LUMIGAN- bimatoprost solution/ drops
Allergan, Inc.

Lumigan 03

Principal Display Panel - 0.3 mg/1 mL Carton Label

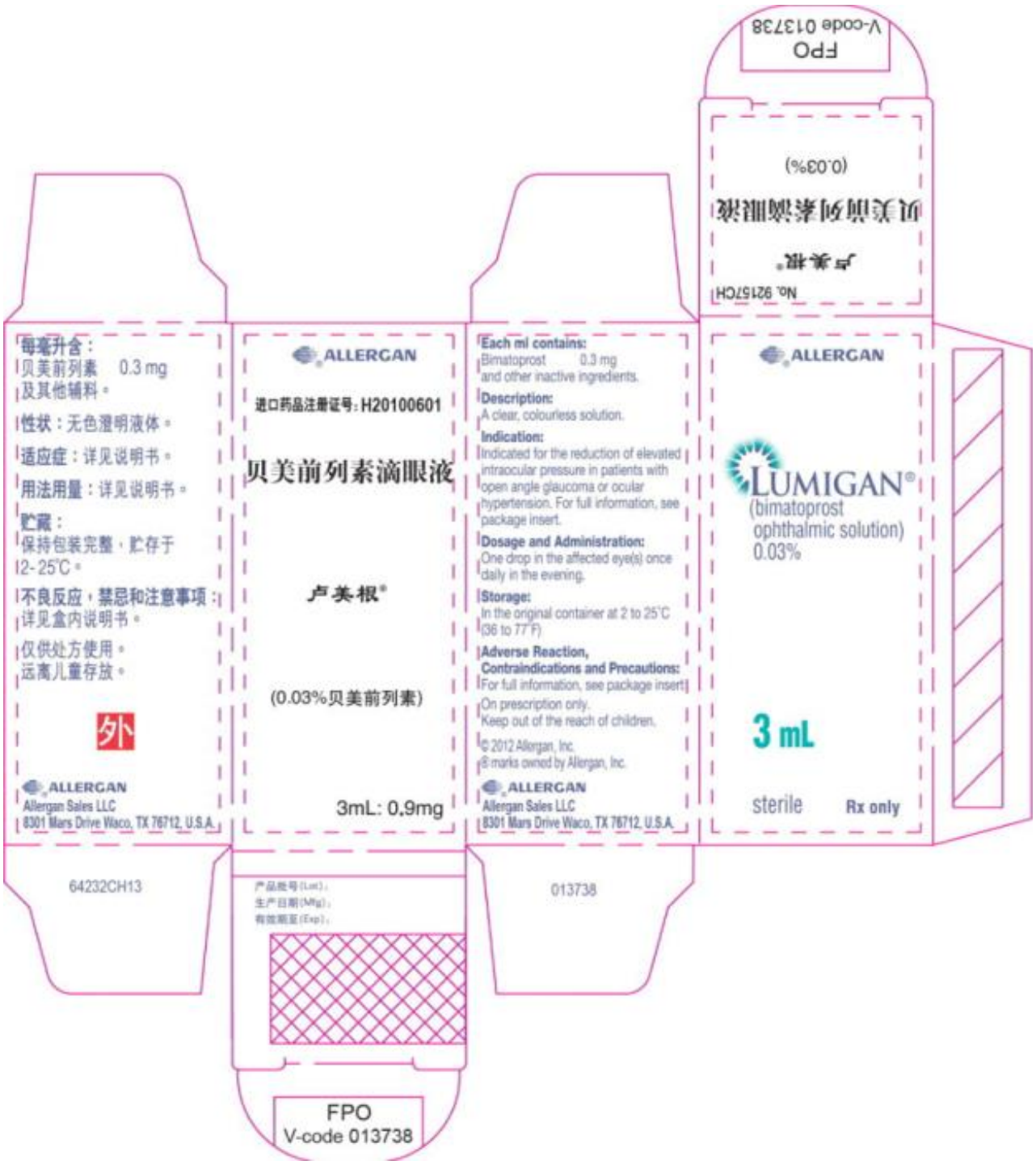
®ALLERGAN

LUMIGAN®
(bimatoprost
ophthalmic solution)

