

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	Basis of Strength	Strength
BETAMETHASONE DIPROPIONATE (UNII: 826Y60901U) (BETAMETHASONE - UNII:9842X06Q6M)	BETAMETHASONE	0.5 mg in 1 g

Inactive Ingredients

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:70771-			

1	NDC:70771-1550-1	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