

AFRIN NO DRIP PUMP MIST BUNDLE PACK- oxymetazoline hydrochloride
Bayer HealthCare LLC.

Afrin ® No Drip Original/Night Pump Mist Bundle Pack

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.

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.

Other information

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

Inactive ingredients

No Drip Original

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

No Drip Night

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?

Questions or comments? 1-800-317-2165

PRINCIPAL DISPLAY PANEL - Bundle Pack Carton



Drug Facts	
Active ingredient	Purpose
Oxymetazoline hydrochloride 0.05%	Nasal decongestant
Uses	
<ul style="list-style-type: none"> temporarily relieves nasal congestion due to: <ul style="list-style-type: none"> 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: <ul style="list-style-type: none"> heart disease high blood pressure thyroid disease diabetes 	
When using this product: <ul style="list-style-type: none"> 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	
<ul style="list-style-type: none"> 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	
<ul style="list-style-type: none"> store between 20° to 25°C (68° to 77°F) retain carton for future reference on full labeling 	
Inactive ingredients	
No Drip Original: benzalkonium chloride solution, benzyl alcohol, edetate disodium, flunar, microcrystalline cellulose and carboxymethylcellulose sodium, polyethylene glycol, povidone, purified water, sodium phosphate dibasic, sodium phosphate monobasic	

BUNDLE PACK 2 X 15 mL Bottles

Oxymetazoline HCl Nasal Solution-Nasal Decongestant

Afrin®

No Drip

Original

NASAL PUMP MIST

- Relieves Nasal Congestion & Sinus Congestion/Pressure

Unblocks in

SECONDS*

Lasts 12 HRS

1/2 FL OZ (15 mL)

Oxymetazoline HCl Nasal Solution-Nasal Decongestant

Afrin®

No Drip

Night

NASAL PUMP MIST

- Nasal Congestion Relief for a More Restful Night
- Chamomile Scent

Unblocks in

SECONDS*

Lasts 12 HRS

1/2 FL OZ (15 mL)

AFRIN NO DRIP PUMP MIST BUNDLE PACK				
oxymetazoline hydrochloride kit				
Product Information				
Product Type		HUMAN OTC DRUG	Item Code (Source)	
			NDC:11523-0147	
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11523-0147-1	1 in 1 CARTON; Type 0: Not a Combination Product	04/19/2024	
Quantity of Parts				
Part #	Package Quantity		Total Product Quantity	
Part 1	1 BOTTLE, SPRAY		15 mL	
Part 2	1 BOTTLE, PUMP		15 mL	
Part 1 of 2				
AFRIN NO DRIP ORIGINAL PUMP MIST				
oxymetazoline hydrochloride spray, metered				
Product Information				
Item Code (Source)		NDC:11523-3142		
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
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311)	
WATER (UNII: 059QF0KO0R)	
SODIUM PHOSPHATE, MONOBASIC (UNII: 3980JIH2SW)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
POVIDONE (UNII: FZ989GH94E)	
SODIUM PHOSPHATE, DIBASIC (UNII: GR686LBA74)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	

Product Characteristics

Color		Score	
Shape		Size	
Flavor	LEMON	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		15 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	08/31/1999	

Part 2 of 2

AFRIN NO DRIP NIGHT CHAMOMILE

oxymetazoline hydrochloride spray, metered

Product Information

Item Code (Source)	NDC:11523-1160
Route of Administration	NASAL

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
OXYMETAZOLINE HYDROCHLORIDE (UNII: K89MJ0S5VY) (OXYMETAZOLINE - UNII:8VLN5B44ZY)	OXYMETAZOLINE HYDROCHLORIDE	0.5 mg in 1 mL

Inactive Ingredients	
Ingredient Name	Strength
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
WATER (UNII: 059QF0KO0R)	
SODIUM PHOSPHATE, MONOBASIC (UNII: 3980JIH2SW)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
POVIDONE (UNII: FZ989GH94E)	
SODIUM PHOSPHATE, DIBASIC (UNII: GR686LBA74)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		15 mL in 1 BOTTLE, PUMP; 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	08/31/1999	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	04/19/2024	

Labeler - Bayer HealthCare LLC. (112117283)