

**SEVERE CONGESTION AND COUGH- dextromethorphan hbr,
guaifenesin tablet, film coated
CVS WOONSOCKET PRESCRIPTION CENTER, INCORPORATED**

CVS 44-808 RESERVED NDC 51316-808-09

Active ingredients (in each caplet)

Dextromethorphan HBr 10 mg
Guaifenesin 200 mg

Purpose

Cough suppressant|
Expectorant

Uses

- helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus and make coughs more productive
- temporarily relieves:
 - cough due to minor throat and bronchial irritation as may occur with the common cold or inhaled irritants
 - the intensity of coughing
 - the impulse to cough to help you get to sleep

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.

Ask a doctor before use if you have

- persistent or chronic cough such as occurs with smoking, asthma, chronic bronchitis, or emphysema
- cough that occurs with too much phlegm (mucus)

Stop use and ask a doctor if

cough persists more than 1 week, tends to recur, 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

- **do not take more than directed**
- do not take more than 12 caplets in any 24-hour period
- adults and children 12 years and over: take 2 caplets every 4 hours
- children under 12 years: do not use

Other information

- **each caplet contains:** sodium 3 mg
- **TAMPER EVIDENT: DO NOT USE IF OUTER PACKAGE IS OPENED OR BLISTER IS TORN OR BROKEN**
- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
- see end flap for expiration date and lot number

Inactive ingredients

FD&C blue #2 aluminum lake, FD&C red #40 aluminum lake, magnesium stearate, maltodextrin, mica, microcrystalline cellulose, polyethylene glycol, polysorbate 80, polyvinyl alcohol, povidone, pregelatinized starch, silicon dioxide, sodium starch glycolate, stearic acid, talc, titanium dioxide

Questions or comments?

1-800-426-9391

Principal display panel

♥CVS™

NDC 51316-808-09

actual size

**Maximum Strength†
SEVERE CONGESTION
& COUGH**

Dextromethorphan HBr, Cough Suppressant
Guaifenesin, Expectorant

Compare to Maximum Strength Mucinex® FAST-MAX
Severe Congestion & Cough active ingredients*

Multi-Symptom
Relieves Chest Congestion
Controls Cough | Thins & Loosens Mucus

20 CAPLETS

†Under OTC monograph.

TAMPER EVIDENT: DO NOT USE IF PACKAGE IS OPENED OR IF BLISTER UNIT IS TORN, BROKEN OR SHOWS ANY SIGNS OF TAMPERING

PARENTS:

**Learn about teen medicine abuse
www.StopMedicineAbuse.org**

Distributed by: CVS Pharmacy, Inc.

