NEO-SYNEPHRINE REGULAR- phenylephrine hydrochloride spray
BF ASCHER AND CO INC

Neo-Synephrine Regular

Phenylephrine hydrochloride 0.50%

Nasal decongestant

Temporarily relieves nasal congestion due to:
- common cold
- hay fever
- upper respiratory allergies

Temporarily relieves sinus congestion and pressure

Shrinks swollen membranes so you can breathe more freely

Temporarily restores freer breathing through the nose.

When using this product:
- temporary discomfort such as burning, stinging, sneezing, or an increase in nasal discharge may occur.
- use of this container by more than one person may spread infection.
- use only as directed. Frequent or prolonged use may cause nasal congestion to recur or worsen.

Do not use if you have ever had an allergic reaction to this product or any of its ingredients.

Do not use more than directed.

Do not use for more than 3 days.

Ask a doctor before use if you have:
- heart disease
- high blood pressure
- thyroid disease
- diabetes
- trouble urinating due to an enlarged prostate gland

Stop use and as a doctor if symptoms persist.

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If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

- to spray, squeeze bottle quickly and firmly
- adults and children 12 years of age and older: 2 or 3 sprays in each nostril not more often then every 4 hours
- Children under 12 years of age: ask a doctor

- Store at room temperature, 59º - 86º F (15º -30º C)
- Retain carton for future reference on full labeling

anhydrous citric acid, benazalkonium chloride, purified water, sodium chloride, sodium citrate

Call 1-800-324-1880, 7:30 - 4:00 Central, Monday - Friday, or visit www.bfascher.com

NEO-SYNEPHRINE REGULAR

phenylephrine hydrochloride spray

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0225-0805
Route of Administration	NASAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	0.5 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
WATER (UNII: 059QF0KO0R)	
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0225-0805-47	15 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	07/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	07/01/2020	

Labeler - BF ASCHER AND CO INC (003854403)

Revised: 11/2023 BF ASCHER AND CO INC