# STONARHINI- chlorpheniramine maleate, phenylephrine tablet Sato Pharmaceutical Co., Ltd.

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#### Stonarhini

**Active ingredients** (In each tablet)

Chlorpheniramine Maleate 2 mg Phenylephrine HCl 5 mg

#### Purpose

Chlorpheniramine Maleate Antihistamine Phenylephrine HCl Nasal decongestant

#### Uses

Temporarily relieves these symptoms due to common cold, hay fever or other upper respiratory allergies (allergic rhinitis).

■ sneezing ■ runny nose ■ itchy, watery eyes

■ nasal congestion, stuffy nose
■ itchy throat

### Warnings

Enter section text here

#### Do not use this product

■ if you are now taking a prescription monoamine oxidase inhibitor (MAOI) (certain drugs for depression, psychiatric or emotional conditions, or Parkinson's disease) or for 2 weeks after stopping the MAOI drug. If you do not know if your prescription contains an MAOI, ask a doctor or pharmacist before taking this priduct.

## Ask a doctor before use if you have

- a breathing problem such as emphysema or chronic bronchitis
- glaucoma heart disease thyroid disease
- diabetes high blood pressure
- ■difficulty in urination due to enlargement of the prostate

# Ask a doctor or pharmacist before use

■ if you are taking sedatives or tranquilizers

# When using this product

- do not exceed recommended dosage. If nervousness, dizziness, or sleeplessness occur, discontinue use and consult a doctor.
- you may get drowsy.
- may cause excitability, especially in children.
- alcohol, sedatives and tranquilizers may increase the drowsiness effect.
- avoid alcoholic beverages.
- use caution when driving a motor vehicle or operating machinery.

## Stop use and ask a doctor if

- symptoms do not improve within seven days
- symptoms are accompanied by fever

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. Prompt attention is critical even if you do not notice any signs or symptoms.

#### **Directions**

adults and children 12 years and over: 2 tablets every 4 hours, not to exceed 12 tablets in 24 hours

children under 12 years of age: ask a doctor

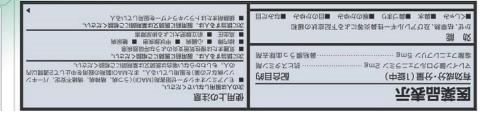
**Inactive ingredients** aspartame, colloidal silicone dioxide, FD&C Red No. 40 aluminum Lake, flavor, hydroxypropyl cellulose, magnesium stearate, mannitol.

**Product Information** HUMAN OTC DRUG NDC:49873-109 **Product Type** Item Code (Source) **Route of Administration ORAL** 

chlorpheniramine maleate, phenylephrine tablet

NDC 48873-109-01

# **STONARHINI**





量用· 去用

SATABLETS(錠)

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Tablets individually blister sealed. If seal is broken, do not use.
 経剤を包装しているソールが飲れている場合 は服用しないでください。

ANTIHISTAMINE NASAL DECONGESTANT STONARHINI

ナースクーア・発展業分の際、FD&C 赤色40号 アルニニウムレーキース・大力では、プロインローク・カー・エースト・ファットのできない。 代気の他の成分

Drug Facts (continued) Stop use and ask a doctor if
symptoms do not improve within seven days
symptoms are accompanied by fever

Reports of serious side effects associate with use of this product can be sent to: SATO PHARMACEUTICAL, INC. 20695 S. Western Ave., Sulte 240, Torrance, CA 90501

PUSH

Active ingredients (in each tablet)
Chlorpheniramine Maleate 2 mg · · · · Phenylephrine HCl 5 mg · · · · · · ·

Manufactured by SATO PHARMACEUTICAL CO., LTD. TOKYO, JAPAN

| Active Ingredient/Active Moiety   |                                |          |  |  |
|---|--------------------------------|----------|--|--|
| Ingredient Name   | Basis of Strength              | Strength |  |  |
| CHLORPHENIRAMINE MALEATE (UNII: V1Q0090J9Z) (CHLORPHENIRAMINE - UNII: 3U6I01965U) | CHLORPHENIRAMINE<br>MALEATE    | 2 mg     |  |  |
| PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII: 1WS297W6MV) | PHENYLEPHRINE<br>HYDROCHLORIDE | 5 mg     |  |  |

| Inactive Ingredients                                 |          |  |  |  |
|--|----------|--|--|--|
| Ingredient Name                                      | Strength |  |  |  |
| ASPARTAME (UNII: Z0H242BBR1)                         |          |  |  |  |
| SILICON DIOXIDE (UNII: ETJ7Z6XBU4)                   |          |  |  |  |
| FD&C RED NO. 40 (UNII: WZB9127XOA)                   |          |  |  |  |
| HYDROXYPROPYL CELLULOSE (TYPE EL) (UNII: 8VAB711C5E) |          |  |  |  |
| MAGNESIUM STEARATE (UNII: 70097M6I30)                |          |  |  |  |
| MANNITOL (UNII: 30WL53L36A)                          |          |  |  |  |
| ALUMINUM OXIDE (UNII: LMI26O6933)                    |          |  |  |  |

| Product Characteristics |                |              |          |  |
|-------------------------|----------------|--------------|----------|--|
| Color                   | red (pale red) | Score        | no score |  |
| Shape                   | ROUND          | Size         | 10mm     |  |
| Flavor                  | MENTHOL        | Imprint Code | S10      |  |
| Contains                |                |              |          |  |

| F | Packaging            |  |                         |                       |  |  |
|---|----------------------|--|-------------------------|-----------------------|--|--|
| # | tem Code             | Package Description                                    | Marketing Start<br>Date | Marketing End<br>Date |  |  |
| 1 | NDC:49873-109-<br>01 | 4 in 1 CARTON  | 11/16/2001              |                       |  |  |
| 1 |                      | 6 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  | 11/16/2001              |                       |  |
|                       |   |                         |                       |  |

# Labeler - Sato Pharmaceutical Co., Ltd. (690575642)

| Establishment                 |         |           |   |
|-------------------------------|---------|-----------|---|
| Name                          | Address | ID/FEI    | Business Operations   |
| Sato Pharmaceutical Co., Ltd. |         | 715699133 | manufacture(49873-109) , label(49873-109) , pack(49873-109) |

Revised: 12/2023 Sato Pharmaceutical Co., Ltd.