SATO CLEAR- naphazoline hydrochloride solution/ drops Sato Pharmaceutical Co., Ltd.

Sato Clear

Active ingredient

Naphazoline hydrochloride 0.012%

Purpose

Naphazoline hydrochloride Vasoconstrictor

Uses relieves this symptom due to minor eye irritations

redness of the eye

Warnings

For external use only

Do not use

if solution changes color or becomes cloudy

Ask a doctor before use if you have

narrow angle glaucoma

When using this product

- pupils may become enlarged temporarily
- overuse of this product may produce increased redness of the eyes
- to avoid contamination, do not touch tip of container to any surface
- replace cap after using

Stop use and ask a doctor if you experience

- eye pain
- changes in vision
- continued redness or irritation of the eye
- the condition worsens or persists for more than 72 hours

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

■ Place 1 or 2 drops in the affected eyes up to 4 times daily.

Other information

- use only if imprinted tamper-resistant seal on the bottle cap is intact
- keep container tightly closed

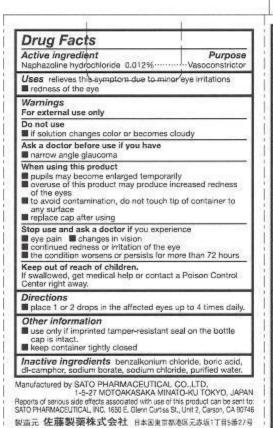
Inactive ingredients benzalkonium chloride, boric acid, dl-camphor, sodium borate, sodium chloride, purified water.



*Use only if imprinted carton overwrap with
"("91-088 Sato SAFETY SEAL" is intact at
time of purchase.

配合目的

US-K





SATO CLEAR

naphazoline hydrochloride solution/ drops

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Product Type HUMAN OTC DRUG Item Code (Source) NDC:49873-044

LOT

Route of Administration OPHTHALMIC

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
NAPHAZOLINE HYDROCHLORIDE (UNII: MZ1131787D) (NAPHAZOLINE -	NAPHAZ OLINE	12 mg
UNII:H231GF11BV)	HYDROCHLORIDE	in 100 mL

Inactive Ingredients

Ingredient Name	Strength
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
BORIC ACID (UNII: R57ZHV85D4)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
WATER (UNII: 059QF0KO0R)	

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49873- 044-01	1 in 1 CARTON	05/31/1989	
1		15 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M018	05/31/1989	

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

Establishment			
Name	Address	ID/FEI	Business Operations
Sato Pharmaceutical Co., Ltd.		715699133	manufacture(49873-044) , label(49873-044) , pack(49873-044)

Revised: 11/2023 Sato Pharmaceutical Co., Ltd.