FOCUS TEARS PF- carboxymethylcellulose sodium solution/ drops Covalent Medical

Carboxymethylcellulose sodium

Active ingredient

Carboxymethylcellulose sodium (0.5%)

Purpose

Eye Lubricant

Warnings

For use in the eyes only.

When using this product

- do not touch tip of container to any surface to avoid contamination.
- do not reuse; once opened discard.

Stop use and ask a doctor if

- you experience eye pain, changes in vision, continued redness or irritation of the eye.
- tile 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

■ Instill 1 or 2 drops into affected eye(s) as needed and discard container.

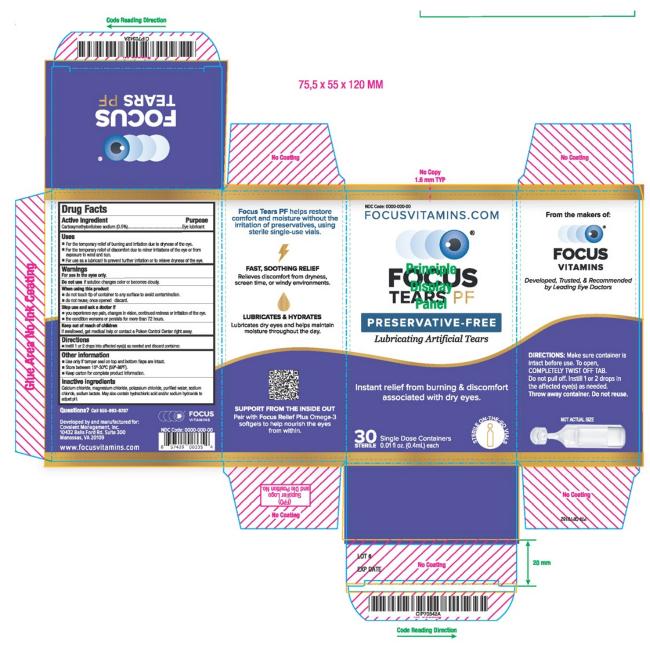
Other information

- Use only if tamper seal on top and bottom flaps are intact.
- Store between 15°-30°C (59°-86°F).
- Keep carton for complete product information.

Inactive ingredients

Calcium chloride, magnesium chloride, potassium chloride, purified water, sodium chloride, sodium lactate. May also contain hydrochloric acid and/or sodium hydroxide to

Principal Display Panel



FOCUS TEARS PF

carboxymethylcellulose sodium solution/ drops

Product Information					
Product Type	HUMAN OTC DRUG	Item Code (S	ource)	NDC:8272	24-009
Route of Administration	OPHTHALMIC				
Active Ingredient/Active Moiety					
Ingredient Name			Basis of Str	ength	Strength

CARBOXYMETHYLCELLULOSE	SODIUM (UNII: K6790BS311)
(CARRONAMETINA CELLUL OCE I	INIII OFIZIZD10V)

(CARBOXYMETHYLCELLULOSE - UNII:05JZ17B19X)

CARBOXYMETHYLCELLULOSE 5 mg in 1 mL

Inactive Ingredients				
Ingredient Name	Strength			
CALCIUM CHLORIDE (UNII: M4I0D6VV5M)				
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)				
POTASSIUM CHLORIDE (UNII: 660YQ98I10)				
WATER (UNII: 059QF0KO0R)				
SODIUM CHLORIDE (UNII: 451W47IQ8X)				
SODIUM LACTATE (UNII: TU7HW0W0QT)				
HYDROCHLORIC ACID (UNII: QTT17582CB)				
SODIUM HYDROXIDE (UNII: 55X04QC32I)				

ı	Packaging				
7	# Item Code	Package Description	Marketing Start Date	Marketing End Date	
:	NDC:82724- 009-20	30 in 1 BOX	09/01/2025		
:	L	0.4 mL in 1 VIAL, SINGLE-USE; 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	09/01/2025	

Labeler - Covalent Medical (112467517)

Establishment				
Name	Address	ID/FEI	Business Operations	
Unither Manufacturing LLC		079176615	manufacture(82724-009)	

Revised: 7/2025 Covalent Medical