LIDOCAINE- lidocaine ointment
Aleor Dermaceuticals Limited

Lidocaine Ointment USP, 5%
FOR TOPICAL USE ONLY. DO NOT USE IN THE EYES
Rx Only

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

NDC 71589-001-30

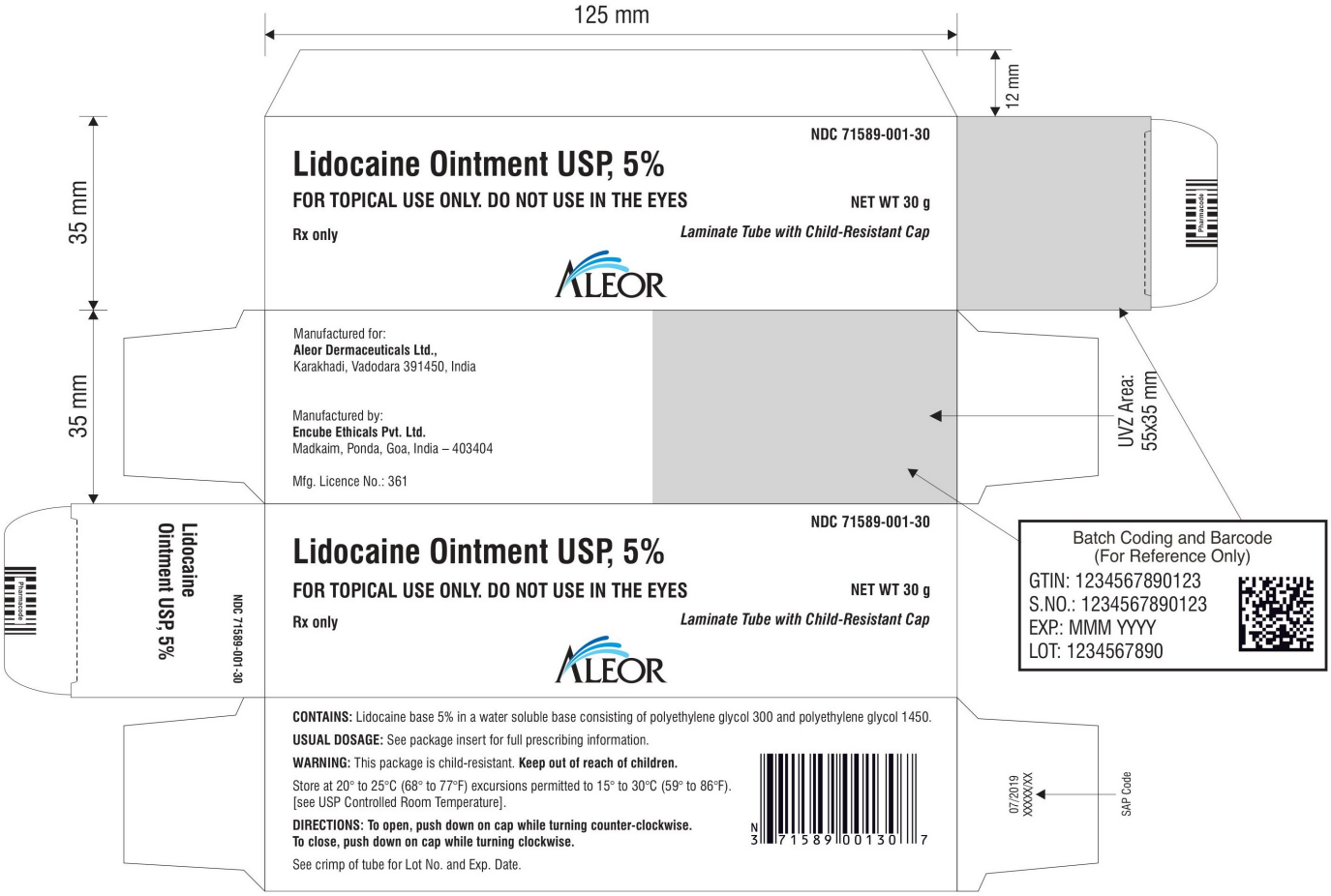
Rx only

Lidocaine Ointment USP, 5%

Laminate Tube with Child-Resistant Cap

FOR TOPICAL USE ONLY. DO NOT USE IN THE EYES

NET WT 30 g



NDC 71589-001-35

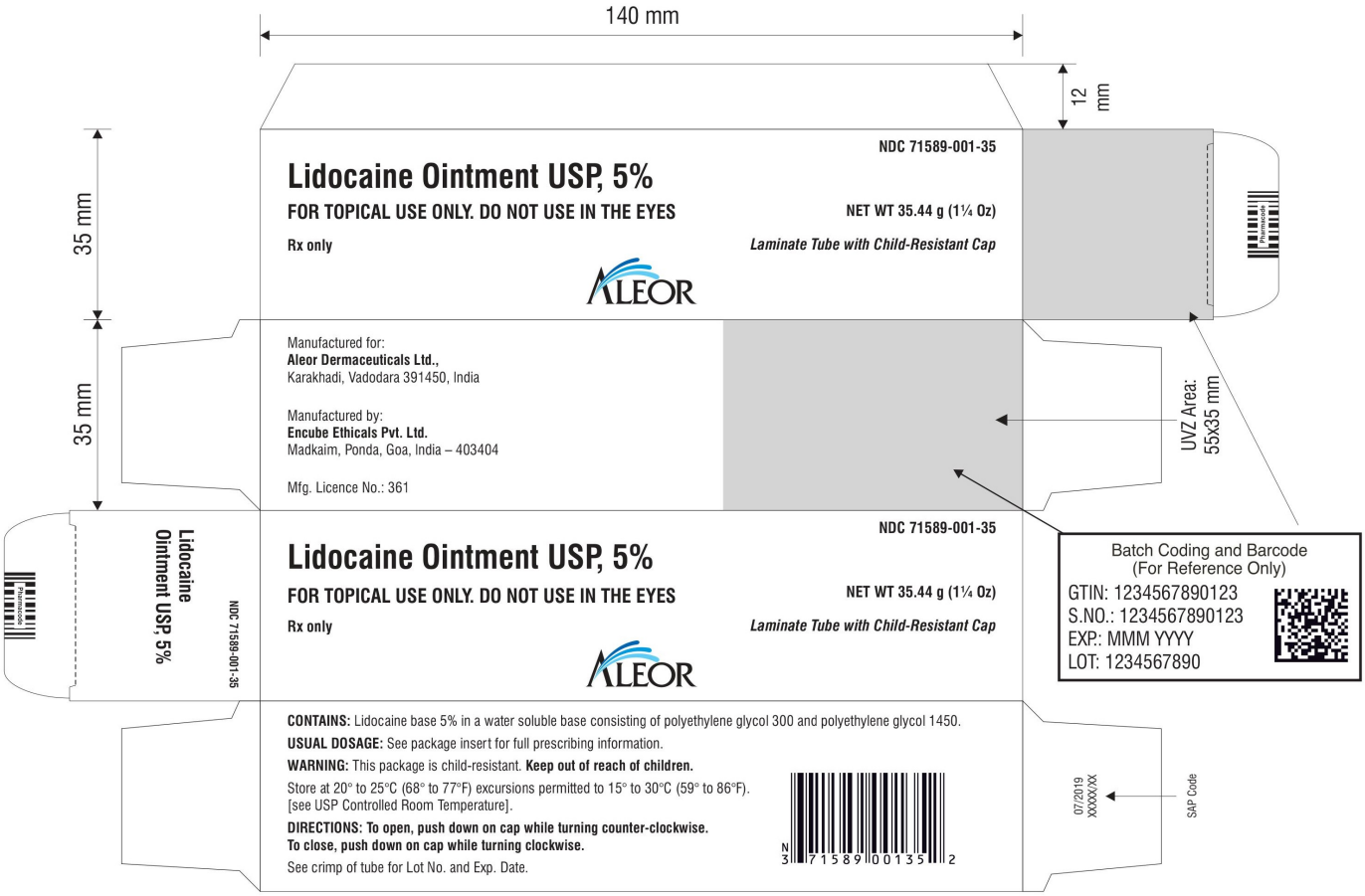
Rx only

Lidocaine Ointment USP, 5%

Laminate Tube with Child-Resistant Cap

FOR TOPICAL USE ONLY. DO NOT USE IN THE EYES

NET WT 35.44 g (1¼ Oz)



NDC 71589-001-50

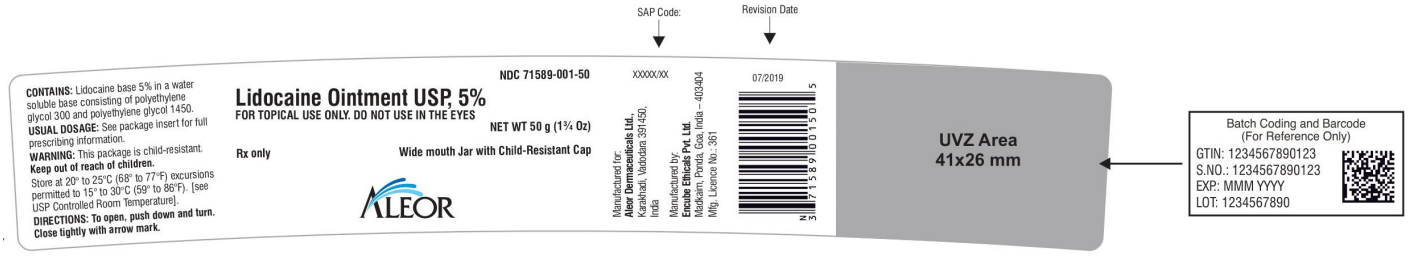
Rx only

Lidocaine Ointment USP, 5%

Wide mouth Jar with Child-Resistant Cap

FOR TOPICAL USE ONLY. DO NOT USE IN THE EYES

NET WT 50 g (1¾ Oz)



LIDOCAINE

lidocaine ointment

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:71589-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	50 mg in 1 g

Inactive Ingredients	
Ingredient Name	**Strength**
POLYETHYLENE GLYCOL 300 (UNII: 5655G9Y8AQ)	
POLYETHYLENE GLYCOL 1450 (UNII: OJ4Z5Z32L4)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:71589-001-30	1 in 1 CARTON	11/30/2018	
1		30 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:71589-001-35	1 in 1 CARTON	11/30/2018	
2		35.44 g in 1 TUBE; Type 0: Not a Combination Product		
3	NDC:71589-001-50	50 g in 1 JAR; Type 0: Not a Combination Product	11/30/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA211469	11/30/2018	

Labeler - Aleor Dermaceuticals Limited (871411532)

Registrant - Aleor Dermaceuticals Limited (871411532)

Establishment

Name	Address	ID/FEI	Business Operations
Encube Ethicals Pvt. Ltd.		725076298	MANUFACTURE(71589-001)

Revised: 7/2019 Aleor Dermaceuticals Limited