

**BREAZE NASAL MOISTURIZING- oxymetazoline hydrochloride spray**  
**Samson Pharmaceuticals, Inc.**

*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.*

-----  
**Breaze®**  
**Nasal Spray Moisturizing**

DRUG FACT

**ACTIVE INGREDIENT**

Oxymetazoline hydrochloride, 0.05%

**PURPOSE**

Nasal Decongestant

**Uses**

- ◆ Temporarily relieves nasal congestion due to:
- ◆ Common cold. Hay fever. Sinusitis
- ◆ Upper respiratory allergies
- ◆ Shrinks swollen nasal membranes so you can breathe more freely

**Warnings**

**Ask a doctor before use if you have**

- ◆ **High blood pressure**
- ◆ **Heart disease**
- ◆ **Diabetes**
- ◆ **Thyroid diseases**
- ◆ **Trouble urinating due to 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**

Adult & children 6 to under 12 years of age (with adult supervision): 2 to 3 sprays in age nostril not more often than every 10 to 12 hours. Do not exceed 2 doses in any 24 -hr period.

Children under 6 years of age: ask a doctor. To spray, squeeze bottle quickly and firmly. Do not tilt head backward while spraying. Wipe nozzle clean after use.

## **Other information**

- ◆ Store between 20C to 25C (68 to 77F).
- ◆ Retain carton for future reference on full labeling.

## **Inactive Ingredients**

Benzalkonium Chloride, Edetate Disodium, Glycerin, Polyethelene Glycol, Povidone, Propylene Glycol, Sodium Diphospahte Dibasic, Sodium Phosphate Monobasic, Water.

## **Question or Comments?**

1-888-995-9935

Manufactured by:  
Samson Pharmaceutical  
Commerce, CA 90040  
Made in USA

Visit our website at  
[www.samson.pharmaceutical.com](http://www.samson.pharmaceutical.com)  
Made in USA

## **PRINCIPAL DISPLAY PANEL - 30 mL Bottle Carton**

MADE IN THE U.S.A.  
WITH PRIDE

**Breaze<sup>®</sup>**

**Relieves Congestion**

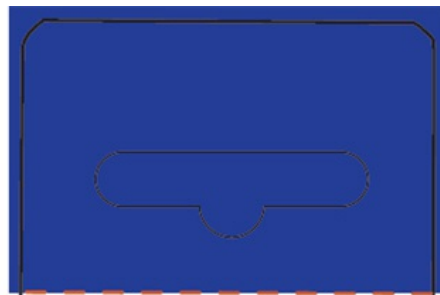
**Nasal  
Spray**

**Moisturizing**

**12 Hour Relief**

- ***Oxymetazoline HCl  
Nasal Solution***
- ***Nasal Decongestant***

***1 FL OZ (30 mL)***



DO NOT USE IF PROTECTIVE SEAL OVER  
CAP IS TORN OR MISSING

**Drug Facts**

**Active Ingredient** Purpose  
Oxymetazoline hydrochloride, 0.05% Nasal decongestant

**Uses**

Temporarily relieves nasal congestion due to:  
• common cold • hay fever • sinusitis  
• upper respiratory allergies  
• Shrinks swollen nasal membranes so you can breathe more freely

**Warnings**

Ask a doctor before use if you have

- Heart disease • High blood pressure
- Diabetes • Thyroid disease
- 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 spray, squeeze bottle quickly and firmly. Do not tilt head backward while spraying. Wipe nozzle clean after use. ▶

**Drug Facts**

(continued)

**Other information**

- Store between 20° to 25 C° (68° to 77° F).
- Retain carton for future reference on full labeling.

**Inactive Ingredients**

Benzalkonium Chloride, Edetate Disodium, Glycerin, Polyethylene Glycol, Povidone, Propylene Glycol, Sodium Phosphate Dibasic, Sodium Phosphate Monobasic, Water.

Keep carton for future reference on full labeling.

Relieves Congestion

**Nasal Spray**

Moisturizing

**Breaze**



Relieves Congestion

**Nasal Spray**

Moisturizing

**12 Hour Relief**

- Oxymetazoline HCl Nasal Solution
- Nasal Decongestant

1 FL OZ (30 mL)

Relieves Congestion

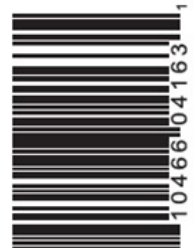
**Nasal Spray**

Moisturizing

**12 Hour Relief**

Manufactured by:  
Samson Pharmaceuticals, Inc.  
Commerce, CA 90040 USA

www.samsonpharmaceutical.com  
Questions or Comments?  
1-888-995-9935



6 10466 04163 1

**BREAZE NASAL MOISTURIZING**

oxymetazoline hydrochloride spray

## Product Information

|                                |                |                           |                |
|--------------------------------|----------------|---------------------------|----------------|
| <b>Product Type</b>            | HUMAN OTC DRUG | <b>Item Code (Source)</b> | NDC:20146-4000 |
| <b>Route of Administration</b> | NASAL          |                           |                |

## Active Ingredient/Active Moiety

| <b>Ingredient Name</b>  | <b>Basis of Strength</b>    | <b>Strength</b>   |
|---|-----------------------------|-------------------|
| <b>Oxymetazoline hydrochloride</b> (UNII: K89MJ0S5VY) (Oxymetazoline - UNII:8VLN5B44ZY) | Oxymetazoline hydrochloride | 0.5 mg<br>in 1 mL |

## Inactive Ingredients

| <b>Ingredient Name</b>   | <b>Strength</b> |
|--|-----------------|
| <b>Benzalkonium Chloride</b> (UNII: F5UM2KM3W7)                  |                 |
| <b>Edetate Disodium</b> (UNII: 7FLD91C86K)                       |                 |
| <b>Glycerin</b> (UNII: PDC6A3C0OX)                               |                 |
| <b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ0SDW1A)       |                 |
| <b>POVIDONE, UNSPECIFIED</b> (UNII: FZ989GH94E)                  |                 |
| <b>Propylene Glycol</b> (UNII: 6DC9Q167V3)                       |                 |
| <b>Sodium Phosphate, Dibasic, Anhydrous</b> (UNII: 22ADO53M6F)   |                 |
| <b>Sodium Phosphate, Monobasic, Anhydrous</b> (UNII: KH7I04HPUU) |                 |
| <b>Water</b> (UNII: 059QF0KO0R)                                  |                 |

## Packaging

| # | <b>Item Code</b> | <b>Package Description</b>                                  | <b>Marketing Start Date</b> | <b>Marketing End Date</b> |
|---|------------------|---|-----------------------------|---------------------------|
| 1 | NDC:20146-4000-2 | 1 in 1 CARTON   | 01/01/2015                  |                           |
| 1 |                  | 30 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product |                             |                           |

## Marketing Information

| <b>Marketing Category</b> | <b>Application Number or Monograph Citation</b> | <b>Marketing Start Date</b> | <b>Marketing End Date</b> |
|---------------------------|---|-----------------------------|---------------------------|
| OTC MONOGRAPH FINAL       | part341   | 01/01/2015                  |                           |

**Labeler** - Samson Pharmaceuticals, Inc. (088169581)

## Establishment

| <b>Name</b>                  | <b>Address</b> | <b>ID/FEI</b> | <b>Business Operations</b> |
|------------------------------|----------------|---------------|----------------------------|
| Samson Pharmaceuticals, Inc. |                | 088169581     | MANUFACTURE(20146-4000)    |

