CORICIDIN HBP MAXIMUM STRENGTH MULTI SYMPTOM FLU- acetaminophen, chlorpheniramine maleate, dextromethorphan hydrobromide tablet
Bayer HealthCare LLC.

Coricidin HBP Max Strength Multi-symptom Flu (tablet) new product name Perrigo Old name Coricidin HBP Max Strength (tablet)

Coricidin HBP

Maximum Strength Multi-Symptom Flu

Drug Facts

Active Ingredient

Active ingredients (in each tablet) Purposes

Acetaminophen 325 mg……Pain reliever/fever reducer

Chlorpheniramine maleate 2 mg……Antihistamine

Dextromethorphan hydrobromide 10 mg……Cough suppressant

Uses

Uses

- temporarily relieves these symptoms due to a cold or flu:
- minor aches and pains
- headache
- cough
- runny nose
- sneezing
- sore throat
- 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

- 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

Ask a doctor before use if you have

- liver disease
- trouble urinating due to an enlarged prostate gland
- cough that occurs with excessive phlegm (mucus)
- a breathing problem or persistent or chronic cough as occurs with smoking, asthma, chronic bronchitis, or emphysema

When using this product

- excitability may occur, especially in children
- marked drowsiness may occur
- avoid alcoholic beverages
- alcohol, sedatives and tranquilizers may increase drowsiness
- use caution when driving a motor vehicle or operating machinery

Ask a doctor or pharmacist before use if you are

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

Stop use and ask a doctor if

- pain or cough 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.

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children. Abuse of this product can lead to serious injury.

Overdose warning

Overdose warning: Taking more than the recommended dose may cause liver damage. 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 children even if you do not notice any signs or symptoms.

Directions

do not use more than directed (see overdose warning)

adults and children 12 years and over: take 2 tablets every 4 hours, while symptoms persist. Do not take more than 10 tablets in 24 hours or as directed by a doctor.

