

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

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**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



4  
15843  
67296  
3

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

**IBUPROFEN**

ibuprofen tablet, film coated

**Product Information**

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

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
<b>IBUPROFEN</b> (UNII: WK2XYI10QM) (IBUPROFEN - UNII:WK2XYI10QM)	IBUPROFEN	600 mg

**Inactive Ingredients**

Ingredient Name	Strength
<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)	

<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	
<b>CELLULOSE, MICROCRYSTALLINE</b> (UNII: OP1R32D61U)	
<b>STARCH, PREGELATINIZED CORN</b> (UNII: O8232NY3SJ)	
<b>CROSCARMELOSE SODIUM</b> (UNII: M28OL1HH48)	
<b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ0SDW1A)	
<b>POLYVINYL ALCOHOL, UNSPECIFIED</b> (UNII: 532B59J990)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	
<b>TALC</b> (UNII: 7SEV7J4R1U)	

### Product Characteristics

<b>Color</b>	white	<b>Score</b>	no score
<b>Shape</b>	CAPSULE	<b>Size</b>	18mm
<b>Flavor</b>		<b>Imprint Code</b>	122
<b>Contains</b>			

### 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)

Revised: 1/2022

RedPharm Drug, Inc.