MUCUS RELIEF COUGH AND CONGESTION DM- guaifenesin and dextromethorphan hbr tablet

Advance Pharmaceutical 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.

Guaifenes in 400 mg - DextromethorphanCaplets 20 mg Caplets

Active ingredients (in each caplet)

Dextromethorphan HBr 20 mg Guaifenesin 400 mg

Purpose

Cough suppressant Expectorant

Uses

- temporarily relieves cough due to minor throat and bronchial irritation associated with the common cold
- helps loosen phlegm (mucus) and thin bronchial secretions to make coughs more productive

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

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

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 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

- take with a full glass of water
- adults and children 12 years of age and over: 1 tablet every 4 hours. Do not take more than 6 tablets in 24 hours.
- children under 12 years: do not use

Other information

store at 15°-30°C (59°-86°F)

Inactive ingredients

Colloidal silicon dioxide, magnesium stearate, maltodextrin, microcrystalline cellulose, povidone, stearic acid

Questions or comments?

call 631-981-4600, 8.30 am-4.30 pm ET, Monday - Friday TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING

Principal Display Panel

Mucus Relief Cough and Congestion DM

NDC 17714-150-60



MUCUS RELIEF COUGH AND CONGESTION DM

guaifenesin and dextromethorphan hbr tablet

Product Information					
Product T ype	HUMAN OTC DRUG	Item Code (Source)	NDC:17714-150		
Route of Administration	ORAL				
Active Ingredient/Active Moiety					

Ingredient Name				Basis of S	Basis of Strength		
	ROMETHORPHAN HYDROBROMIDE (UNII: 9 D2RTI9 KYH)DIROMETHORPHAN - UNII:7355X3ROTS)H					20 mg	
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ) GUAIFENESI						400 mg	
Inactive Ingredien	ts						
Ingredient Name						Strength	
MAGNESIUM STEARATE (UNII: 70097M6I30)							
MALTODEXTRIN (UNII	: 7CVR7L4A2I))					
POVIDONE (UNII: FZ98	9 GH9 4E)						
SILICON DIO XIDE (UN	II: ETJ7Z6XBU	J4)					
STEARIC ACID (UNII: 41	ELV7Z65AP)						
MICRO CRYSTALLINE	CELLULOSE	(UNII: OP1R32D6	1U)				
Product Character	ristics						
Color	white	2	Score		2 pieces		
Shape	OVA	L	Size		17mm		
Flavor			Imprint Code		AP;150		
Contains							
Packaging							
		Package Descr	iption	Marketing Start Date	Marketin	ng End Date	
# Item Code		•	iption Combination Product	Marketing Start Date 08/01/2017	Marketii	ng End Date	
# Item Code		•	•		Marketii	ng End Dat	
# Item Code		•	•		Marketii	ıg End Dato	
# Item Code 1 NDC:17714-150-60 6	0 in 1 BOTTLI	•	•		Marketin	ng End Date	
 # Item Code 1 NDC:17714-150-60 6 Marketing Info 	0 in 1 BOTTLI	E; Type 0: Not a C	Combination Product	08/01/2017			
	0 in 1 BOTTLI	E; Type 0: Not a C	•			ng End Date ng End Date	

Labeler - Advance Pharmaceutical Inc. (078301063)

Registrant - Advance Pharmaceutical Inc. (078301063)

Establishment									
Name	Address	ID/FEI	Business Operations						
Advance Pharmaceutical Inc.		078301063	manufacture(17714-150)						

Revised: 10/2017

Advance Pharmaceutical Inc.