GONIOSOFT - hypromellose 2.5% liquid
OCuSOFT, Inc.

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Drug Facts

Active ingredient

Hypromellose 2.5%

Purpose

Demulcent

Use

● For professional use in Gonioscopic examinations.

Warnings

● **For use in the eyes only**. ● To avoid contamination do not touch tip of container to any surface. ● Replace cap after using. ● Not for use in conjunction with hot laser treatment.

Do not use if solution changes color or becomes cloudy

KEEP OUT OF REACH OF CHILDREN.

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

Directions

● Fill Gonioscopic prism with solution as necessary.

Other information

● Store between 15º-30ºC (59º-86ºF). ● Keep tightly closed. ● If this solution dries on optical surfaces, let stand in cool water before cleansing.

DO NOT USE IF IMPRINTED SEAL ON CAP IS TORN, BROKEN OR MISSING.

Inactive ingredients

Benzalkonium Chloride, Boric Acid, Edetate Disodium, Sodium Borate, Water for Injection, Hydrochloric Acid and/or Sodium Hydroxide may be added to adjust pH.

Questions or Comments?

800-233-5469 or www.ocusoft.com

PRINCIPAL DISPLAY PANEL

GONIOSOFT™

NDC 54799-503-15

Hypromellose 2.5% Ophthalmic Demulcent Solution (Sterile)

0.5 fl oz (15mL)

Distributed by OCuSOFT Inc.
Rosenberg, TX 77471 USA
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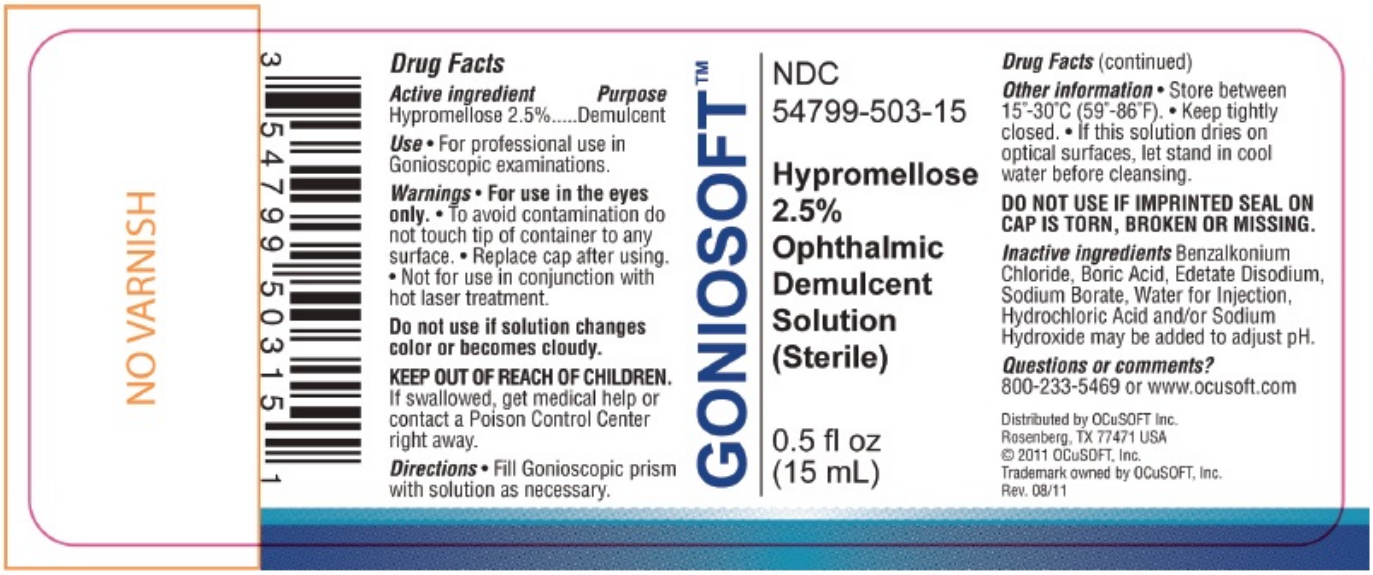
Rev. 08/11



GONIOSOFT

hypromellose 2.5% liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:54799-503
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Hypromelloses (UNII: 3NXW29V3WO) (Hypromelloses - UNII:3NXW29V3WO)	Hypromelloses	25 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
Benzalkonium Chloride (UNII: F5UM2KM3W7)	

Boric Acid (UNII: R57ZHV85D4)	
Edetate Disodium (UNII: 7FLD91C86K)	
Sodium Borate (UNII: 91MBZ8H3QO)	
Water (UNII: 059QF0KO0R)	
Hydrochloric Acid (UNII: QTT17582CB)	
Sodium Hydroxide (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54799-503-15	15 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/02/1989	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		10/02/1989	

Labeler - OCuSOFT, Inc. (174939207)

Establishment

Name	Address	ID/FEI	Business Operations
Altaire Pharmaceuticals, Inc.		786790378	MANUFACTURE(54799-503)

Establishment

Name	Address	ID/FEI	Business Operations
OCuSOFT, Inc.		174939207	MANUFACTURE(54799-503)

Revised: 12/2012 OCuSOFT, Inc.