THERATEARS LUBRICANT- carboxymethylcellulose sodium solution/ drops
MEDTECH PRODUCTS INC

TheraTears MD 58790-001

Drug Facts

Active ingredient

Sodium carboxy-
methylcellulose 0.25%

Purpose

Eye lubricant

Uses

- As a lubricant to relieve dryness of the eye.
- As a protectant against further irritation of the eye.
- For temporary relief of burning, irritation, and discomfort including exposure to wind or sun.

Warnings

For external use only

- To avoid contamination do not touch tip of opened container to any surface. Replace cap after using.

Do not use

- If solution changes color or becomes cloudy.

Stop use and ask a doctor if

- You experience eye pain, changes in vision, continued redness or irritation.
- 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 immediately.

Directions

- Instill 1 or 2 drops in the affected eye(s) as needed.
- After use and prior to re-capping, shake any residual product from the dropper tip.

Other information

- Store between 15-25C (59-77F)
- Discard 45 days after opening.

Inactive ingredients

boric acid, calcium chloride, diethylenetriamine pentamethylene phosphonic acid, hydrochloric acid, magnesium chloride, phosphonic acid, potassium chloride, sodium bicarbonate, sodium chloride, sodium perborate, sodium phosphate, water

Questions or comments?

1-800-579-8327

Principal Display Panel Text for Carton Label:

thera

tears®

LUBRICANT

EYE DROPS

1 FL OZ (30 mL)

THERATEARS LUBRICANT

carboxymethylcellulose sodium solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:58790-001
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
carboxymethylcellulose sodium (UNII: K679OBS311) (carboxymethylcellulose - UNII:05JZI7B19X)	carboxymethylcellulose sodium	2.5 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
boric acid (UNII: R57ZHV85D4)	
sodium perborate (UNII: Y52BK1W96C)	

calcium chloride (UNII: M4I0D6VV5M)	
diethylenetriamine pentamethylene phosphonic acid (UNII: 0Q75589TM3)	
magnesium chloride (UNII: 02F3473H9O)	
potassium chloride (UNII: 660YQ98I10)	
water (UNII: 059QF0KO0R)	
sodium bicarbonate (UNII: 8MDF5V39QO)	
sodium chloride (UNII: 451W47IQ8X)	
sodium phosphate (UNII: SE337SVY37)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:58790-001-30	1 in 1 CARTON	07/01/1999	
1		30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
2	NDC:58790-001-15	1 in 1 CARTON	07/01/1999	
2		15 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
3	NDC:58790-001-31	2 in 1 CARTON	12/01/2020	
3		30 mL in 1 BOTTLE, DROPPER; 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	07/01/1999	

Labeler - MEDTECH PRODUCTS INC (114707784)

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