BIOGLO- fluorescein sodium strip
HUB Pharmaceuticals, 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.

BIO GLO (fluorescein sodium) strip

Product Facts

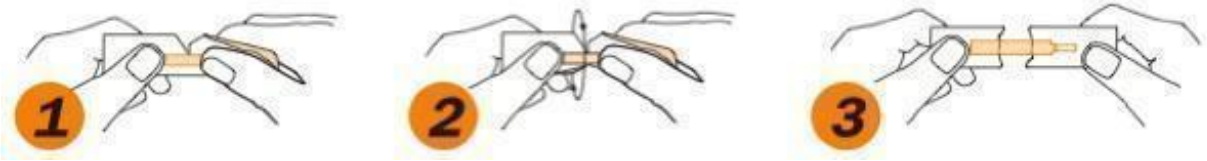
Each sterile strip is impregnated with 1 mg. of fluorescein sodium U.S.P.

INDICATIONS:

For staining the anterior segment of the eye when fitting contact lenses, in disclosing corneal injury and in applanation tonometry.

DIRECTIONS FOR USE:

To insure full fluorescence and patient comfort, the BioGlo impregnated tip should be moistened before application. One or two drops of sterile irrigating or saline solution should be used for this purpose. Touch conjunctiva or fornix as required with moistened tip. It is recommended that the patient blink several times after application.



1. Grasp tabs between thumbs & index fingers
2. Gently pull tabs apart
3. Remove strip taking care to touch only the white portion of the strip

STORAGE:

Keep at room temperature.

NOTE:

For external use only. Contents may not be sterile if individual strips have been damaged or previously opened.

Keep out of reach of children

HOW SUPPLIED:

Dispenser carton containing 100 or 300 strips.

HUB Pharmaceuticals, LLC

Rancho Cucamonga, CA 91730

www.hubrx.com

Mfr. Lic. No. G/1197

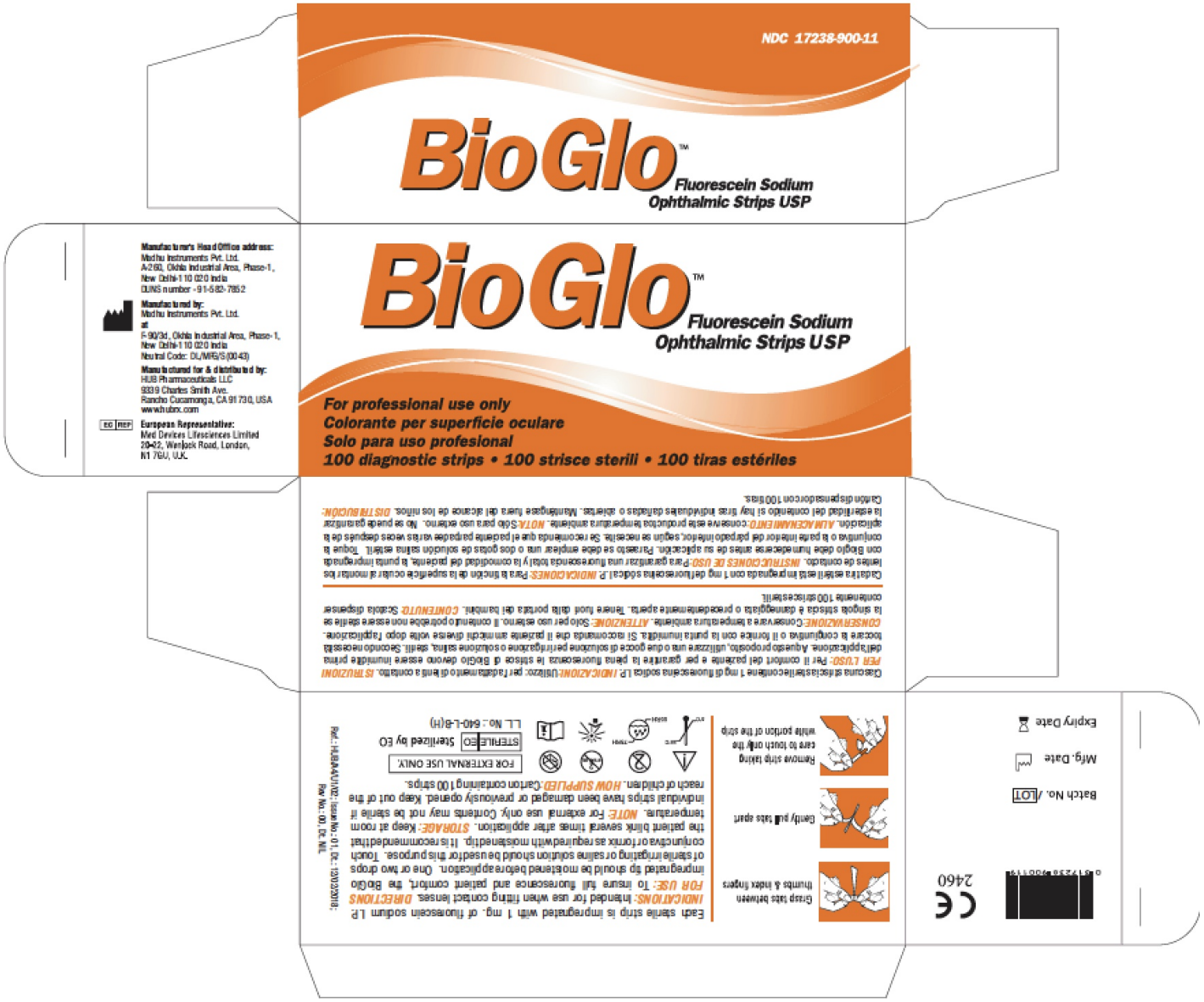
European Representative:

