MUCUS RELIEF- guaifenesin tablet, extended release BJWC (Berkley & Jensen / BJ's)

BJs Guaifenesin Extended Release Tablets 1200 mg

Drug Facts

Active ingredient (in each extended-release tablet)

Guaifenesin, USP 1200 mg

Purpose

Expectorant

Uses

 helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus and make coughs more productive

Warnings

Do not use

• for children under 12 years of age

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)

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 (1-800-222-1222) right away.

Directions

- do not crush, chew, or break extended-release tablet
- take with a full glass of water
- this product can be administered without regard for the timing of meals
- adults and children 12 years of age and over: 1 extended-release tablet every 12 hours. Do not exceed 2 extended-release tablets in 24 hours.
- children under 12 years of age: do not use

Other information

store at 20-25°C (68-77°F)

Inactive ingredients colloidal silicon dioxide, copovidone, FD&C blue no. 1 aluminum lake, hypromellose, magnesium stearate, maltodextrin, microcrystalline cellulose, povidone (K-30), sodium starch glycolate, stearic acid.

Questions or comments? Call 1-877-290-4008

724R 84ct label



724R 84ct Carton



MUCUS RELIEF

quaifenesin tablet, extended release

Prod		

Product Type HUMAN OTC DRUG Item Code (Source) NDC:68391-724

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	1200 mg

Inactive Ingredients			
Ingredient Name	Strength		
STEARIC ACID (UNII: 4ELV7Z65AP)			
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)			
COPOVIDONE K25-31 (UNII: D9C330MD8B)			
FD&C BLUE NO. 1 ALUMINUM LAKE (UNII: J9EQA3S2JM)			
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)			
POVIDONE K30 (UNII: U725QWY32X)			
SODIUM STARCH GLYCOLATE TYPE A (UNII: H8AV0SQX4D)			
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)			
MAGNESIUM STEARATE (UNII: 70097M6I30)			
MALTODEXTRIN (UNII: 7CVR7L4A2D)			

Product Characteristics			
Color	blue (Blue and White)	Score	no score
Shape	OVAL	Size	22mm
Flavor		Imprint Code	41;1200
Contains			

F	Packaging				
#	tem Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:68391-724- 95	1 in 1 CARTON	08/20/2025		
1		84 in 1 BOTTLE; Type 0: Not a Combination Product			

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA217780	08/20/2025	

Labeler - BJWC (Berkley & Jensen / BJ's) (159082692)

Registrant - TIME CAP LABORATORIES, INC. (037052099)

Establishment				
Name	Address	ID/FEI	Business Operations	
MARKSANS PHARMA LIMITED		925822975	manufacture(68391-724)	