

**AFRIN NODRIP ALLERGY SINUS NIGHT- oxymetazoline hydrochloride spray**  
**Bayer HealthCare LLC.**

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**Afrin NoDrip Allergy Sinus Night UI1612461**

*Drug Facts*

**Active ingredient Purpose**

Oxymetazoline hydrochloride 0.05%.....Nasal decongestant

**Uses**

- temporarily relieves nasal congestion due to:
- common cold ■ hay fever
- upper respiratory allergies
- reduces swelling of nasal passages so you can breathe more freely
- temporarily relieves sinus congestion and pressure

**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 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 between 20° to 25°C (68° to 77°F)
- retain carton for future reference on full labeling

***Inactive ingredients*** benzalkonium chloride solution, benzyl alcohol, dibasic sodium phosphate, edetate disodium, flavor, glycerin, microcrystalline cellulose and carboxymethylcellulose sodium, monobasic sodium phosphate, polyethylene glycol, povidone, purified water

***Questions or comments? 1-800-317-2165***

**Carton label 15 mL**



Oxymetazoline HCl  
 Nasal Solution-Nasal Decongestant  
**AFRIN**  
**NODRIP**  
 Won't drive from nose or down throat  
 Allergy Sinus  
 Night  
 with soothing chamomile scent  
 PUMP MIST  
 Instant Congestion

Relief from Allergies  
 for a More Restful Night  
 Reduces  
 Swelling of  
 Nasal  
 Passages  
 1/2 FL OZ (15mL)

## AFRIN NODRIP ALLERGY SINUS NIGHT

oxymetazoline hydrochloride spray

### Product Information

|                                |                |                           |                |
|--------------------------------|----------------|---------------------------|----------------|
| <b>Product Type</b>            | HUMAN OTC DRUG | <b>Item Code (Source)</b> | NDC:11523-0034 |
| <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 | 0.5 mg in 1 mL |

### Inactive Ingredients

| Ingredient Name  | Strength |
|--|----------|
| <b>CARBOXYMETHYLCELLULOSE SODIUM</b> (UNII: K679OBS311)    |          |
| <b>SODIUM PHOSPHATE, MONOBASIC</b> (UNII: 3980JIH2SW)      |          |
| <b>WATER</b> (UNII: 059QF0KO0R)                            |          |
| <b>EDETATE DISODIUM</b> (UNII: 7FLD91C86K)                 |          |
| <b>GLYCERIN</b> (UNII: PDC6A3C0OX)                         |          |
| <b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7)            |          |
| <b>POVIDONE</b> (UNII: FZ989GH94E)                         |          |
| <b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ0SDW1A) |          |
| <b>BENZYL ALCOHOL</b> (UNII: LKG8494WBH)                   |          |
| <b>MICROCRYSTALLINE CELLULOSE</b> (UNII: OP1R32D61U)       |          |
| <b>HIBISCUS BIFURCATUS WHOLE</b> (UNII: 60F5JKG79P)        |          |

### Packaging

| # | Item Code        | Package Description                                  | Marketing Start Date | Marketing End Date |
|---|------------------|--|----------------------|--------------------|
| 1 | NDC:11523-0034-1 | 15 mL in 1 BOTTLE; Type 0: Not a Combination Product | 11/09/2020           |                    |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|--------------------|--|----------------------|--------------------|
| OTC Monograph Drug | M012                                     | 11/09/2020           |                    |

**Labeler** - Bayer HealthCare LLC. (112117283)

Revised: 12/2025

Bayer HealthCare LLC.