

**FLUTICASONE PROPIONATE- fluticasone propionate spray, metered**  
**Apotex Corp.**

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**Drug Facts**

***Active ingredient (in each spray)***

Fluticasone propionate (glucocorticoid) \*50 mcg.

\*read the Question & Answer Leaflet

***Purpose***

Allergy symptom reliever

***Uses***

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**

### ***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 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 (single)**

**CARTON LABEL - PRINCIPAL DISPLAY PANEL - 50 mcg per spray**

**Apothecare Plus** NDC 60505-6205-3

**Allergy Nasal Spray**

**Fluticasone Propionate 50 mcg per spray**

**Allergy Symptom Reliever**

**(Glucocorticoid)**

**120 sprays**

**24 Hour Relief of:**

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



**Principal Display Panel - Bottle**

**BOTTLE LABEL - PRINCIPAL DISPLAY PANEL - 50 mcg per spray**

**Apothecare Plus NDC 60505-6205-3**

**Allergy Nasal Spray**

**Fluticasone Propionate 50 mcg**

**120 sprays**

**Allergy Symptom Reliever**

**(Glucocorticoid)**

0.54 fl oz (15.8 mL)



**Fluticasone Propionate  
Nasal Spray, USP**

50 mcg Per Spray  
Allergy Symptom Reliever  
(Glucocorticoid) **120 sprays**

**IMPORTANT:** Read Drug Facts label 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°C and 30°C (39°F and 86°F) Shake gently before each use.

Distributed by:  
Apotex Corp.  
Weston, Florida 33326  
MADE IN CANADA

408827

## FLUTICASONE PROPIONATE

fluticasone propionate spray, metered

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:60505-6205
<b>Route of Administration</b>	NASAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>FLUTICASONE PROPIONATE</b> (UNII: O2GMZ0LF5W) (Fluticasone - UNII:CUT2W21N7U)	FLUTICASONE PROPIONATE	50 ug

### Inactive Ingredients

Ingredient Name	Strength
<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7)	
<b>CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED</b> (UNII: K679OBS311)	
<b>MICROCRYSTALLINE CELLULOSE</b> (UNII: OP1R32D61U)	
<b>DEXTROSE, UNSPECIFIED FORM</b> (UNII: IY9XDZ35W2)	
<b>PHENYLETHYL ALCOHOL</b> (UNII: ML9LGA7468)	
<b>POLYSORBATE 80</b> (UNII: 6OZP39ZG8H)	
<b>WATER</b> (UNII: 059QF0KO0R)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:60505-6205-3	1 in 1 CARTON	01/11/2019	
1		120 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	01/11/2019	

**Labeler** - Apotex Corp. (845263701)

**Registrant** - Apotex Inc. (243805095)

### Establishment

Name	Address	ID/FEI	Business Operations
Apotex Inc.		255092496	analysis(60505-6205) , manufacture(60505-6205)

Revised: 9/2025

Apotex Corp.