

IBUPROFEN- ibuprofen tablet, film coated
RedPharm Drug, Inc.

ibuprofen

HOW SUPPLIED

600mg (white to off white, capsule shaped, biconvex, film coated tablets debossed with '122' on one side and plain on the other side) Bottles of 30, 50, 100 & 500

PACKAGE LABEL

NDC: 67296-1584-1

IBUPROFEN

600 MG

10 Tablets

Rx Only

Lot: _____

Exp: _____

Usual adult dosage: See package insert

Store at controlled room temperature: 20-25 C (68-77 F)

Mfg By: Marksans Pharma Limited
Verna Goa 403 722 India
49483-603-01

Dist. by: Redpharm Drug Eden Prairie, MN 55344

SIN 10814



0
15841
67296
3

NDC: 67296-1584-1
IBUPROFEN
 600 MG
 10 Tablets

Rx Only

Lot: _____ Exp: _____

Usual adult dosage: See package insert
 Store at controlled room temperature: 20-25 C (68-77 F)

Mfg By: Marksans Pharma Limited
 Verna Goa 403 722 India
 49483-603-01

Dist. by: Redpharm Drug Eden Prairie, MN 55344 SIN 10814



0
15841
67296
3



NDC: 67296-1584-3
IBUPROFEN
 600MG
 30 Tablets

Rx Only

Lot: A125GX1 Exp: 07/21

Usual adult dosage: See package insert
 Store at controlled room temperature: 20-25 C (68-77 F)

Mfg: Marksans Pharma LTD
 Verna GOA 403 722 India
 49483-603-50

Dist. by: Redpharm Drug Eden Prairie, MN 55344 SIN 70733



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15843
67296
3

(01)00367296158434
 (21)100000000000204
 (17)210731
 (10)A125GX1

IBUPROFEN

ibuprofen tablet, film coated

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:67296-1584(NDC:49483-603)
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
IBUPROFEN (UNII: WK2XYI10QM) (IBUPROFEN - UNII:WK2XYI10QM)	IBUPROFEN	600 mg

Inactive Ingredients

Ingredient Name	Strength
MAGNESIUM STEARATE (UNII: 70097M6I30)	

CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
STARCH, PREGELATINIZED CORN (UNII: O8232NY3SJ)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
TALC (UNII: 7SEV7J4R1U)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	

Product Characteristics

Color	white	Score	no score
Shape	CAPSULE	Size	18mm
Flavor		Imprint Code	122
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67296-1584-1	10 in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2018	
2	NDC:67296-1584-3	30 in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA090796	01/01/2018	

Labeler - RedPharm Drug, Inc. (828374897)

Establishment

Name	Address	ID/FEI	Business Operations
RedPharm Drug, Inc.		828374897	repack(67296-1584)