

DESOXIMETASONE - desoximetasone cream
Zydus Lifesciences Limited

Desoximetasone Cream USP, 0.25%

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

NDC 70771-1437-1 in tube of 15gm

Desoximetasone Cream USP, 0.25%

Rx only


15 gm

NDC 70771-1437-1

Desoximetasone Cream, USP

0.25%

FOR TOPICAL USE ONLY. NOT FOR ORAL, OPHTHALMIC, OR INTRAVAGINAL USE.
Keep this and all medications out of the reach of children.

 **zydus**
pharmaceuticals

15 g
Rx only

Each gram contains: 2.5 mg of desoximetasone, USP in an emollient cream base consisting of cetostearyl alcohol, edetate disodium dihydrate, isopropyl myristate, lanolin alcohols, mineral oil, purified water and white petrolatum.


Usual dosage: Apply a thin film to affected skin area twice daily. Rub in gently.
See package insert for full prescribing information.

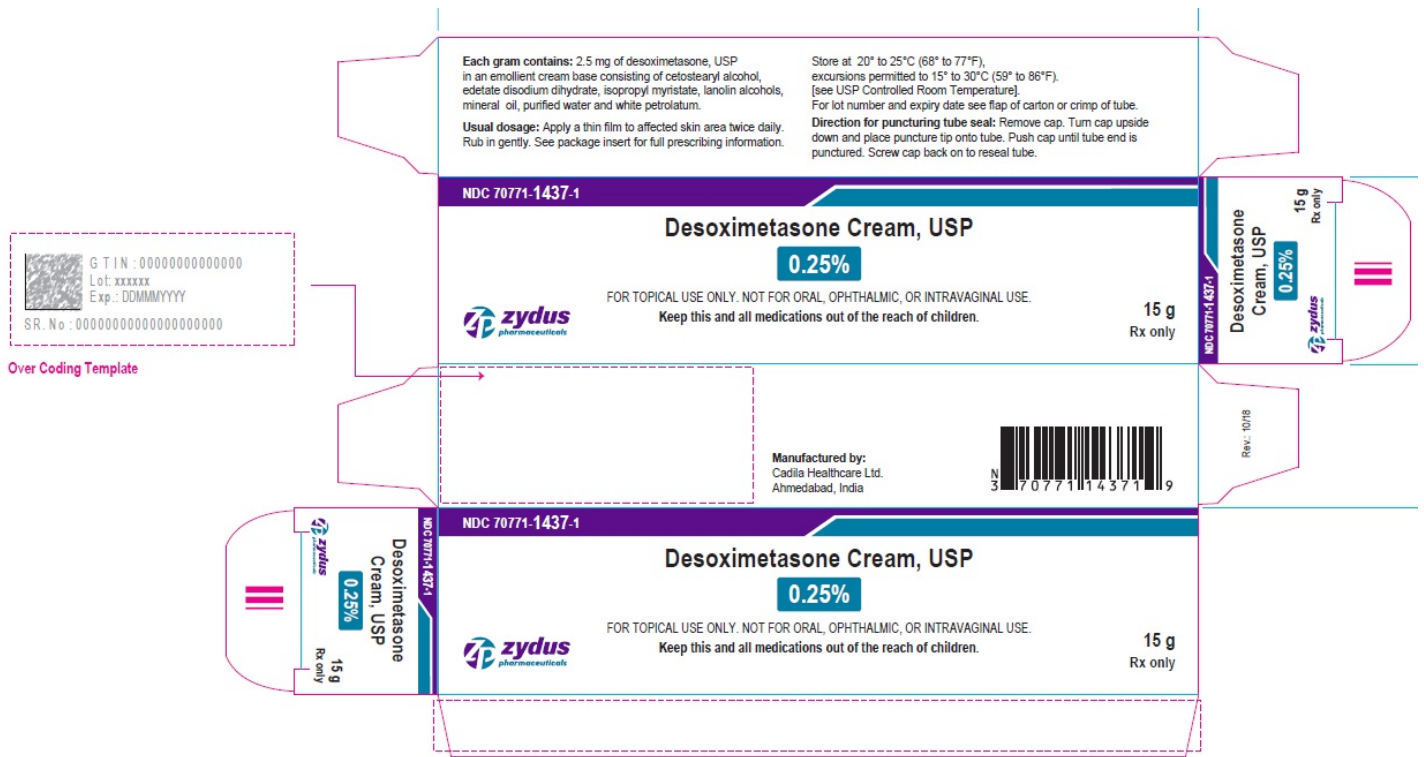
Store at 20° to 25°C (68° to 77°F), excursions permitted to 15° to 30°C (59° to 86°F).
[see USP Controlled Room Temperature].

To open: Remove cap. Turn cap upside down and place puncture tip onto tube.
Push cap until tube end is punctured. Screw cap back on to reseal tube.
For lot number and expiry date see crimp of tube.

Manufactured by:
Cadila Healthcare Ltd.
Ahmedabad, India

Rev: 10/18


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DESOXIMETASONE

desoximetasone cream

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:70771-1437
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DESOXIMETASONE (UNII: 4E07GXB7AU) (DESOXIMETASONE - UNII:4E07GXB7AU)	DESOXIMETASONE	2.5 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
LANOLIN ALCOHOLS (UNII: 884C3FA9HE)	
MINERAL OIL (UNII: T5L8T28FGP)	
WATER (UNII: 059QF0KO0R)	
PETROLATUM (UNII: 4T6H12BN9U)	

Product Characteristics

Color	WHITE (WHITE TO OFF-WHITE)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70771-1437-1	1 in 1 CARTON	02/12/2019	
1		15 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:70771-1437-3	1 in 1 CARTON	02/12/2019	
2		60 g in 1 TUBE; Type 0: Not a Combination Product		
3	NDC:70771-1437-0	1 in 1 CARTON	02/12/2019	
3		100 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	ANDA205620	02/12/2019	

Labeler - Zydus Lifesciences Limited (918596198)

Registrant - Zydus Lifesciences Limited (650650802)

Establishment

Name	Address	ID/FEI	Business Operations
Zydus Lifesciences Limited		650650802	ANALYSIS(70771-1437) , MANUFACTURE(70771-1437)

Revised: 11/2022

Zydus Lifesciences Limited