

EQUALINE NASAL- phenylephrine hydrochloride spray
United Natural Foods, Inc. dba UNFI

SuperValu Inc. Nasal Spray Drug Facts

Active ingredient

Phenylephrine hydrochloride 1%

Purpose

Nasal decongestant

Uses

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

Warnings

Ask a doctor before use if you have

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

When using this product

- **do not use more than directed**
- do not use for more than 3 days. Frequent or prolonged use may cause nasal congestion to come back or get worse.
- use only as directed
- 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

Stop use and ask a doctor if

symptoms persist

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. (1-800-222-1222)

Directions

- adults and children 12 years and over: 2 or 3 sprays in each nostril not more often than every 4 hours
- children under 12 years: ask a doctor
- Use instructions: with head in a normal, upright position, put atomizer tip into nostril. Squeeze bottle with firm, quick pressure while inhaling. Wipe nozzle clean after each use.

Other information

- store at 20°-25°C (68°-77°F)

Inactive ingredients

benzalkonium chloride solution, benzyl alcohol, boric acid, purified water, sodium borate

Questions or comments?

1-855-423-2630

Principal Display Panel

compare to 4 Way® Fast Acting Nasal Spray active ingredient

EQUALINE®

fast acting nasal spray

phenylephrine hydrochloride 1 % (nasal decongestant)

fast acting relief of:

- nasal congestion
- sinus congestion
- sinus pressure
- swollen nasal membranes

1 FL OZ (30mL)



EQUALINE NASAL

phenylephrine hydrochloride spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:41163-648
Route of Administration	NASAL		

Active Ingredient/Active Moiety

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

Inactive Ingredients

Ingredient Name	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
BORIC ACID (UNII: R57ZHV85D4)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41163-648-10	1 in 1 CARTON	07/26/2018	
1		30 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	07/26/2018	

Labeler - United Natural Foods, Inc. dba UNFI (943556183)

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