

SOUND BODY SEVERE COLD AND FLU- acetaminophen, dextromethorphan hbr, phenylephrine hcl powder, for solution

Big Lots Stores, Inc.

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.

Big Lots, Inc Severe Cold & Flu Drug Facts

Active ingredients (in each packet)

Acetaminophen 500 mg

Dextromethorphan HBr 20 mg

Phenylephrine HCl 10 mg

Purposes

Pain reliever/fever reducer

Cough suppressant

Nasal decongestant

Uses

- temporarily relieves these symptoms due to a cold:
- minor aches and pains
- minor sore throat pain
- headache
- nasal and sinus congestion
- cough due to minor throat and bronchial irritation
- 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
- 3 or more alcoholic drinks every day while using this product

Allergy alert: Acetaminophen may cause severe skin reactions. Symptoms may include:

- skin reddening
- blisters
- rash

If a skin 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

- in a child under 12 years of age
- if you have ever had an allergic reaction to this product or any of its ingredients
- 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.

Ask a doctor before use if you have

- liver disease
- heart disease
- high blood pressure
- thyroid disease
- diabetes
- trouble urinating due to an enlarged prostate gland
- cough that occurs with too much phlegm (mucus)
- cough that lasts or is chronic 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

When using this product

do not exceed recommended dosage

Stop use and ask a doctor if

- nervousness, dizziness, or sleeplessness occurs
- fever gets worse or lasts more than 3 days
- redness or swelling is present
- new symptoms occur
- pain, cough or nasal congestion gets worse or lasts more than 7 days
- cough comes back or occurs with rash or headache that lasts. These could be signs of serious condition.

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

Overdose warning: In case of overdose, get medical help or contact a Poison Control Center right away (1-800-222-1222). Prompt medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

Directions

- do not use more than directed (see overdose warning)
- take every 4 hours, while symptoms persist. Do not take more than 6 packets in 24 hours unless directed by a doctor.

Age	Dose
adults and children 12 years of age and over	one packet
children under 12 years of age	do not use

- dissolve contents of one packet into 8 oz. hot water: sip while hot. Consume entire drink within 10-15 minutes.
- if using a microwave, add contents of one packet to 8 oz. of cool water: stir briskly before and after heating. Do not overheat.

Other information

- each packet contains: potassium 10 mg and sodium 25 mg
- phenylketonurics: contains phenylalanine 22 mg per packet
- store at 20-25°C (68-77°F). Protect product from heat and moisture.

Inactive ingredients

acesulfame potassium, anhydrous citric acid, aspartame, colloidal silicon dioxide, D&C yellow #10, FD&C blue #1, FD&C red #40, flavors, maltodextrin, pregelatinized starch, sodium citrate, sucrose, tribasic calcium phosphate

Questions or comments?

1-800-719-9260

Package/Label Principal Display Panel

Compare to the Active Ingredients in Theraflu® Multi-Symptom Severe Cold

Nasal & Sinus Congestion

Cough

Body Ache

Sore Throat Pain

Headache

Fever

DAYTIME MULTI-SYMPTOM

Severe Cold & Flu

Pain Reliever / Fever Reducer

Acetaminophen

Cough Suppressant

Dextromethorphan HBr

Nasal Decongestant

Phenylephrine HCl

Green Tea & Honey Lemon Flavors

6 PACKETS



Compare to the Active
Ingredients in Theraflu®
Multi-Symptom Severe Cold*

- Nasal & Sinus Congestion • Cough
- Body Ache • Sore Throat Pain
- Headache • Fever

DAYTIME
MULTI-SYMP TOM
Severe
Cold & Flu

Pain Reliever / Fever Reducer
Acetaminophen

Cough Suppressant
Dextromethorphan HBr

Nasal Decongestant
Phenylephrine HCl

Green Tea & Honey
Lemon Flavors

6 PACKETS



DAYTIME
MULTI-SYMP TOM
Severe
Cold & Flu

Pain Reliever / Fever Reducer
Acetaminophen

Cough Suppressant
Dextromethorphan HBr

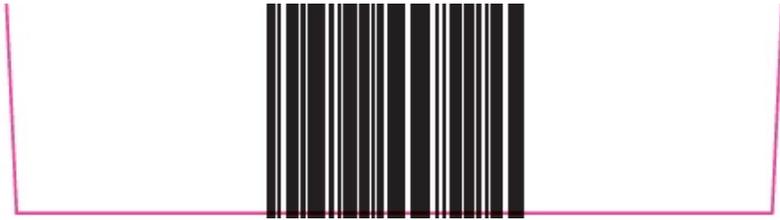
Nasal Decongestant
Phenylephrine HCl

PARENTS:

Learn about teen medicine abuse

www.StopMedicineAbuse.org

*This product is not manufactured or distributed by Novartis Consumer Health, Inc., distributor of Theraflu® Multi-Symptom Severe Cold.



CONVENIENT RECLOSEING TAB

DO NOT USE IF PRINTED PACKETS ARE
TORN OR PUNCTURED

Drug Facts

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Drug Facts (continued)

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1-800-719-9260

GLUTEN FREE

Distributed By:
Perrigo Company
515 Eastern Avenue,
Allegan, MI 49010
V#5003185
MADE IN MEXICO
ITEM#8Y491

OPEN OTHER END





LOT NO.

EXP.

: 8Y491 TM C3

SOUND BODY SEVERE COLD AND FLU

acetaminophen, dextromethorphan hbr, phenylephrine hcl powder, for solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:50594-022
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	500 mg
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	20 mg
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	10 mg

Inactive Ingredients

Ingredient Name	Strength
ACESULFAME POTASSIUM (UNII: 23OV73Q5G9)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
ASPARTAME (UNII: Z0H242BBR1)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
SUCROSE (UNII: C151H8M554)	
TRIBASIC CALCIUM PHOSPHATE (UNII: 91D9GV0Z28)	

Product Characteristics

Color		Score	
Shape		Size	
Flavor	HONEY (green tea) , LEMON (green tea)	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50594-022-91	6 in 1 CARTON; Type 0: Not a Combination Product	02/16/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	02/16/2018	

Labeler - Big Lots Stores, Inc. (017885351)

Revised: 8/2019

Big Lots Stores, Inc.