SODIUM HYALURONATE EYE DROPS- sodium hyaluronate liquid
Guangzhou Hechuang Commercial Service Co.,Ltd.

Sodium Hyaluronate Eye Drops

ACTIVE INGREDIENT

Sodium Hyaluronate 0.1%

Purpose

Eye lubricant

Inactive Ingredients

Water□sodium chloride□Pyridoxine□Cyanocobalamin

Used for dry eye syndrome
Relieve dry eye symptoms

•The temporary relief of burning and irritation due to dryness of the eye.
•The temporary relief of discomfort due to minor irritations of the eye or exposure to wind or sun.
•Use as a protectant against further irritation or to relieve dryness of the eye.
•Use as a lubricant to prevent further irritation or to relieve dryness of the eye.

Directions
To open, Twist and pull tab to remove.
•Instill 1 or 2 drops as needed.
•If used for dryness and discomfort associated with LASIK, follow your eye doctor's instructions.

Warnings

•For external use only.
•To avoid contamination, do not touch tip of container to any surface.
•Use within 30 days after opening. Maintain good hygiene during use and reseal tightly after each use.
•If solution changes color, do not use.

Stop use and ask a doctor

Stop use and ask a doctor.
If you experience eye pain, changes in vision, continued redness or irritation of the eye, or if the condition worsens or persists for more than 72 hours.

Keep out of reach of children

Keep out of reach of children.
If swallowed, get medical help or contact a Poison Control Center right away.

Other Information

- Use only if container is intact.
- Use before expiration date markedon container.
- Store at 59°-86°F (15°-30°C).
- Protect from sunlight.
- Retain this carton for future reference.

Distributed by:Mitsui Pharmaceutical Co., Ltd.

SODIUM HYALURONATE EYE DROPS

sodium hyaluronate liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:87298-037
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM HYALURONATE (UNII: YSE9PPT4TH) (HYALURONIC ACID - UNII:S270N0TRQY)	SODIUM HYALURONATE	0.1 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
PYRIDOXINE (UNII: KV2JZ1BI6Z)	
CYANOCOBALAMIN (UNII: P6YC3EG204)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:87298-037-01	15 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/11/2026	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M018	05/11/2026	

Labeler - Guangzhou Hechuang Commercial Service Co.,Ltd. (704398144)

Establishment

Name	Address	ID/FEI	Business Operations
Guangzhou Hechuang Commercial Service Co.,Ltd.		704398144	label(87298-037) , manufacture(87298-037)

Revised: 5/2026 Guangzhou Hechuang Commercial Service Co.,Ltd.