# FLONASE NIGHTTIME ALLERGY RELIEF- triprolidine hcl tablet Haleon US Holdings LLC

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### **Drug Facts**

### Active ingredient (in each tablet)

Triprolidine HCl 2.5 mg

### **Purpose**

#### Uses

- temporarily relieves these symptoms due to hay fever (allergic rhinitis) or other upper respiratory allergies:
- 1. runny nose
- 2. sneezing
- 3. itching of the nose or throat
- 4. itchy, watery eyes

### Warnings

### Ask a doctor before use if you have

- glaucoma
- difficulty in urination due to enlargement of the prostate gland
- a breathing problem such as emphysema or chronic bronchitis

## Ask a doctor before use if you are taking sedatives or tranquilizers.

# When using this product

- do not exceed recommended dosage
- excitability may occur, especially in children
- drowsiness may occur
- alcohol, sedatives, and tranquilizers may increase drowsiness
- avoid alcoholic beverages while taking this product
- be careful when driving a motor vehicle or operating machinery

# Stop use and ask a doctor if new symptoms occur.

# 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.

Quick medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

#### **Directions**

- do not use more than directed
- adults and children 12 years of age and over:1 tablet (2.5 mg) every 4 to 6 hours; not to exceed 4 tablets (10 mg) in 24 hours or as directed by a doctor
- children under 12 years of age: do not use

#### Other information

store at controlled room temperature 20-25°C (68-77°F)

### Inactive ingredients

hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, pregelatinized starch, titanium dioxide

#### Questions or comments?

1-844-356-6273

#### Additional Information

#### **KEEP CARTON FOR COMPLETE INFORMATION**

Do not use if blister is broken or damaged.

\*Dosage is 1 tablet every 4 to 6 hours

Distributed by:

#### **GSK Consumer Healthcare**

Warren, NJ 07059

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### **Principal Display Panel**

**FLONASE** 

**NIGHTTIME ALLERGY RELIEF** 

**NEW INGREDIENT** 

**Triprolidine HCI 2.5 mg - Antihistamine** 

UP TO 6 HOURS\* RELIEF FROM

Sneezing

- Runny nose
- Itchy throat
- Itchy, watery eyes

#### **36**TABLETS

080320 - Front Carton



## FLONASE NIGHTTIME ALLERGY RELIEF

triprolidine hcl tablet

| Product Information     |                |                    |               |  |
|-------------------------|----------------|--------------------|---------------|--|
| Product Type            | HUMAN OTC DRUG | Item Code (Source) | NDC:0135-1329 |  |
| Route of Administration | ORAL           |                    |               |  |

| Active Ingredient/Active Moiety   |                               |          |
|---|-------------------------------|----------|
| Ingredient Name   | <b>Basis of Strength</b>      | Strength |
| TRIPROLIDINE HYDROCHLORIDE (UNII: YAN7R5L890) (TRIPROLIDINE - UNII: 2L8T9S52QM) | TRIPROLIDINE<br>HYDROCHLORIDE | 2.5 mg   |
|   |                               |          |

# **Inactive Ingredients**

| Ingredient Name                                     | Strength |
|---|----------|
| HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)        |          |
| MAGNESIUM STEARATE (UNII: 70097M6I30)               |          |
| MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)       |          |
| POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A) |          |
| STARCH, CORN (UNII: O8232NY3SJ)                     |          |
| TITANIUM DIOXIDE (UNII: 15FIX9V2JP)                 |          |

| Product Characteristics |       |              |          |
|-------------------------|-------|--------------|----------|
| Color                   | white | Score        | no score |
| Shape                   | ROUND | Size         | 7mm      |
| Flavor                  |       | Imprint Code | FN       |
| Contains                |       |              |          |

| Packaging |           |   |                         |                       |
|-----------|-----------|---|-------------------------|-----------------------|
| # Item    | Code      | Package Description                                     | Marketing Start<br>Date | Marketing End<br>Date |
| 1 NDC:01  | 135-<br>1 | 1 in 1 CARTON   | 12/15/2022              |                       |
| 1         |           | 36 in 1 BLISTER PACK; Type 0: Not a Combination Product |                         |                       |

| Marketing Information |   |                         |                       |
|-----------------------|---|-------------------------|-----------------------|
| Marketing<br>Category | Application Number or Monograph<br>Citation | Marketing Start<br>Date | Marketing End<br>Date |
| OTC Monograph Drug    | M012  | 12/15/2022              |                       |
|                       |   |                         |                       |

# Labeler - Haleon US Holdings LLC (079944263)

Revised: 3/2024 Haleon US Holdings LLC