	Total Product Quantity			
Part 1	4 BLISTER PACK	24			
Part 2	2 BLISTER PACK	12			

Part 1 of 2

ALKA-SELTZER PLUS MAXIMUM STRENGTH DAY COLD AND FLU POWERMAX GEL

acetaminophen, dextromethorphan hydrobromide, phenylephrine hydrochloride capsule, liquid filled

Product Information

Route of Administration ORAL

Active Ingredient/Active Moiety Ingredient Name Strength **Basis of Strength PHENYLEPHRINE** PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE -5 mg **HYDROCHLORIDE** UNII:1WS297W6MV) ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D) | ACETAMINOPHEN 325 mg **DEXTROMETHORPHAN HYDROBROMIDE** (UNII: 9D2RTI9KYH) DEXTROMETHORPHAN 10 mg (DEXTROMETHORPHAN - UNII:7355X3ROTS) **HYDROBROMIDE**

Inactive Ingredients	
Ingredient Name	Strength
POVIDONE (UNII: FZ989GH94E)	
SORBITOL (UNII: 506T60A25R)	
GELATIN (UNII: 2G86QN327L)	
FERROSOFERRIC OXIDE (UNII: XM0M87F357)	

GLYCERIN (UNII: PDC6A3C0OX)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
SHELLAC (UNII: 46N107B710)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
SORBITAN (UNII: 6092ICV9RU)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
WATER (UNII: 059QF0KO0R)	

Product Characteristics			
Color	orange	Score	no score
Shape	OVAL	Size	16mm
Flavor		Imprint Code	ASP;DC
Contains			

Packaging					
#	Item Package Description		Marketing Start Date	Marketing End Date	
1		4 in 1 CARTON			
1		6 in 1 BLISTER PACK; 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	07/02/2018		

Part 2 of 2

ALKA-SELTZER PLUS MAXIMUM STRENGTH NIGHT COLD AND FLU POWERMAX GEL

acetaminophen, dextromethorphan hydrobromide, doxylamine succinate, phenylephrine hydrochloride capsule, liquid filled

Product Information Route of Administration ORAL

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)	ACETAMINOPHEN	325 mg		
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg		

DOXYLAMINE SUCCINATE (UNII: V9BI9B5YI2) (DOXYLAMINE - UNII:95QB77JKPL)	DOXYLAMINE SUCCINATE	6.25 mg	
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg	

Inactive Ingredients				
Ingredient Name	Strength			
GLYCERIN (UNII: PDC6A3C0OX)				
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)				
SHELLAC (UNII: 46N107B710)				
SORBITAN (UNII: 6092ICV9RU)				
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)				
SODIUM HYDROXIDE (UNII: 55X04QC32I)				
SORBITOL (UNII: 506T60A25R)				
POVIDONE (UNII: FZ989GH94E)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
WATER (UNII: 059QF0KO0R)				
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)				
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)				
GELATIN (UNII: 2G86QN327L)				

Product Characteristics				
Color	green	Score	no score	
Shape	OVAL	Size	16mm	
Flavor		Imprint Code	ASP;N	
Contains				

Pa	Packaging						
#	ltem Code	Package Description	Marketing Start Date	Marketing End Date			
1		2 in 1 CARTON					
1		6 in 1 BLISTER PACK; 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	07/02/2018		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M012	07/02/2018		

Labeler - Bayer HealthCare LLC. (112117283)

Revised: 8/2024 Bayer HealthCare LLC.