children under 12 years of age: ask a doctor

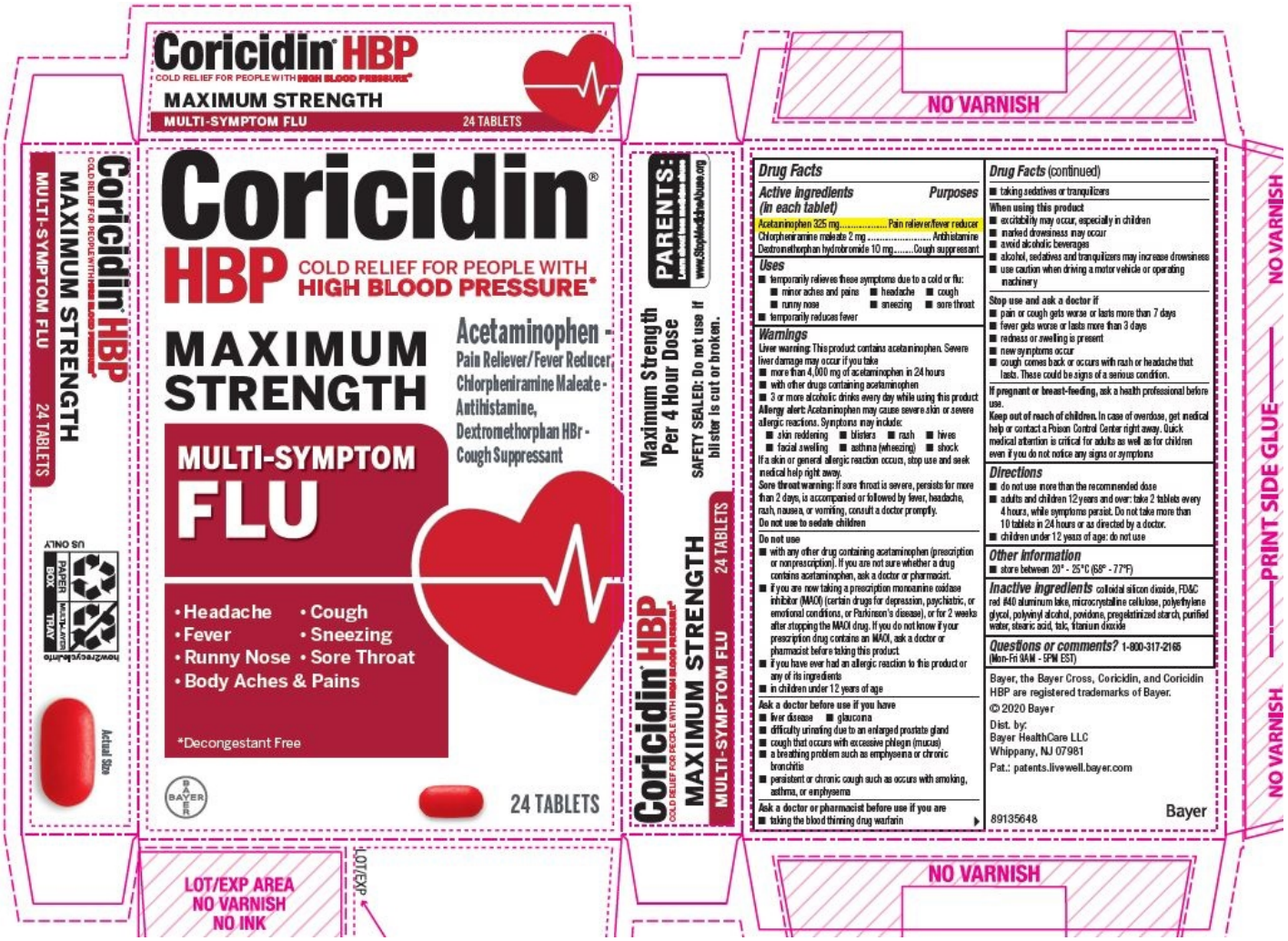
Other information

store at room temperature. Avoid excessive heat.

Inactive ingredients FD&C red #40 aluminum lake, lecithin, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, povidone, pregelatinized starch, stearic acid, talc, titanium dioxide

Questions or comments? 1-800-317-2165 (Mon-Fri 9AM - 5PM EST)

Carton



Coricidin®

HBP COLD RELIEF FOR PEOPLE WITH

HIGH BLOOD PRESSURE‡

MAXIMUM STRENGTH

MULTI-SYMPTOM

FLU

Relieves

- Body Aches & Pains
- Headache
- Fever
- Cough
- Runny Nose & Sneezing
- Sore Throat

‡Decongestant Free

24 tablets

Acetaminophen - Pain Reliever/Fever Reducer

Chlorpheniramine Maleate - Antihistamine

Dextromethorphan HBr - Cough Supressant

LEARN MORE

American Heart Association

Information Enclosed

CORICIDIN HBP MAXIMUM STRENGTH MULTI SYMPTOM FLU

acetaminophen, chlorpheniramine maleate, dextromethorphan hydrobromide tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11523-0038
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
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg
CHLORPHENIRAMINE MALEATE (UNII: V1Q0O9OJ9Z) (CHLORPHENIRAMINE - UNII:3U6IO1965U)	CHLORPHENIRAMINE MALEATE	2 mg

Inactive Ingredients	
Ingredient Name	**Strength**
TALC (UNII: 7SEV7J4R1U)	
STARCH, CORN (UNII: O8232NY3SJ)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
POVIDONE (UNII: FZ989GH94E)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

Product Characteristics			
Color	red	**Score**	no score
Shape	OVAL	**Size**	17mm
Flavor		**Imprint Code**	c;flu
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:11523-0038-1	1 in 1 CARTON	09/04/2020	
1		10 in 1 BLISTER PACK; Type 0: Not a Combination		

1		Product		
2	NDC:11523-0038-2	2 in 1 CARTON	09/04/2020	
2		12 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	07/01/2020	

Labeler - Bayer HealthCare LLC. (112117283)

Revised: 12/2024 Bayer HealthCare LLC.