

**NEO-SYNEPHRINE MAXIMUM- phenylephrine hydrochloride spray**  
**BF ASCHER AND CO INC**

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**Neo-Synephrine Extra**

Phenylephrine hydrochloride 1.0%

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 ask 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 than every 4 hours
- Children under 12 years of age: ask a doctor

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

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

Call 1-800-324-1880, 7:30 - 4:00 Central, Monday - Friday, or visit [www.bfascher.com](http://www.bfascher.com)



**NEO-SYNEPHRINE MAXIMUM**

phenylephrine hydrochloride spray

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:0225-0810
<b>Route of Administration</b>	NASAL		

### Active Ingredient/Active Moiety

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

### Inactive Ingredients

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

### Packaging

<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:0225-0810-47	15 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	07/01/2020	

### Marketing Information

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

**Labeler** - BF ASCHER AND CO INC (003854403)

Revised: 11/2023

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