#### NASAL- oxymetazoline hydrochloride spray H E B

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# **HEB Nasal Spray Drug Facts**

### **Active ingredient**

Oxymetazoline hydrochloride 0.05%

# **Purpose**

Nasal decongestant

#### Uses

- temporarily relieves nasal congestion due to:
- common cold
- hay fever
- upper respiratory allergies
- temporarily relieves sinus congestion and pressure
- shrinks swollen nasal membranes so you can breathe more freely

# Warnings

# 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 use more than directed
- do not use for more than 3 days. Use only as directed. Frequent or prolonged use may cause nasal congestion to recur or worsen.
- 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 6 to under 12 years of age (with adult supervision): 2 or 3 sprays in each nostril not more often than every 10 to 12 hours. Do not exceed 2 doses in any 24-hour period.
- children under 6 years of age: ask a doctor

**To Use:** Shake well before use. Hold white tabs, <u>SQUEEZE</u> grooved area of cap <u>FIRMLY</u> and turn counter clockwise. Before using the first time, prime metered pump by depressing pump firmly several times. To spray, hold bottle with thumb at base and nozzle between first and second fingers. Without tilting head, insert nozzle into nostril. Fully depress rim with a firm, even stroke and sniff deeply. Wipe nozzle clean after use. Secure cap after use.

#### Other information

- store at 20-25°C (68-77°F)
- · retain carton for future reference on full labeling

# Inactive ingredients

benzalkonium chloride solution, benzyl alcohol, camphor, dibasic sodium phosphate, edetate disodium, eucalyptol, menthol, microcrystalline cellulose and carboxymethylcellulose sodium, monobasic sodium phosphate, polyethylene glycol, povidone, propylene glycol, purified water

#### Questions or comments?

1-800-719-9260

# Package/Label Principal Display Panel

Compare to Afrin® No Drip Severe Congestion active ingredient

No Drip

Maximum Strength plus Menthol

**Nasal Spray** 

Oxymetazoline HCl 0.05%

Nasal Decongestant

Won't Drip From Nose or Down Throat Severe Congestion Fast, Powerful Congestion Relief For Colds & Allergies Pump Mist 12 Hour

1 FL OZ (30 mL)



# NASAL

oxymetazoline hydrochloride spray

# Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:37808-752 Route of Administration NASAL

l	Active Ingredient/Active Moiety		
l	Ingredient Name	<b>Basis of Strength</b>	Strength
	<b>OXYMETAZOLINE HYDROCHLORIDE</b> (UNII: K89MJ0S5VY) (OXYMETAZOLINE - UNII:8VLN5B44ZY)	OXYMETAZ OLINE HYDROCHLORIDE	.05 g in 100 mL

Inactive Ingredients		
Ingredient Name	Strength	
BENZYL ALCOHOL (UNII: LKG8494WBH)		
SODIUM PHOSPHATE, DIBASIC, UNSPECIFIED FORM (UNII: GR686LBA74)		
EDETATE DISODIUM (UNII: 7FLD91C86K)		
EUCALYPTOL (UNII: RV6J6604TK)		
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A)		
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)		
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K6790BS311)		
SODIUM PHOSPHATE, MONOBASIC, ANHYDROUS (UNII: KH7I04HPUU)		
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)		
POVIDONE, UNSPECIFIED (UNII: FZ 989GH94E)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
WATER (UNII: 059QF0KO0R)		
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)		

Product Characteristics			
Color	WHITE (off white)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
NDC:37808- 752-10	1 in 1 CARTON	06/11/2021	
	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	06/11/2021	

# **Labeler -** H E B (007924756)

Revised: 11/2024 H E B