

**NASAL ALLERGY- triamcinolone acetonide spray, metered**  
**Rugby Laboratories, Inc.**

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***Drug Facts***

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

Triamcinolone acetonide 55 mcg (glucocorticoid)

***Purpose***

Allergy symptom reliever

***Uses***

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

- nasal congestion
- runny nose
- sneezing
- itchy nose

***Warnings***

**Do not use**

- In children under 2 years of age
- If you have ever had an allergic reaction to any of the ingredients

**Ask a doctor before use if you**

- have had recent nose ulcers or nose surgery
- have had a nose injury that has not healed
- are using a steroid medicine for asthma, allergies or skin rash
- have an eye infection
- have or had glaucoma or cataracts

**When using this product**

- the growth rate of some children may be slower
- some symptoms may get better on the first day of treatment. It may take up to one week of daily use to feel the most symptom relief.
- 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 in contact with someone who has, chickenpox, measles or tuberculosis

- you have or develop symptoms of an infection such as persistent fever
- you have any change in vision
- 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 (1800-222-1222) right away.

**Direction**

**Read insert (\*inside package) on how to:**

- **get a new bottle ready (primed before first use**
- **prime bottle again if not used for more than 2 weeks**
- **use the spray nozzle**

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**ADULTS AND CHILDREN 12 YEARS OF AGE AND OLDER**

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**adults and children 12 years of age and older**

- once daily, spray 2 times into each nostril while sniffing gently
- once your allergy symptoms improve, reduce to 1 spray in each nostril per day

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**CHILDREN 2 TO UNDER 12 YEARS OF AGE**

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- the growth rate of some children may be slower while using this product. Talk to your child's doctor if your child to use the spray for longer than two months a year

**children 6 to under 12 years of age**

- an adult should supervise use
- once daily, spray 1 time into each nostril while sniffing gently
- if allergy symptoms do not improve, increase to 2 sprays in each nostril per day. Once allergy symptoms improve, reduce to 1 spray in each nostril per day.

**children 2 to under 6 years of age**

- an adult should supervise use
- once daily, spray 1 time into each nostril while sniffing gently

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**CHILDREN UNDER 2 YEARS OF AGE**

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- do not use
- do not use more than directed
- if you forget a dose, do not double the next dose

- do not spray into the eyes or mouth
- if allergy symptoms do not improve after one weeks, stop using and talk to a doctor
- do not use for the common cold
- shake well before each use

### ***Other information***

- **TAMPER EVIDENT: DO NOT USE IF SEALS ON CARTON ARE BROKEN OR MISSING**
- keep package and insert. They contain important information.
- Store upright 20°C to 25°C (68°F to 77°F)

### ***Inactive ingredients***

benzalkonium chloride solution (50% W/V) carboxymethylcellulose sodium, dextrose monohydrate, edetate disodium dihydrate, hydrochloric acid, microcrystalline cellulose, polysorbate 80 purified water, sodium hydroxide

### ***Questions or comments?***

Call **1-877-753-3935** Monday-Friday 9AM-5PM EST

### **Principal Display Panel**

compare to the active ingredient in Nasacort® Allergy 24 HR\*

nasal allergy spray, USP

triamcinolone

acetonide (glucocorticoid)\*\*

55 mcg per spray

nasal allergy

symptom reliever

original prescription strength

24-hour relief of:

- nasal congestion
- runny nose
- itchy nose
- sneezing

non-drowsy

scent-free

alcohol-free

|              |                |                    |               |
|--------------|----------------|--------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:0536-1457 |
|--------------|----------------|--------------------|---------------|

|                                                                                        |                  |                                                                                              |                      |                    |
|----------------------------------------------------------------------------------------|------------------|----------------------------------------------------------------------------------------------|----------------------|--------------------|
| Route of Administration                                                                |                  | NASAL                                                                                        |                      |                    |
|                                                                                        |                  |                                                                                              |                      |                    |
| Active Ingredient/Active Moiety                                                        |                  |                                                                                              |                      |                    |
| Ingredient Name                                                                        |                  | Basis of Strength                                                                            | Strength             |                    |
| TRIAMCINOLONE ACETONIDE (UNII: F446C597KA) (Triamcinolone Acetonide - UNII:F446C597KA) |                  | TRIAMCINOLONE ACETONIDE                                                                      | 55 ug                |                    |
|                                                                                        |                  |                                                                                              |                      |                    |
| Inactive Ingredients                                                                   |                  |                                                                                              |                      |                    |
| Ingredient Name                                                                        |                  |                                                                                              | Strength             |                    |
| BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)                                               |                  |                                                                                              |                      |                    |
| DEXTROSE MONOHYDRATE (UNII: LX22YL083G)                                                |                  |                                                                                              |                      |                    |
| EDETATE DISODIUM (UNII: 7FLD91C86K)                                                    |                  |                                                                                              |                      |                    |
| HYDROCHLORIC ACID (UNII: QTT17582CB)                                                   |                  |                                                                                              |                      |                    |
| SODIUM HYDROXIDE (UNII: 55X04QC32I)                                                    |                  |                                                                                              |                      |                    |
| WATER (UNII: 059QF0KO0R)                                                               |                  |                                                                                              |                      |                    |
| MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)                                          |                  |                                                                                              |                      |                    |
| POLYSORBATE 80 (UNII: 6OZP39ZG8H)                                                      |                  |                                                                                              |                      |                    |
| CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K679OBS311)                          |                  |                                                                                              |                      |                    |
|                                                                                        |                  |                                                                                              |                      |                    |
| Packaging                                                                              |                  |                                                                                              |                      |                    |
| #                                                                                      | Item Code        | Package Description                                                                          | Marketing Start Date | Marketing End Date |
| 1                                                                                      | NDC:0536-1457-69 | 1 in 1 CARTON                                                                                | 11/11/2025           |                    |
| 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                                                                                   |                  | ANDA214615                                                                                   | 11/11/2025           |                    |

**Labeler** - Rugby Laboratories, Inc. (079246066)

Revised: 11/2025

Rugby Laboratories, Inc.