0.03%

3 mL

sterile **Rx only**



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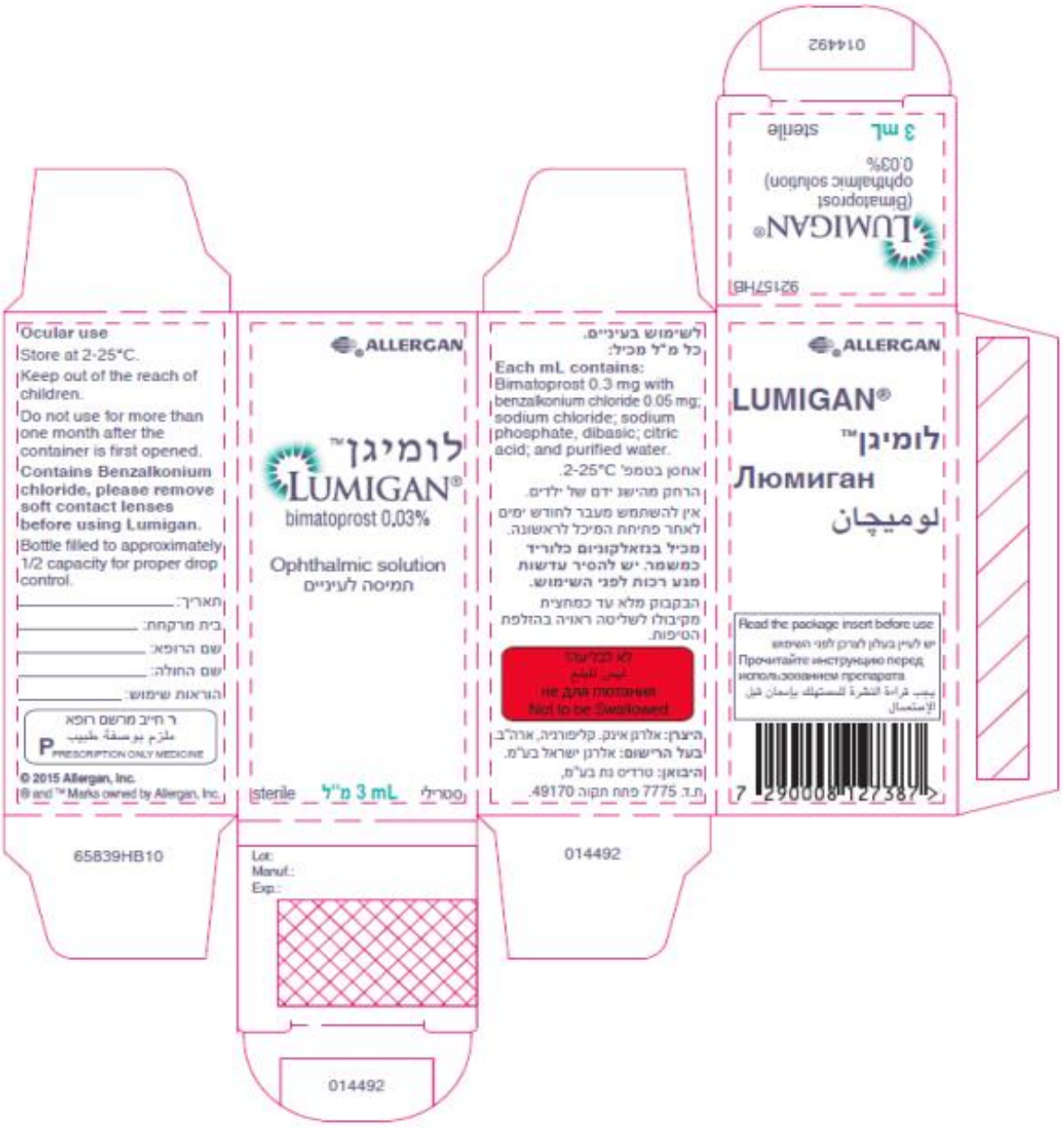

LUMIGAN

bimatoprost solution/ drops

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:0023-9187
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BIMATOPROST (UNII: QXS94885MZ) (BIMATOPROST - UNII:QXS94885MZ)	BIMATOPROST	0.3 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
SODIUM PHOSPHATE, DIBASIC (UNII: GR686LBA74)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
WATER (UNII: 059QF0KO0R)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0023-9187-01	1 in 1 CARTON	03/22/2001	
1		2.5 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
2	NDC:0023-9187-03	1 in 1 CARTON	03/22/2001	
2		2.5 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
3	NDC:0023-9187-05	1 in 1 CARTON	03/22/2001	
3		5 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
4	NDC:0023-9187-07	1 in 1 CARTON	03/22/2001	
4		7.5 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
Export only		03/22/2001	

Labeler - Allergan, Inc. (144796497)

Revised: 12/2018 Allergan, Inc.