Biovision Limited

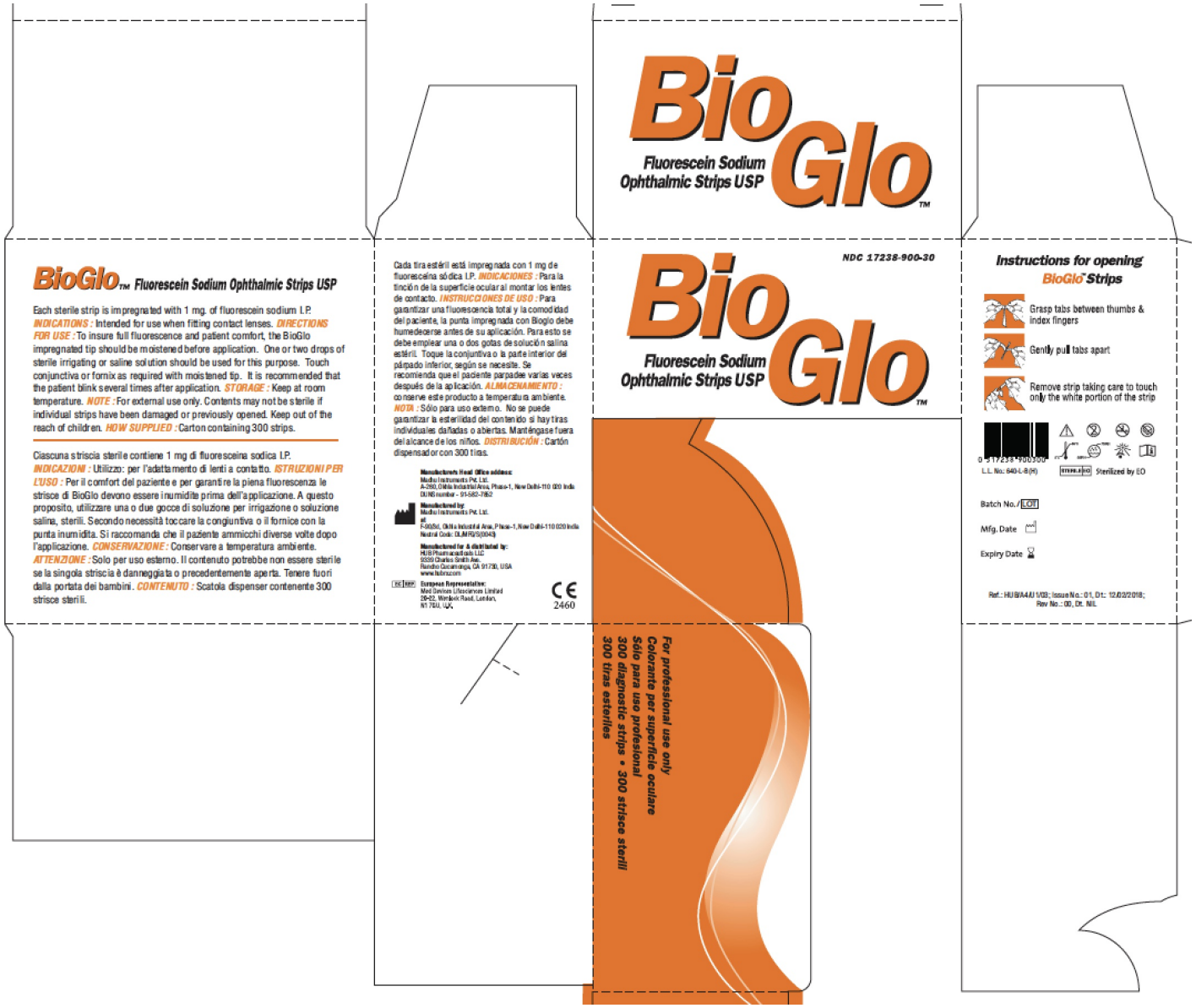
Wayside, Tring Road, Wellhead, Dunstable, BEDS LU6 2JU, UK

Representative Packaging:

17238-900-99: BioGlo Individual Pouch (above)



17238-900-11: Dispenser Carton containing 100 strips (above)



17238-900-30: Dispenser Carton containing 300 strips (above)

BIOGLO

fluorescein sodium strip

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:17238-900
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
FLUORESCEIN SODIUM (UNII: 93X55PE38X) (FLUORESCEIN - UNII:TPY09G7XIR)	FLUORESCEIN SODIUM	1 mg in 1 mg

Packaging

		Marketing Start	Marketing End

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:17238-900-11	100 in 1 BOX	04/01/2018	
1	NDC:17238-900-99	1 mg in 1 PACKET; Type 0: Not a Combination Product		
2	NDC:17238-900-30	300 in 1 BOX	04/01/2018	
2	NDC:17238-900-99	1 mg in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		04/01/2012	

Labeler - HUB Pharmaceuticals, Inc. (611747945)

Establishment

Name	Address	ID/FEI	Business Operations
OMNI LENS PRIVATE LIMITED		862170057	manufacture(17238-900)

Establishment

Name	Address	ID/FEI	Business Operations
Madhu Instruments Pvt.Ltd.		915827852	manufacture(17238-900)

Revised: 1/2024 HUB Pharmaceuticals, Inc.