MEDI-FIRST IBUPROFEN- ibuprofen tablet, coated MEDI-FIRST PLUS IBUPROFEN- ibuprofen tablet, coated MEDIQUE IPRIN- ibuprofen tablet, coated DOVER ADDAPRIN- ibuprofen tablet, coated OTIS CLAPP ULTRAPRIN- ibuprofen tablet, coated Unifirst First Aid Corporation

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## **UniFirst First Aid Ibuprofen**

#### **Drug Facts**

#### Active ingredient

Ibuprofen 200 mg (NSAID)

\*nonsteroidal antinflamatory drug

## Purpose

Pain reliever/fever reducer

#### Uses

Temporarily relieves minor aches and pains associated with

- headache toothache backache menstrual cramps
- common cold muscular aches minor arthritis pain

Temporarily reduces fever.

## Warnings

**Allergy alert:** Ibuprofen may cause a severe allergic reaction, especially in people allergic to aspirin. Symptoms may include:

■ hives ■ skin reddening ■ asthma (wheezing) ■ facial swelling ■ rash ■ shock ■ blisters

If an allergic reaction occurs, stop use and seek medical help right away.

**Stomach bleeding warning:** This product contains an NSAID, which may cause severe stomach bleeding. The chance is higher if you:

- are age 60 or older
- have had stomach ulcers or bleeding problems
- take a blood thinning (anticoagulant) or steroid drug
- take other drugs containing prescription or nonprescription NSAIDs (aspirin, ibuprofen, naproxen, or others)
- have 3 or more alcoholic drinks every day while using this product
- take more or for a longer time than directed

**Heart attack or stroke warning:** NSAIDS, except aspirin, increase the risk of heart attack, heart failure, and stroke. These can be fatal. The risk is higher if you use more

than directed or for longer than directed.

#### Do not use

- if you have ever had an allergic reaction to any other pain reliever/fever reducer
- right before or after heart surgery

#### Ask a doctor before use if

- you have problems or serious side effects from taking pain relievers or fever reducers
- stomach bleeding warning applies to you
- you have a history of stomach problems such as heartburn
- you have high blood pressure, heart disease, liver cirrhosis, kidney disease, asthma or had a stroke
- you are taking a diuretic

## Ask a doctor or pharmacist before use if you are

- taking aspirin for heart attack or stroke, because ibuprofen may decrease this benefit of aspirin
- under a doctor's care for any serious condition
- taking any other drug

## When using this product

■ take with food or milk if stomach upset occurs

## Stop use and ask a doctor if

- you experience any of the following signs of stomach bleeding:
- feel faint vomit blood have bloody or black stools have stomach pain that does not get better
- you have symptoms of heart problems or stroke
- $\blacksquare$  chest pain  $\blacksquare$  trouble breathing  $\blacksquare$  weakness in one part or side of body  $\blacksquare$  slurred speech  $\blacksquare$  leg swelling
- pain gets worse or lasts for more than 10 days
- fever gets worse or lasts for more than 3 days
- redness or swelling is present in the painful area
- any new or unexpected symptoms occur

**If pregnant or breast-feeding,** ask a health professional before use. It is especially important not to use ibuprofen at 20 weeks or later in pregnancy unless specifically directed to do so by a doctor because it may cause problems in the unborn child or complications during delivery.

**Keep out of reach of children.** In case of overdose, get medical help or contact a Poison Control Center right away.

#### **Directions**

- do not use more than directed
- **■** the smallest effective dose should be used
- do not take longer than 10 days, unless directed by a doctor (see Warnings)

#### Adults and children: (12 years and older)

Take 1 tablet every 4 to 6 hours while symptoms persist. If pain or fever does not respond to 1 tablet, 2 tablets may be used.

Do not exceed 6 tablets in 24 hours, unless directed by a doctor.

## Children under 12 years:

Ask a doctor

#### Other information

- read all product information before using
- store at 68-77°F (20-25°C)
- avoid excessive heat 104°F (above 40°C)
- tamper evident sealed packets
- do not use any opened or torn packets

## Inactive ingredients

carnauba wax\*, corn starch, hypromellose\*, iron oxide red, lactose\*, magnesium stearate\*, microcrystalline cellulose\*, polydextrose\*, polyethylene glycol, povidone (K30)\*, silicon dioxide, sodium starch glycolate, stearic acid, talc\*, titanium dioxide

### Questions or comments? 1-800-634-7680

## Medi-First Ibuprofen Label

Medi-First®

Ibuprofen 200 mg

100 tablets (50 x 2)

Pain Reliever/Fever Reducer

Aches, Fever • Ibuprofen (NSAID) 200 mg

Pull to Open

Compare active ingredient to:

**Advil®** 

Registered Trademark of Pfizer Consumer Healthcare

This Package is for Households without Young Children.