One CVS Drive, Woonsocket, RI 02895

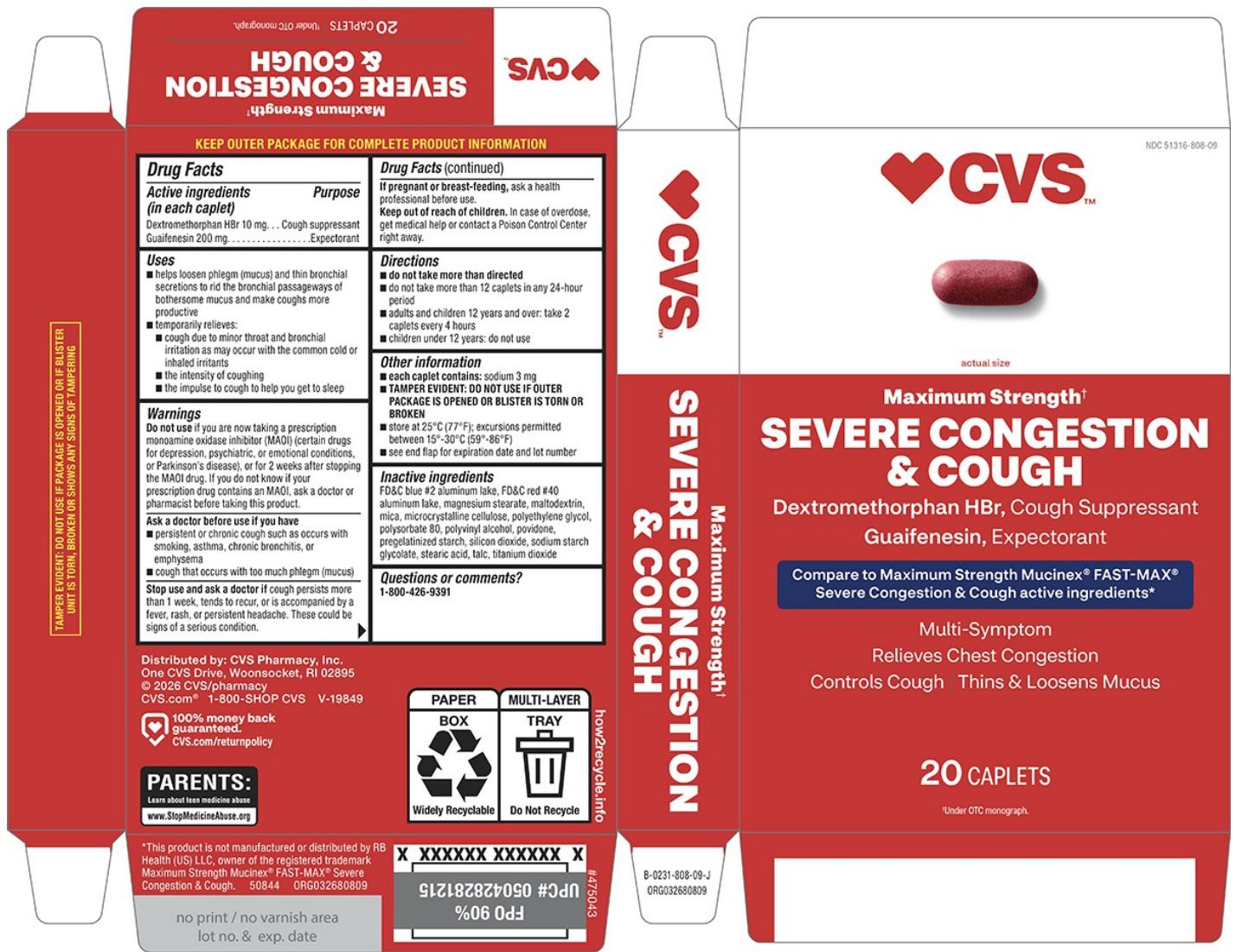
© 2026 CVS /pharmacy

CVS.com® 1- 80 0-SHOP CVS V-19849

*This product is not manufactured or distributed by RB Health (US) LLC, owner of the registered trademark Maximum Strength Mucinex® FAST-MAX® Severe Congestion & Cough. 50844 ORG032680809

100% money back
guaranteed.

CVS.com/returnpolicy



CVS 44-808

SEVERE CONGESTION AND COUGH

dextromethorphan hbr, guaifenesin tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:51316-808
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

FD&C BLUE NO. 2 ALUMINUM LAKE (UNII: 4AQJ3LG584)
FD&C RED NO. 40 ALUMINUM LAKE (UNII: 6T47AS764T)
MAGNESIUM STEARATE (UNII: 70097M6I30)
MALTODEXTRIN (UNII: 7CVR7L4A2D)
MICA (UNII: V8A1AW0880)
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)
POLYSORBATE 80 (UNII: 6OZP39ZG8H)
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)
STARCH, CORN (UNII: O8232NY3SJ)
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)
STEARIC ACID (UNII: 4ELV7Z65AP)
TALC (UNII: 7SEV7J4R1U)
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)

Product Characteristics

Color	red	Score	no score
Shape	OVAL	Size	19mm
Flavor		Imprint Code	44;808
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51316-808-09	2 in 1 CARTON	05/11/2026	
1		10 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	05/11/2026	

Labeler - CVS WOONSOCKET PRESCRIPTION CENTER, INCORPORATED (062312574)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	pack(51316-808)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(51316-808) , pack(51316-808)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867894	manufacture(51316-808)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		117025878	manufacture(51316-808)

Revised: 5/2026

CVS WOONSOCKET PRESCRIPTION CENTER, INCORPORATED