

**MUCINEX NASAL CONGESTION AND ALLERGY RELIEF- fluticasone propionate spray, metered
RB Health (US) LLC**

MUCINEX® Nasal Congestion + Allergy Relief

Drug Facts

Fluticasone propionate (glucocorticoid)¹ 50 mcg

Fluticasone propionate (glucocorticoid)¹ 50 mcg Allergy symptom reliever

Uses

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Temporarily relieves these symptoms of hay fever or other upper respiratory allergies:

- nasal congestion ■ itchy nose ■ runny nose ■ itchy, watery eyes ■ sneezing

Warnings

Only for use in the nose. Do not spray into your eyes or mouth.

Do not use

- in children under 4 years of age
- to treat asthma
- if you have an injury or surgery to your nose that is not fully healed
- if you have ever had an allergic reaction to this product or any of the ingredients

Ask a doctor before use if you have or had glaucoma or cataracts

Ask a doctor or pharmacist before use if you are taking

- medicine for HIV infection (such as ritonavir)
- a steroid medicine for asthma, allergies or skin rash
- ketoconazole pills (medicine for fungal infection)

When using this product

- the growth rate of some children may be slower
- stinging or sneezing may occur for a few seconds right after use
- do not share this bottle with anyone else as this may spread germs
- remember to tell your doctor about all the medicines you take, including this one

Stop use and ask a doctor if

- you have, or come into contact with someone who has, chicken pox, measles or tuberculosis
- your symptoms do not get better within 7 days of starting use or you get new symptoms such as severe

facial pain or thick nasal discharge. You may have something more than allergies, such as an infection.

- you get a constant whistling sound from your nose. This may be a sign of damage inside your nose.
- you get an allergic reaction to this product. Seek medical help right away.
- you get new changes to your vision that develop after starting this product
- you have severe or frequent nosebleeds

If pregnant or breast-feeding, ask a health professional before use. Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center right away.

Directions

- read the Quick Start Guide for how to:
- prime the bottle ■ use the spray
- clean the spray nozzle
- shake gently before each use
- use this product only once a day
- do not use more than directed

ADULTS AND CHILDREN 12 YEARS

OF AGE AND OLDER

- Week 1 - use 2 sprays in each nostril once daily
- Week 2 through 6 months - use 1 or 2 sprays in each nostril once daily, as needed to treat your symptoms
- After 6 months of daily use - ask your doctor if you can keep using

CHILDREN 4 TO 11 YEARS OF AGE

- the growth rate of some children may be slower while using this product. **Children should use for the shortest amount of time necessary to achieve symptom relief. Talk to your child's doctor if your child needs to use the spray for longer than two months a year.**
- an adult should supervise use
- use 1 spray in each nostril once daily

CHILDREN UNDER 4 YEARS OF AGE

- do not use

IMPORTANT: Read Drug Facts on package and enclosed materials for important information.

Children 4-11: 1 spray per nostril per day.

Talk to a doctor if your child needs to use for longer than two months a year. Keep out of reach of children.

Only for use in the nose, do not spray into eyes. Store between 4°-30°C (39°-86°F). Shake gently before each use.

Other information

- you may start to feel relief the first day and full effect after several days of regular, once-a-day use
- store between 4° and 30°C (39° and 86°F)
- keep this package and enclosed materials. They contain important additional information.

Inactive ingredients

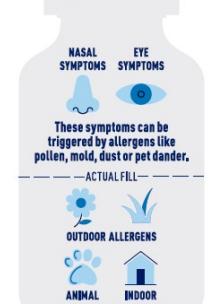
benzalkonium chloride, dextrose anhydrous, microcrystalline cellulose and carboxymethylcellulose sodium, phenylethyl alcohol, polysorbate 80, purified water

**Questions? 1-866-MUCINEX
(1-866-682-4639)**



What problems can Mucinex Nasal Congestion + Allergy Relief Nasal Spray help with?

Mucinex Nasal Congestion + Allergy Relief helps relieve a broad range of uncomfortable symptoms like congestion and itchy eyes.



ACTUAL SIZE
Package Contains 1 Bottle

TAMPER-EVIDENT features for your protection. The product is packaged in a sealed carton. Under the cap and nozzle, each bottle has an aluminum seal around bottle neck. Do not use if any of these features are torn or damaged.

IMPORTANT: Children 4-11: do not use for more than 2 months a year.
Be sure to read the Quick Start Guide and Question & Answer Leaflet inside package.

Drug Facts

Active ingredient (in each spray) Fluticasone propionate (glucocorticoid)* 50 mcg **Purpose** Allergy symptom reliever *read the Question & Answer Leaflet

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Drug Facts (continued)

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Inactive ingredients

benzalkonium chloride, dextrose anhydrous, microcrystalline cellulose and carboxymethylcellulose sodium, phenylethyl alcohol, polysorbate 80, purified water

**Questions? 1-866-MUCINEX
(1-866-682-4639)**

MUCINEX[®]

NASAL CONGESTION + ALLERGY RELIEF

NDC72854-227-60

Fluticasone Propionate Nasal Spray
50 mcg Per Spray Allergy Symptom Reliever
(Glucocorticoid)



IMPORTANT: Read Drug Facts on package and enclosed materials for important information.
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Dist. by: RB Health (US)
Parsippany, NJ 07054-0224
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Made in India
3344971 111025

Unvarnished Zone
(dotted line not for printing)
30 x 12 mm

MUCINEX NASAL CONGESTION AND ALLERGY RELIEF

fluticasone propionate spray, metered

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72854-227
Route of Administration	NASAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
FLUTICASONE PROPIONATE (UNII: O2GMZ0LF5W) (FLUTICASONE - UNII:CUT2W21N7U)	FLUTICASONE PROPIONATE	50 ug

Inactive Ingredients

Ingredient Name	Strength
PHENYLETHYL ALCOHOL (UNII: ML9LGA7468)	
POLYSORBATE 80 (UNII: 60ZP39ZG8H)	
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311)	
DEXTROSE, UNSPECIFIED FORM (UNII: IY9XDZ35W2)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
WATER (UNII: 059QF0KO0R)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72854-227-60	1 in 1 CARTON	07/01/2025	
1		60 in 1 BOTTLE; 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
ANDA	ANDA217088	07/01/2025	

Labeler - RB Health (US) LLC (081049410)

Revised: 1/2026

RB Health (US) LLC