SEVERE COLD AND COUGH DAYTIME- acetaminophen, dextromethorphan, phenylephrine powder, for solution
Chain Drug Manufacturing Assn

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.

QC Quality Choice Severe Cold and Cough Daytime Berry infused with Menthol and Green Tea Flavors 6 Packets

Active ingredients (in each packet)

Acetaminophen 650 mg

Dextromethorphan hydrobromide 20 mg

Phenylephrine hydrochloride 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 are allergic to acetaminophen
- with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or a 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 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

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
- symptoms do not get better or worsen
- pain, cough or nasal congestion gets worse or lasts more than 7 days
- cough comes back or occurs fever, rash or headache that lasts. These could be

signs of a serious condition.

If pregnant or breast-feeding,

ask a health care professional before use.

Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center right away at 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**
- take every 4 hours; while symptoms persist. Do not to exceed 5 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 8 oz. of cool water: Stir briskly before and after heating: Do not overheat.

Other information

- each packet contains: **potassium 6mg**
- store at room temperature. Protect from excessive heat and moisture.

Inactive ingredients

citric acid, FD &C blue # 1, FD & C red # 40,flavors, maltodextrin, potassium chloride, silica ,sucralose, sucrose

Questions or Comments?

1-866-467-2748

Principal Display

QC Quality Choice

*Compare to the Active Ingredients in Theraflu® Daytime Severe Cold &Cough

SEVERE COLD & COUGH

Daytime

Acetaminophen

Pain Reliever/ Fever Reducer

Dextromethorphan HBr

Cough suppressant

Phenylephrine HCl

Nasal Decongestant

Aspartame Free ●Sodium Free

Relieves:

Nasal Congestion
Cough/Body Ache
Sore Throat Pain
Headache/Fever

Berry Infused with Menthol & Green Tea Flavors

6 Packets

READ ALL WARNINGS AND DIRECTIONS ON CARTON BEFORE USE, KEEP CARTON FOR REFERENCE, DO NOT DISCARD,

*This product is not manufactured or distributed by GSK Consumer Healthcare, distributor of Theraflu® Daytime Severe Cough & Cold

TAMPER EVIDENT INNER UNIT: DO NOT USE IF SEALED PACKET IS TORN OR BROKEN.

Distributed by: C.D.M.A., Inc

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Novi, MI 48375

www.qualitychoice.com

Questions: 248-449-9300

SEVERE COLD AND COUGH DAYTIME

acetaminophen, dextromethorphan, phenylephrine powder, for solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63868-290
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	650 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
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
SUCROSE (UNII: C151H8M554)	

Product Characteristics

Color		**Score**	
Shape		**Size**	
Flavor	BERRY (Infused with Menthol and Green Tea Flavors)	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63868-290-06	6 in 1 CARTON	07/25/2018	
1		1 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 final	part341	07/25/2018	

Labeler - Chain Drug Manufacturing Assn (011920774)

Revised: 1/2022 Chain Drug Manufacturing Assn