

**SIGNATURE CARE ORIGINAL- oxymetazoline hydrochloride spray**  
**SAFEWAY, INC.**

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**Signature Care Original Nasal Spray 1oz 76095**

Active ingredient Purpose

Oxymetazoline hydrochloride 0.05%..... 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 (1-800-222-1222) 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: consult a doctor
- shake well before use. Remove safety seal. To open, rotate cap to align the marks. Squeeze cap on both sides in a counter-clockwise turn and pull off to remove. To spray, remove clamp and hold bottle with thumb at base and nozzle between first and second fingers. Without tilting the head, insert nozzle into nostril. Fully depress rim with a firm, even stroke and sniff deeply. Wipe nozzle clean after use and snap

cap onto bottle

Other information

- store at room temperature 20-25° (68-77°F)

Inactive ingredients

benzalkonium chloride, benzyl alcohol, edetate disodium, polyethylene glycol 1500, povidone k30, purified water, sodium phosphate dibasic, sodium phosphate monobasic monohydrate

**Distributed by:**

Better Living Brands LLC

P.O. Box 99

Pleasanton, CA 94566-0009

Made in China



Quality  
Guaranteed

Pump Mist  
Original Congestion  
**Nasal Relief**  
Oxymetazoline Hydrochloride 0.05%  
(Nasal Decongestant)

**Drug Facts (continued)**

**Inactive ingredients**

benzalkonium chloride, benzyl alcohol,  
edetate disodium, polyethylene glycol  
1500, povidone K30, purified water,  
sodium phosphate dibasic, sodium  
phosphate monobasic monohydrate

**Questions or comments?**  
Call 1-888-645-8204 (Mon - Fri 9AM -  
5PM PST)

**KEEP OUTER CARTON FOR COMPLETE  
WARNINGS AND PRODUCT INFORMATION.**

**TAMPER EVIDENT: DO NOT USE IF PRINTED SEAL  
AROUND PUMP AND CAP IS BROKEN OR MISSING.**

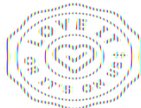
\*This product is not manufactured or  
distributed by Bayer®, distributor of Afrin®  
Original Nasal Spray.

DISTRIBUTED BY:  
BETTER LIVING BRANDS LLC  
P.O. BOX 99  
PLEASANTON, CA 94566-0009  
\*1-888-723-3929

MADE IN CHINA  
76095-ZDP-BX-R1  
NAS-TP-ZDP105-R4



Scan here  
for more  
information



SignatureCare.com

Compare To Afrin®  
Original Nasal Spray  
active ingredient\*

NDC 21130-956-28



Quality  
Guaranteed

CHILD SAFETY CAP  
Pump Mist  
Original Congestion  
**Nasal Relief**  
Oxymetazoline  
Hydrochloride 0.05%  
(Nasal Decongestant)

- Maximum strength
- Fast, powerful  
congestion relief
- Up to 12 hour relief



1 FL OZ (30 mL)



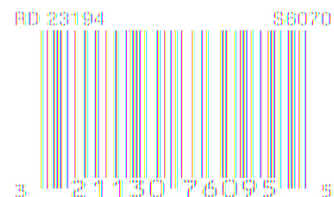
**DIRECTIONS FOR USE:**

**STEP 1:** To open, rotate cap to  
align the marks.  
Squeeze cap on both sides in a  
counter-clockwise turn and  
pull off to remove.

**STEP 2:** To spray, remove  
clamp and hold bottle with  
thumb at base and nozzle  
between first and second  
fingers. Without tilting the  
head, insert nozzle into nostril.  
Fully depress rim in a firm,  
even stroke and sniff deeply.

**STEP 3:** Wipe nozzle clean  
after use, and snap cap back  
onto the bottle.

FULL DIRECTIONS IN THE  
DRUG FACTS PANEL



**Drug Facts**  
**Active ingn**  
Oxymetazoline h

**Uses**  
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**Warnings**

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Keep out of rea  
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**Other inform**

• store at room

Lot:  
Exp:

**SIGNATURE CARE ORIGINAL**

oxymetazoline hydrochloride spray

| Product Information   |                  |   |                      |                    |
|---|------------------|---|----------------------|--------------------|
| Product Type  | HUMAN OTC DRUG   | Item Code (Source)  | NDC:21130-956        |                    |
| Route of Administration   | NASAL            |   |                      |                    |
|   |                  |   |                      |                    |
| Active Ingredient/Active Moiety   |                  |   |                      |                    |
| Ingredient Name   |                  | Basis of Strength   | Strength             |                    |
| OXYMETAZOLINE HYDROCHLORIDE (UNII: K89MJ0S5VY)<br>(OXYMETAZOLINE - UNII:8VLN5B44ZY) |                  | OXYMETAZOLINE<br>HYDROCHLORIDE                              | 0.05 g<br>in 100 mL  |                    |
|   |                  |   |                      |                    |
| Inactive Ingredients  |                  |   |                      |                    |
| Ingredient Name   |                  |   | Strength             |                    |
| POVIDONE K30 (UNII: U725QWY32X)   |                  |   |                      |                    |
| WATER (UNII: 059QF0KO0R)  |                  |   |                      |                    |
| EDETATE DISODIUM (UNII: 7FLD91C86K)   |                  |   |                      |                    |
| BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)  |                  |   |                      |                    |
| SODIUM PHOSPHATE DIBASIC DIHYDRATE (UNII: 94255I6E2T)                               |                  |   |                      |                    |
| POLYETHYLENE GLYCOL 1500 (UNII: 1212Z7S33A)   |                  |   |                      |                    |
| SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE (UNII: 593YOG76RN)                         |                  |   |                      |                    |
| BENZYL ALCOHOL (UNII: LKG8494WBH)   |                  |   |                      |                    |
|   |                  |   |                      |                    |
| Packaging   |                  |   |                      |                    |
| #   | Item Code        | Package Description   | Marketing Start Date | Marketing End Date |
| 1   | NDC:21130-956-28 | 1 in 1 CARTON   | 10/18/2023           |                    |
| 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  | 10/18/2023           |                    |

**Labeler -**
SAFEWAY, INC. (009137209)