ROBITUSSIN MAXIMUM STRENGTH 12 HOUR COUGH AND MUCUS RELIEFdextromethorphan hydrobromide, guaifenes in tablet, extended release Wyeth Consumer Healthcare LLC

Robitussin[®] Maximum Strength 12 Hour Cough and Mucus Relief Extended-Release

Drug Facts

Active ingredients (in each
extended-release tablet)PurposesDextromethorphan HBr 60 mgCough suppressantGuaifenesin 1200 mgExpectorant

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

- for children under 12 years of age
- 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 accompanied by too much phlegm (mucus)

When using this product

• do not use more than directed

Stop use and ask a doctor if

• cough lasts more than 7 days, comes back, or occurs with fever, rash, or persistent headache. These could be signs of a serious illness.

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 (1-800-222-1222).

Directions

- do not crush, chew, or break tablet
- take with a full glass of water
- this product can be administered without regard for timing of meals
- adults and children 12 years and older: 1 tablet every 12 hours; not more than 2 tablets in 24 hours

• children under 12 years of age: do not use

Other information

- each tablet contains: magnesium 25 mg
- store between 20-25°C (68-77°F)

Inactive ingredients

carbomer homopolymer type B, copovidone, D&C yellow #10 aluminum lake, hypromellose, magnesium hydroxide, magnesium stearate, microcrystalline cellulose, silicon dioxide

Questions?

call weekdays from 9 AM to 5 PM EST at 1-800-762-4675

Distributed by: Pfizer, Madison, NJ 07940 USA

PRINCIPAL DISPLAY PANEL - 1200 mg/60 mg Tablet Blister Pack

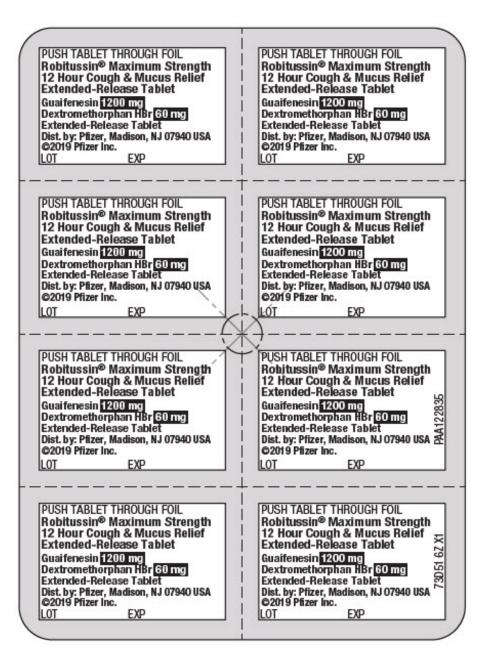
PUSH TABLET THROUGH FOIL

Robitussin[®] Maximum Strength 12 Hour Cough & Mucus Relief Extended-Release Tablet

Guaifenesin 1200 mg Dextromethorphan HBr 60 mg Extended-Release Tablet

Dist. by: Pfizer, Madison, NJ 07940 USA ©2019 Pfizer Inc.

LOT EXP



PRINCIPAL DISPLAY PANEL - 8 Tablet Blister Pack Carton

NEW!

Robitussin®

MAXIMUM STRENGTH 12 Hour Cough & Mucus Relief EXTENDED-RELEASE TABLETS

GUAIFENESIN & DEXTROMETHORPHAN HYDROBROMIDE 1200 mg/60 mg EXTENDED-RELEASE TABLETS

Expectorant & Cough Suppressant

Actual Size

Controls Cough

OThins & Loosens Mucus

8 EXTENDED-RELEASE TABLETS





ROBITUSSIN MAXIMUM STRENGTH 12 HOUR COUGH AND MUCUS RELIEF

dextromethorphan hydrobromide, guaifenesin tablet, extended release

Product Information					
Product T ype	HUMAN OTC DRUG	Item Code (S	ource)	NDC:0031-8765	
Route of Administration	ORAL				
Active Ingredient/Active Moi	ety				
Ingredient Name			Basis of St	rength	Strength
DEXTROMETHORPHAN HYDROBRO (DEXTROMETHORPHAN - UNII:7355X3		DEXTROMETHORI HYDROBROMIDE	PHAN	60 mg	
GUAIFENESIN (UNII: 495W7451VQ) (C	Q)	GUAIFENESIN		1200 mg	