<sup>\*</sup>may contain

#### Tamper Evident Unit Dose Packets



## Medi-First Plus Ibuprofen Label

Medi-First® Plus

**Ibuprofen** 

Ibuprofen 200 mg (NSAID)

100 Tablets (50 x 2's)

Pull To Open

This Package is for Households without Young Children.

Pain Reliever/Fever Reducer

Compare active ingredient to:

**Advil®** 

Registered Trademark of Pfizer Consumer Healthcare

Tamper Evident Unit Dose Packets



## **Medique Iprin Label**

 $\text{Medique} \, \mathbb{B}$ 

I-Prin

Ibuprofen 200 mg

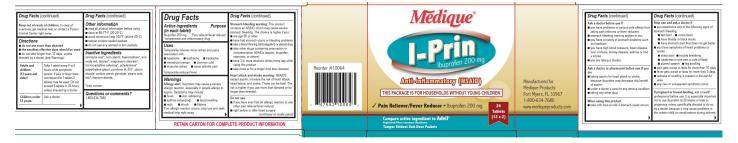
Anti-Inflammatory (NSAID)

This Package is for Households without Young Children.

Pain Reliever/Fever Reducer • Ibuprofen 200 mg

24 Tablets (12 x 2)

Tamper Evident Unit Dose Packets.



## **Dover Addaprin Label**

Dover Addaprin™

Ibuprofen 200 mg Tablets (NSAID)

Pain Reliever-Fever Reducer

This Package is for Households Without Young Children.

**Dover Pharmaceutical** 

Products of the highest quality and effectiveness

Tamper Evident

**Sealed Packets** 

**Unit Dose Packs** 

500 Tablets

(250 Packets of 2)



OC Otis Clapp

Quality & Integrity Since 1840

Ultraprin ™

Pain Reliever-Fever Reducer (NSAID)

Ibuprofen Tablets USP 200 mg

For Deep Seated Pain

See Warnings and Directions on Side Panel

Tear Out Along Perforation To Dispense

Professional Healthcare

500 Tablets (250 Packets of 2)



#### **MEDI-FIRST IBUPROFEN**

ibuprofen tablet, coated

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:47682-718	
Route of Administration	ORAL			

Active Ingredient/Active Molety				
Ingredient Name	<b>Basis of Strength</b>	Strength		
IBUPROFEN (UNII: WK2XYI10QM) (IBUPROFEN - UNII:WK2XYI10QM)	IBUPROFEN	200 mg		

Inactive Ingredients	
Ingredient Name	Strength
STARCH, CORN (UNII: O8232NY3SJ)	
POVIDONE K30 (UNII: U7250WY32X)	

STEARIC ACID (UNII: 4ELV7Z65AP)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
TALC (UNII: 7SEV7J4R1U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	

Product Characteristics				
Color	red (Reddish Brown)	Score	no score	
Shape	ROUND	Size	10mm	
Flavor		Imprint Code	G;2	
Contains				

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:47682-718- 13	250 in 1 BOX	01/26/2017		
1	NDC:47682-718- 99	2 in 1 PACKET; Type 0: Not a Combination Product			
2	NDC:47682-718- 48	125 in 1 BOX	01/26/2017		
2		2 in 1 PACKET; Type 0: Not a Combination Product			
3	NDC:47682-718- 33	50 in 1 BOX	01/26/2017		
3		2 in 1 PACKET; Type 0: Not a Combination Product			
4	NDC:47682-718- 30	4 in 1 BOX	01/26/2017		
4		2 in 1 PACKET; Type 0: Not a Combination Product			
5	NDC:47682-718- 50	25 in 1 BOX	04/16/2019		
5		2 in 1 PACKET; Type 0: Not a Combination Product			

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA079174	01/26/2017	

## **MEDI-FIRST PLUS IBUPROFEN**

ibuprofen tablet, coated

## **Product Information**

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:47682-709
Route of Administration	ORAL		

l	Active Ingredient/Active Moiety		
l	Ingredient Name	<b>Basis of Strength</b>	Strength
l	IBUPROFEN (UNII: WK2XYI10QM) (IBUPROFEN - UNII:WK2XYI10QM)	IBUPROFEN	200 mg

Inactive Ingredients			
Ingredient Name	Strength		
FERRIC OXIDE RED (UNII: 1K09F3G675)			
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)			
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)			
POVIDONE K30 (UNII: U725QWY32X)			
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)			
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)			
STEARIC ACID (UNII: 4ELV7Z65AP)			
STARCH, CORN (UNII: O8232NY3SJ)			
TALC (UNII: 7SEV7J4R1U)			

