DEXTROMETHORPHAN HBR 15 MG- dextromethorphan hbr 15 mg capsule, liquid filled
AMZ789 LLC

DEXTROMETHORPHAN HBR 15 MG

Dextromethorphan HBr, USP 15 mg

Cough suppressant

Use

temporarily relieves cough due to minor throat and bronchial irritation as may occur with a cold

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.

a cough that occurs with too much phlegm (mucus)
a cough that lasts or is chronic as occurs with smoking, asthma, or emphysema

Stop use and ask a doctor if cough lasts for more than 7 days, comes back, or is accompanied by 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.

do not take more than 8 capsules in any 24-hour period
this adult product is not intended for use in children under 12 years of age

adults and children 12 years of age and over	take 2 capsules every 6 to 8 hours, as needed
children under 12 years of age	do not use

store at 20-25°C (68-77°F)
avoid excessive heat above 40°C (104°F)
protect from light

FD&C blue #1, FD&C Red #40, gelatin, glycerin, polyethylene glycol, povidone, propylene glycol, purified water, sorbitol special, and white edible ink

DEXTROMETHORPHAN HBR 15 MG

dextromethorphan hbr 15 mg capsule, liquid filled

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73629-005
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	15 mg

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
POVIDONE (UNII: FZ989GH94E)	
GELATIN (UNII: 2G86QN327L)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

GLYCERIN (UNII: PDC6A3C0OX)	
SORBITOL (UNII: 506T60A25R)	

Product Characteristics

Color	red (CLEAR)	**Score**	score with uneven pieces
Shape	OVAL (CAPSULE)	**Size**	10mm
Flavor		**Imprint Code**	PC6
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73629-005-10	100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/01/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	05/01/2022	

Labeler - AMZ789 LLC (117410213)

Establishment

Name	Address	ID/FEI	Business Operations
Humanwell Puracap Pharmaceuticals (Wuhan) Co., Ltd		421293287	manufacture(73629-005)

Revised: 1/2025 AMZ789 LLC