			Ingre	dient Name			Strength
		OLYME	ER TYPE B (ALLYL PENT	AERYTHRITOL OR AI	LIYL SUCROSE CROSS	LINKED)	
`	III: K6 MOM3T5YL)	4 /11311					
	POVIDONE K25-3	,					
	C YELLOW NO. 1			~			
			FIED (UNII: 3NXW29V3WO)			
		,	INII: NBZ3QY004S)				
	GNESIUM STEAR		ULOSE (UNII: OP1R32D61	II)			
	ICON DIO XIDE (U			.0)			
		л п . L15	/LOADO+)				
Pr	oduct Characte	eristic	8				
Col			YELLOW	Score		no score	
Shape			OVAL	Size		22mm	
Flavor				Imprint Code		2424	
	ntains			Inprint Cour			
001							
Pa	ckaging						
	ckaging Item Code		Package Descr	iption	Marketing Start Date	e Marketing	End Date
#	00	1 in 1 C.	Package Descr	iption	Marketing Start Date	e Marketing	End Date
# 1 N	Item Code		0		J	e Marketing	End Date
# 1 N 1	Item Code	4 in 1 B	ARTON LISTER PACK; Type 0: No		J	e Marketing	End Date
# 1 1 N 1 2 N	Item Code NDC:0031-8765-04	4 in 1 B 1 in 1 C.	ARTON LISTER PACK; Type 0: No	t a Combination Product	0 6/14/20 19 0 6/14/20 19	e Marketing	End Date
# N 1 N 2 N 2 N	Item Code NDC:0031-8765-04	4 in 1 B 1 in 1 C. 8 in 1 B	ARTON LISTER PACK; Type 0: No ARTON LISTER PACK; Type 0: No	t a Combination Product	0 6/14/20 19 0 6/14/20 19	e Marketing	End Data
 # 1 N 2 3 	Item Code NDC:0031-8765-04 NDC:0031-8765-08	4 in 1 B 1 in 1 C. 8 in 1 B 2 in 1 C	ARTON LISTER PACK; Type 0: No ARTON LISTER PACK; Type 0: No	t a Combination Product t a Combination Product	0 6/14/20 19 0 6/14/20 19 0 6/14/20 19	e Marketing	End Date
# 1 N 2 N 2 N 3 N	Item Code NDC:0031-8765-04 NDC:0031-8765-08	4 in 1 B 1 in 1 C. 8 in 1 B 2 in 1 C	ARTON LISTER PACK; Type 0: No ARTON LISTER PACK; Type 0: No ARTON	t a Combination Product t a Combination Product	0 6/14/20 19 0 6/14/20 19 0 6/14/20 19	Marketing	End Dat
 # 1 N 2 3 	Item Code NDC:0031-8765-04 NDC:0031-8765-08	4 in 1 B 1 in 1 C. 8 in 1 B 2 in 1 C	ARTON LISTER PACK; Type 0: No ARTON LISTER PACK; Type 0: No ARTON	t a Combination Product t a Combination Product	0 6/14/20 19 0 6/14/20 19 0 6/14/20 19	e Marketing	End Dat
 # 1 N 1 2 N 3 N 3 	Item Code NDC:0031-8765-04 NDC:0031-8765-08 NDC:0031-8765-16	4 in 1 B 1 in 1 C 8 in 1 B 2 in 1 C 8 in 1 B	ARTON LISTER PACK; Type 0: No ARTON LISTER PACK; Type 0: No ARTON LISTER PACK; Type 0: No	t a Combination Product t a Combination Product	0 6/14/20 19 0 6/14/20 19 0 6/14/20 19	e Marketing	End Date
 # 1 N 2 N 3 N 3 	Item Code NDC:0031-8765-04 NDC:0031-8765-08	4 in 1 B 1 in 1 C. 8 in 1 B 2 in 1 C 8 in 1 B	ARTON LISTER PACK; Type 0: No ARTON LISTER PACK; Type 0: No ARTON LISTER PACK; Type 0: No	t a Combination Product t a Combination Product t a Combination Product	0 6/14/20 19 0 6/14/20 19 0 6/14/20 19	e Marketing	

Labeler - Wyeth Consumer Healthcare LLC (828831730)

Revised: 6/2019

Wyeth Consumer Healthcare LLC