

**CARE ONE NASAL- oxymetazoline hcl spray, metered  
American Sales Company**

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**American Sales Company 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, SQUEEZE grooved area of cap FIRMLY 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, dibasic sodium phosphate, edetate disodium, microcrystalline cellulose and carboxymethylcellulose sodium, monobasic sodium phosphate, polyethylene glycol, povidone, purified water

### **Questions or comments?**

**1-800-719-9260**

### **Principal Display Panel**

CAREone®

Compare to the active ingredient in Afrin® No Drip Original

ORIGINAL NASAL SPRAY

Oxymetazoline HCl 0.05% Nasal Decongestant

12hr

No-Drip - Maximum Strength

Fast, Powerful Congestion Relief

For Colds & Allergies

Won't Drip From Nose or Down Throat

Pump Mist

Gluten Free

1 FL OZ (30mL)

**CAREone**  
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**Actual Size**

**CAREone**  
NDC 41520-388-10  
**ORIGINAL NASAL SPRAY**  
Oxymetazoline HCl 0.05% Nasal Decongestant  
**12hr**  
No Drip Maximum Strength  
Gluten Free  
1 FL OZ (30mL)

DO NOT USE IF PRINTED NECKBAND IS BROKEN OR MISSING  
SEE CARTON FOR FULL LABELING

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DISTRIBUTED BY:  
AUSA DISTRIBUTION, LLC  
SALISBURY, NC 28147  
For product questions or concerns,  
contact us at 1-833-992-3872  
Quality guaranteed or your money back.

**Drug Facts**

Active ingredient	Purpose
Oxymetazoline hydrochloride 0.05%...Nasal decongestant	

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**Questions or comments?**  
1-800-719-9260

\*This product is not manufactured or distributed by Bayer HealthCare LLC, distributor of Afrin® No Drip Original.

**How to use:**  
Hold white tabs, **SQUEEZE** grooved area of cap **FIRMLY** and turn counter clockwise. Secure cap after use.

**SQUEEZE grooves FIRMLY while turning**

Hold white tabs **FIRMLY**

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38810 OF C3

# CARE ONE NASAL

oxymetazoline hcl spray, metered

## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:41520-388
<b>Route of Administration</b>	NASAL		

## Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>OXYMETAZOLINE HYDROCHLORIDE</b> (UNII: K89MJ0S5VY) (OXYMETAZOLINE - UNII:8VLN5B44ZY)	OXYMETAZOLINE HYDROCHLORIDE	0.05 g in 100 mL

## Inactive Ingredients

Ingredient Name	Strength
<b>BENZYL ALCOHOL</b> (UNII: LKG8494WBH)	
<b>SODIUM PHOSPHATE, DIBASIC, UNSPECIFIED FORM</b> (UNII: GR686LBA74)	
<b>EDETATE DISODIUM</b> (UNII: 7FLD91C86K)	
<b>MICROCRYSTALLINE CELLULOSE</b> (UNII: OP1R32D61U)	
<b>CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED</b> (UNII: K679OBS311)	
<b>SODIUM PHOSPHATE, MONOBASIC, UNSPECIFIED FORM</b> (UNII: 3980JIH2SW)	
<b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ0SDW1A)	
<b>POVIDONE, UNSPECIFIED</b> (UNII: FZ989GH94E)	
<b>WATER</b> (UNII: 059QF0K00R)	
<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7)	

## Product Characteristics

<b>Color</b>	WHITE (to off white, viscous)	<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>		<b>Imprint Code</b>	
<b>Contains</b>			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41520-388-10	1 in 1 CARTON	07/20/2004	
1		30 mL in 1 BOTTLE, SPRAY; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)		

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	07/20/2004	

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**Labeler** - American Sales Company (809183973)

Revised: 8/2024

American Sales Company