

FLUTICASONE PROPIONATE- fluticasone propionate spray, metered **Sam's West Inc.**

Drug Facts

Active ingredient (in each spray)

Fluticasone propionate (glucocorticoid) 50 mcg.

Purpose

Allergy symptom reliever

Uses

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

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

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**

Other information

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

Inactive ingredients

0.02% w/w benzalkonium chloride, dextrose, microcrystalline cellulose and carboxymethylcellulose sodium, 0.25% w/w phenylethyl alcohol, polysorbate 80, purified water

Questions or comments?

call toll free **1-800-706-5575**, weekdays, 8:30am – 5:00pm Eastern Standard Time

Principal Display Panel - Carton

CARTON LABEL - PRINCIPAL DISPLAY PANEL - 50 mcg per spray

Aller-Nose Nasal Spray

Fluticasone Propionate Nasal Spray, USP

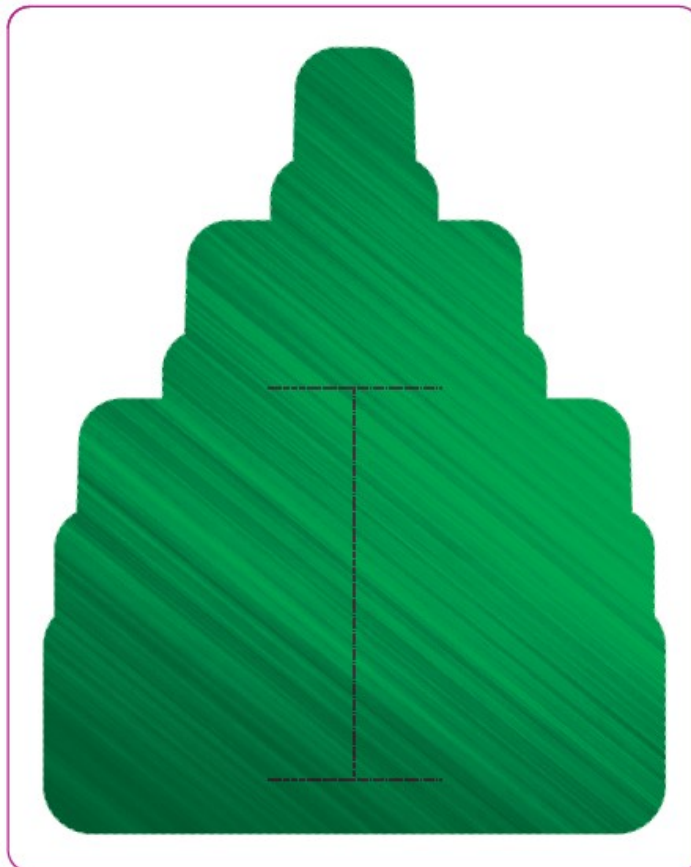
Apotex Corp. NDC 68196-210-01


6 bottles (144 sprays each)

Allergy Symptom Reliever Nasal Spray

24 Hour Relief of:

- Itchy, Watery Eyes
- Nasal Congestion
- Runny Nose
- Itchy Nose
- Sneezing



Member's Mark 

full prescription strength • non-drowsy
Aller-Nose Nasal Spray
Fluticasone Propionate, USP
50 mcg per spray
allergy symptom reliever (Glucocorticoid)**

What problems can Fluticasone Propionate Nasal Spray, USP help with?
Fluticasone Propionate Nasal Spray, USP helps relieve a broad range of uncomfortable symptoms like congestion and itchy eyes.

Nasal symptoms
Eye symptoms
These symptoms can be triggered by allergens like pollen, mold, dust or pet dander.
• Outdoor allergens • Animal allergens • Indoor allergens

**Fluticasone propionate is a steroid and when taken as a glucocorticoid.

Get Further Prescribed Information About Use in Children

Drug Facts	Drug Facts (continued)	Drug Facts (continued)
Active ingredient(s) (for each spray) Fluticasone propionate NDC 68196-210-01 Allergy symptoms relief	Purpose Allergy symptoms relief	CAUTIONS (continued) • Do not use if you are allergic to any of the ingredients listed above or to any other ingredients in this product. Ask your pharmacist for more information. • Do not use if you have had a recent eye infection or if you have had a recent eye surgery. • Do not use if you have had a recent eye injury or if you have had a recent eye operation. • Do not use if you have had a recent eye infection or if you have had a recent eye surgery. • Do not use if you have had a recent eye injury or if you have had a recent eye operation.
Uses Temporarily relieves these signs and symptoms of hay fever or other upper respiratory allergies: • itchy nose • itchy eyes • itchy throat • itchy ears	Warnings • Do not spray into your eyes or mouth. • Do not use if you are allergic to any of the ingredients listed above or to any other ingredients in this product. Ask your pharmacist for more information. • Do not use if you have had a recent eye infection or if you have had a recent eye surgery. • Do not use if you have had a recent eye injury or if you have had a recent eye operation.	Directions • Use as directed. • Do not use if you are allergic to any of the ingredients listed above or to any other ingredients in this product. Ask your pharmacist for more information. • Do not use if you have had a recent eye infection or if you have had a recent eye surgery. • Do not use if you have had a recent eye injury or if you have had a recent eye operation.
Directions • Use as directed. • Do not use if you are allergic to any of the ingredients listed above or to any other ingredients in this product. Ask your pharmacist for more information. • Do not use if you have had a recent eye infection or if you have had a recent eye surgery. • Do not use if you have had a recent eye injury or if you have had a recent eye operation.	Other information • This product is not intended to be used for the treatment of any other condition. • This product is not intended to be used for the treatment of any other condition. • This product is not intended to be used for the treatment of any other condition.	Other information • This product is not intended to be used for the treatment of any other condition. • This product is not intended to be used for the treatment of any other condition. • This product is not intended to be used for the treatment of any other condition.

TAMPER EVIDENT: ON ANY ONE OF PACKAGE'S PANEL, THERE WILL BE A SEAL AND/OR THE PUNCH HOLE LABELS
"SEAL FOR YOUR PROTECTION" IS BROKEN OR MISSING.

Questions or Comments?
We would love to hear about your Member's Mark experience! Call us at: 1-800-706-5575 (toll-free).
8:30am – 5:00pm EST Monday – Friday

2D FPO

DISTIBUTED BY SANI'S WEST, INC.
CENTREVILLE, OH 43016
PRODUCT OF CANADA

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Principal Display Panel - Bottle

BOTTLE LABEL - PRINCIPAL DISPLAY PANEL - 50 mcg per spray

Aller-Nose Nasal Spray

Fluticasone Propionate Nasal Spray, USP

Apotex Corp. NDC 68196-210-02

144 sprays 0.62 fl oz (18.2 mL)

Allergy Symptom Reliever Nasal Spray



FLUTICASONE PROPIONATE

fluticasone propionate spray, metered

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68196-210
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
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K679OBS311)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
DEXTROSE, UNSPECIFIED FORM (UNII: IY9XDZ35W2)	
PHENYLETHYL ALCOHOL (UNII: ML9LGA7468)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68196-210-02	144 in 1 BOTTLE, SPRAY; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)	01/27/2021	

2	NDC:68196-210-01	6 in 1 CARTON	01/27/2021	
2		6 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
ANDA	ANDA208150	01/27/2021	

Labeler - Sam's West Inc. (051957769)

Registrant - Apotex Inc. (209429182)

Revised: 6/2024

Sam's West Inc.