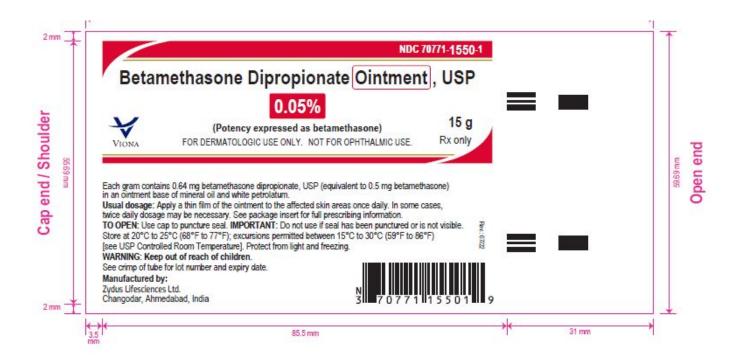
BETAMETHASONE DIPROPIONATE - betamethasone dipropionate ointment Zydus Lifesciences Limited

BETAMETHASONE DIPROPIONATE OINTMENT USP, 0.05%

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

Betamethasone Dipropionate Ointment USP, 0.05% NDC 70771-1550-1 in tube of 15 gm Rx Only





BETAMETHASONE DIPROPIONATE

betamethasone dipropionate ointment

Product Information

Product Type HUMAN PRESCRIPTION DRUG Item Code (Source) NDC:70771-1550

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name

BETAMETHASONE DIPROPIONATE (UNII: 826Y60901U) (BETAMETHASONE - UNII:9842X06Q6M)

Basis of Strength

BETAMETHASONE 0.5 mg in 1 g

Inactive	Ingredients

Ingredient Name	Strength		
MINERAL OIL (UNII: T5L8T28FGP)			
PETROLATUM (UNII: 4T6H12BN9U)			

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Item Code Package Description Marketing Start Date Marketing End Date

1	1550-1	1 in 1 CARTON	07/20/2020	
1		15 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:70771- 1550-4	1 in 1 CARTON	07/20/2020	
2		45 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
ANDA	ANDA214048	07/20/2020		

Labeler - Zydus Lifesciences Limited (918596198)

Establishment				
Name	Address	ID/FEI	Business Operations	
Zydus Lifesciences Limited		650650802	ANALYSIS(70771-1550), MANUFACTURE(70771-1550)	

Revised: 11/2022 Zydus Lifesciences Limited