CYPROHEPTADINE HYDROCHLORIDE - cyproheptadine hydrochloride tablet
Zydus Lifesciences Limited

Cyproheptadine Hydrochloride Tablets, USP

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

NDC 70771-1076-1

Cyproheptadine Hydrochloride Tablets USP, 4 mg

100 Tablets

R_x only

CYPROHEPTADINE HYDROCHLORIDE

cyproheptadine hydrochloride tablet

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:70771-1076
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CYPROHEPTADINE HYDROCHLORIDE (UNII: NJ82J0F8QC) (CYPROHEPTADINE - UNII:2YHB6175DO)	CYPROHEPTADINE HYDROCHLORIDE	4 mg

Inactive Ingredients

Ingredient Name	Strength
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	

Product Characteristics

Color	WHITE (White)	**Score**	2 pieces
Shape	ROUND (Round)	**Size**	7mm
Flavor		**Imprint Code**	1110
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70771-1076-1	100 in 1 BOTTLE; Type 0: Not a Combination Product	07/20/2017	
2	NDC:70771-1076-0	1000 in 1 BOTTLE; Type 0: Not a Combination Product	07/20/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA208938	07/20/2017	

Labeler - Zydus Lifesciences Limited (918596198)

Registrant - Zydus Lifesciences Limited (918596198)

Establishment

Name	Address	ID/FEI	Business Operations
Zydus Lifesciences Limited		863362789	ANALYSIS(70771-1076) , MANUFACTURE(70771-1076)

Revised: 8/2022 Zydus Lifesciences Limited