GENCARE-NASAL DECONGESTANT PE- phenylephrine hydrochloride tablet
Pioneer Life Sciences, LLC

Gencare-Nasal Decongestant PE

Active Ingredient (in each tablet)

Phenylephrine HCl 10mg

Purpose

Nasal decongestant

USES:

- temporarily relieves sinus congestion and pressure
- temporarily relieves nasal congestion due to the common cold, hay fever or other upper respiratory allergies

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

- heart disease
- high blood pressure
- thyroid disease
- diabetes
- trouble urinating due to an enlarged prostate gland.

When using this product do not exceed recommended dose.

Stop use and ask a doctor if

- nervousness, dizziness, or sleeplessness occur symptoms do not improve within 7 days or occur with a fever.

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:

- take 1 tablet every 4 hours
- adults and children 12 years and over: do not take more than 6 tablets in 24 hours
- children under 12 years: ask a doctor

OTHER INFORMATION

- store between 20-25°C (68-77°F)
- do not use if seal under cap is broken or missing

INACTIVE INGREDIENTS

Crosscarmellose Soidum, Dibasic Calcium Phosphate, Magnesium Stearate, Microcrystalline Cellulose, Purified water. Readycoat Red contains: polyvinyl alcohol, polyethylene glycol, hypromellose, methacrylic acid copolymer, talc, titanium dioxide, allura red lake, indigo carmine lake.

QUESTIONS OR COMMENTS?

Call (732) 994-2808 or email support@gencare.health

GENCARE-NASAL DECONGESTANT PE

phenylephrine hydrochloride tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72090-039
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE	10 mg

Inactive Ingredients	
Ingredient Name	**Strength**
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
DICALCIUM PHOSPHATE (UNII: L11K75P92J)	

CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYVINYL ALCOHOL (UNII: 532B59J990)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
METHACRYLIC ACID - METHYL METHACRYLATE COPOLYMER (1:1) (UNII: 74G4R6TH13)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
FD&C RED NO. 40 ALUMINUM LAKE (UNII: 6T47AS764T)	
INDIGOTINDISULFONATE SODIUM (UNII: D3741U8K7L)	

Product Characteristics

Color	red	**Score**	no score
Shape	ROUND	**Size**	7mm
Flavor		**Imprint Code**	3P
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72090-039-01	300 in 1 BOTTLE; Type 0: Not a Combination Product	02/18/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	02/18/2025	

Labeler - Pioneer Life Sciences, LLC (014092742)

Registrant - Pioneer Life Sciences, LLC (014092742)

Revised: 2/2025 Pioneer Life Sciences, LLC