

**SEVERE ANEFRIN- oxymetazoline hydrochloride liquid
WALGREENS**

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

**Severe Anefrin Nasal Spray
No Drip
MAXIMUM STRENGTH
12 Hour
Menthol**

ACTIVE INGREDIENT

Oxymetazoline hydrochloride 0.05%

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.

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, press grooved area of cap firmly and turn counterclockwise. 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.

Questions or comments? 1-800-925-4733.

**DISTRIBUTED BY: WALGREENS CO.
200 WILMOT RD., DEERFIELD, IL 60015**

100% SATISFACTION GUARANTEED

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Inactive Ingredients

Benzalkonium Chloride Solution, Camphor, Carboxymethyl Cellulose Sodium, Cellulose Gum, Disodium EDTA, Disodium Phosphate, Eucalyptol, Glycerin, Menthol, Microcrystalline Cellulose, PEG-6, Povidone, Propylene Glycol, Purified Water, Sodium Phosphate Monobasic

Uses

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

PURPOSE

Nasal Decongestant

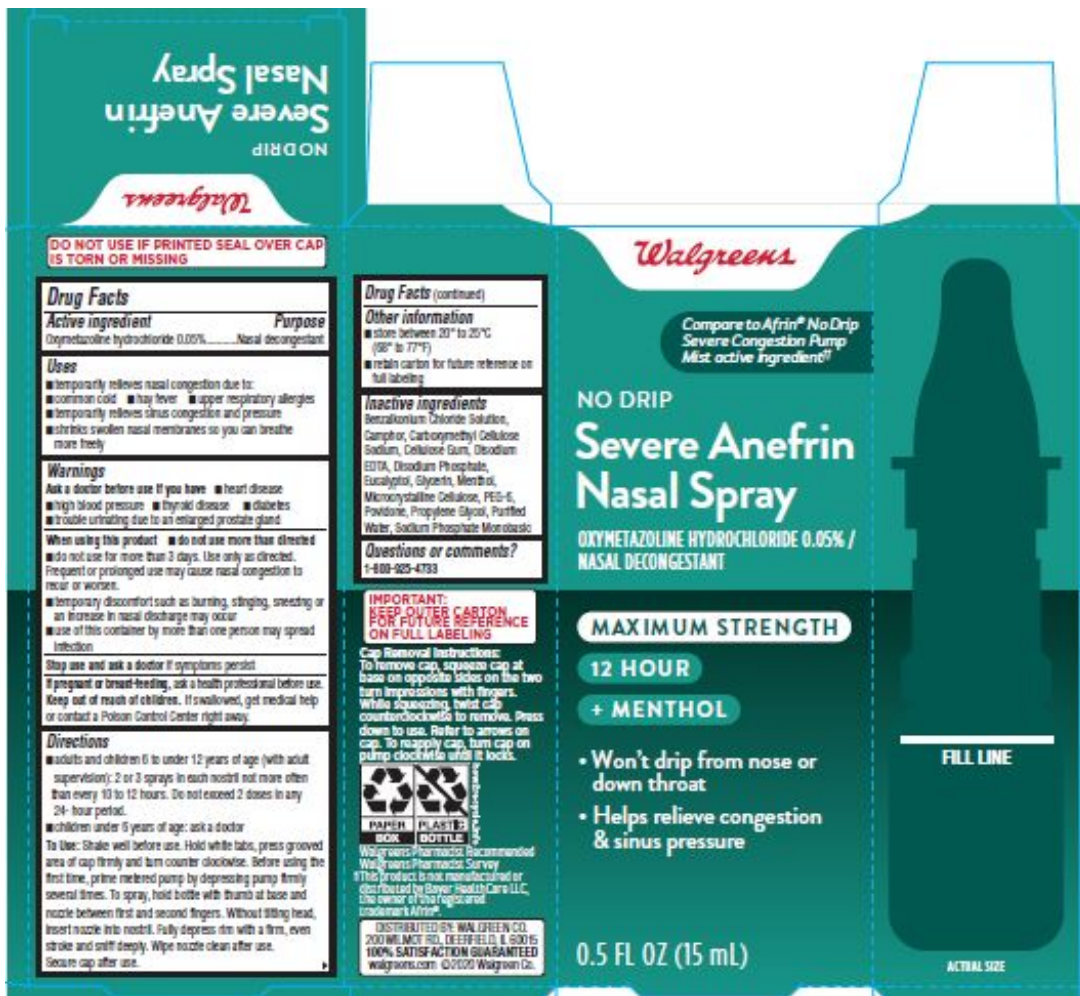
Severe Afrin Nasal Spray

NO DRIP

MAXIMUM STRENGTH

12 HOUR

+ MENTHOL



SEVERE ANEFRIN

oxymetazoline hydrochloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0363-2021
Route of Administration	NASAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OXYMETAZOLINE HYDROCHLORIDE (UNII: K89MJ0S5VY) (OXYMETAZOLINE - UNII: 8VLN5B44ZY)	OXYMETAZOLINE HYDROCHLORIDE	0.05 g in 15 mL

Inactive Ingredients

Ingredient Name	Strength
EDETATE DISODIUM (UNII: 7FLD91C86K)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
EUCALYPTOL (UNII: RV6J6604TK)	
GLYCERIN (UNII: PDC6A3C0OX)	
MENTHOL, (+)- (UNII: C6B1OE8P3W)	
PEG-6 STEARATE (UNII: 8LQC57C6B0)	

CAMPHOR, (-)- (UNII: 213N3S8275)	
POVIDONE K30 (UNII: U725QWY32X)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SODIUM PHOSPHATE, DIBASIC, MONOHYDRATE (UNII: BWZ7K44R51)	
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K679OBS311)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-2021-15	1 in 1 BOTTLE, SPRAY	02/08/2021	
1		15 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 final	part341	02/08/2021	

Labeler - WALGREENS (008965063)

Revised: 2/2021

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