Product Characteristics				
Color	red (Reddish Brown)	Score	no score	
Shape	ROUND	Size	10mm	
Flavor		Imprint Code	G;2	
Contains				

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:47682-709- 48	125 in 1 BOX	01/26/2017		
1		2 in 1 PACKET; Type 0: Not a Combination Product			
2	NDC:47682-709- 33	50 in 1 BOX	01/26/2017		
2		2 in 1 PACKET; Type 0: Not a Combination Product			

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA079174	01/26/2017	

# **MEDIQUE IPRIN**

## ibuprofen tablet, coated

## **Product Information**

Product Type HUMAN OTC DRUG Item Code (Source	NDC:47682-700
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**Route of Administration** ORAL

## **Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
IBUPROFEN (UNII: WK2XYI10QM) (IBUPROFEN - UNII:WK2XYI10QM)	IBUPROFEN	200 mg

Inactive Ingredients		
Ingredient Name	Strength	
STARCH, CORN (UNII: O8232NY3SJ)		
STEARIC ACID (UNII: 4ELV7Z65AP)		
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)		
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)		
POVIDONE K30 (UNII: U725QWY32X)		
FERRIC OXIDE RED (UNII: 1K09F3G675)		
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)		
TALC (UNII: 7SEV7J4R1U)		
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)		

Product Characteristics				
Color	red (Reddish Brown)	Score	no score	
Shape	ROUND	Size	10mm	
Flavor		Imprint Code	G;2	
Contains				

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:47682-700- 69	3 in 1 BOX	01/26/2017		
1	NDC:47682-700- 99	2 in 1 PACKET; Type 0: Not a Combination Product			
2	NDC:47682-700- 64	12 in 1 BOX	01/26/2017		
2		2 in 1 PACKET; Type 0: Not a Combination Product			
3	NDC:47682-700- 47	100 in 1 BOX	01/26/2017		
3		2 in 1 PACKET; Type 0: Not a Combination Product			
4	NDC:47682-700- 13	250 in 1 BOX	01/26/2017		
Л		2 in 1 PACKET; Type 0: Not a Combination			

## **DOVER ADDAPRIN**

ibuprofen tablet, coated

#### **Product Information**

Product Type HUMAN OTC DRUG Item Code (Source) NDC:47682-714

Route of Administration ORAL

Product

## **Active Ingredient/Active Moiety**

Ingredient Name

Basis of Strength

IBUPROFEN (UNII: WK2XYI10QM) (IBUPROFEN - UNII: WK2XYI10QM)

BUPROFEN (UNII: WK2XYI10QM) | IBUPROFEN | 200 mg

Inactive Ingredients	
Ingredient Name	Strength
STARCH, CORN (UNII: O8232NY3SJ)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
POVIDONE K30 (UNII: U725QWY32X)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
TALC (UNII: 7SEV7J4R1U)	

Product Characteristics				
Color	red (Reddish Brown)	Score	no score	
Shape	ROUND	Size	10mm	
Flavor		Imprint Code	G;2	
Contains				

F	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47682-714- 13	250 in 1 BOX	01/26/2017	

NDC:47682-714- 2 in 1 PACKET; Type 0: Not a Combination Product

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA079174	01/26/2017	

## **OTIS CLAPP ULTRAPRIN**

ibuprofen tablet, coated

**Product Information** 

Product Type HUMAN OTC DRUG Item Code (Source) NDC:47682-702

Route of Administration ORAL

**Active Ingredient/Active Moiety** 

FERRIC OXIDE RED (UNII: 1K09F3G675)

Ingredient Name Basis of Strength Strength

IBUPROFEN (UNII: WK2XYI10QM) (IBUPROFEN - UNII: WK2XYI10QM) IBUPROFEN 200 mg

Inactive Ingredients	
Ingredient Name	Strength
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
TALC (UNII: 7SEV7J4R1U)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
<b>SODIUM STARCH GLYCOLATE TYPE A POTATO</b> (UNII: 5856J3G2A2)	
POVIDONE K30 (UNII: U725QW/32X)	
STARCH, CORN (UNII: 08232NY3SJ)	

Product Characteristics			
Color	red (Reddish Brown)	Score	no score
Shape	ROUND	Size	10mm
Flavor		Imprint Code	G;2
Contains			

	Packaging						
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
П							

1 NDC:47082-702-	250 in 1 BOX	02/01/2017	04/03/2017			
1 NDC:47682-702-	2 in 1 PACKET; Type 0: Not a Combination Product					
Marketing Information						
Marketing	Application Number or Monograph	_	Marketing End			
Category	Citation	Date	Date			

# **Labeler -** Unifirst First Aid Corporation (832947092)

Revised: 3/2024 Unifirst First Aid Corporation