

AFRIN ORIGINAL- oxymetazoline hydrochloride spray
Bayer HealthCare LLC.

Afrin®

Original

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
- 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.

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, edetate disodium, polyethylene glycol, povidone, propylene glycol, purified water, sodium phosphate dibasic, sodium phosphate monobasic

Questions or comments?

Call 1-800-317-2165

PRINCIPAL DISPLAY PANEL - 15 mL Bottle Carton

Afrin®

Oxymetazoline HCl Nasal Solution-Nasal Decongestant

Original

NASAL SPRAY

Fast, Powerful

Congestion

Relief

Colds • Allergies

up to 12

Hour

Relief

Maximum

Strength

#1 Doctor & Pharmacist

Recommended Brand

1/2 FL OZ (15 mL)

2 4 3

Child Resistant Cap
(see side panel)

Oxymetazoline HCl Nasal Solution-Nasal Decongestant
Afrin®
Original
NASAL SPRAY
SAFETY SEALED: Do not use if seal imprinted "Afrin® Security Seal" is broken or missing.

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
temporarily relieves sinus congestion and pressure
shrinks swollen nasal membranes so you can breathe more freely

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Ask a doctor before use if you have
■ heart disease ■ high blood pressure
■ thyroid disease ■ diabetes
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Directions
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■ children under 6 years of age: ask a doctor
To Use: Push firmly down on cap and turn counter clockwise. To spray, squeeze bottle quickly and firmly. Do not tilt head backward while spraying. Wipe nozzle clean after use. Secure cap after use.

Afrin® Original Nasal Spray
*It unblocks your nose in seconds, providing 12 hours of nasal congestion relief without drowsiness, so you can breathe more freely all day or all night.

Oxymetazoline HCl Nasal Solution-Nasal Decongestant
Afrin®
Original
NASAL SPRAY

• Relieves Nasal Congestion & Sinus Congestion/Pressure

UNBLOCKS IN 1 SECONDS* LASTS 12 HRS

No1 DOCTOR AND PHARMACIST* RECOMMENDED NASAL SPRAY BRAND

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Whippany, NJ 07981
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www.patents.livewell.bayer.com

Bayer ^{Pharmacy Times survey}

Drug Facts (continued)
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How to use:
Push firmly down on cap and turn counterclockwise. Secure cap after use.

3 1 0 0 0 9 5 4

LOT
EXP

0 4 1 1 0 0 8 1 1 2 3 3

2 4 3

AFRIN ORIGINAL

oxymetazoline hydrochloride spray

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:11523-1167

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 100 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM PHOSPHATE, DIBASIC (UNII: GR686LBA74)	
SODIUM PHOSPHATE, MONOBASIC (UNII: 3980JIH2SW)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POVIDONE (UNII: FZ989GH94E)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11523-1167-2	2 in 1 CARTON	07/14/1975	
1		21 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product		
2	NDC:11523-1167-3	1 in 1 CARTON	07/14/1975	
2		20 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product		
3	NDC:11523-1167-4	3 in 1 CARTON	07/14/1975	12/31/2021
3		30 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product		
4	NDC:11523-1167-5	1 in 1 CARTON	07/14/1975	
4		30 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product		
5	NDC:11523-1167-6	1 in 1 CARTON	07/14/1975	
5		15 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product		
6	NDC:11523-1167-1	6 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/16/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	07/14/1975	

Labeler - Bayer HealthCare LLC. (112117283)

Revised: 12/2025

Bayer HealthCare LLC.