**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**

---

**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

■ chest pain ■ trouble breathing ■ weakness in one part or side of body ■ slurred speech ■ 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

**may contain*

***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

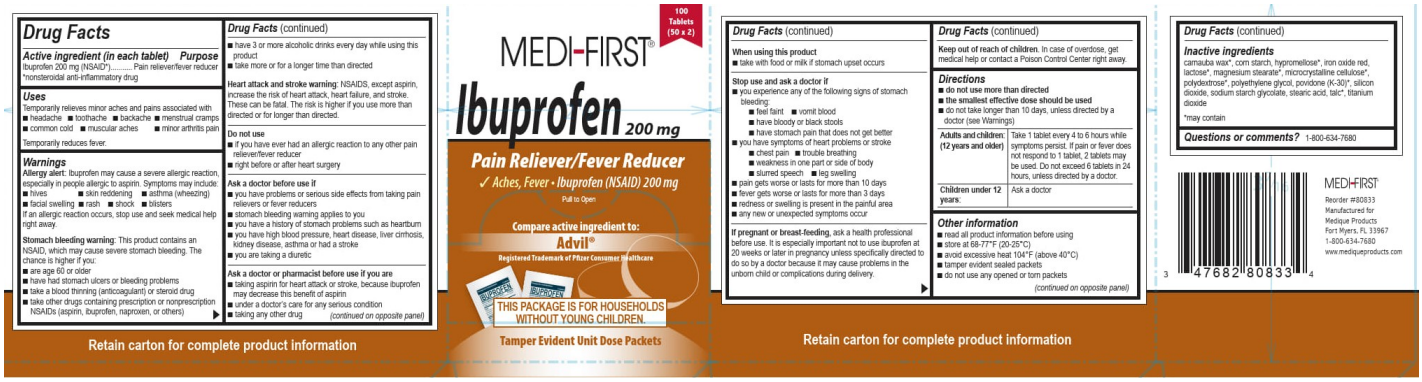
Compare active ingredient to:

Advil®

Registered Trademark of Pfizer Consumer Healthcare

This Package is for Households without Young Children.

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 isfor 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

Medique®

I-Prin

Ibuprofen 200 mg

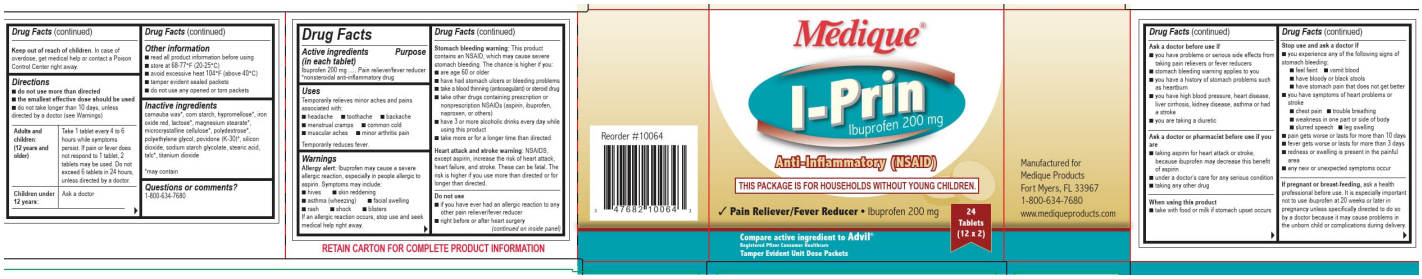
Anti-Inflammatory (NSAID)

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

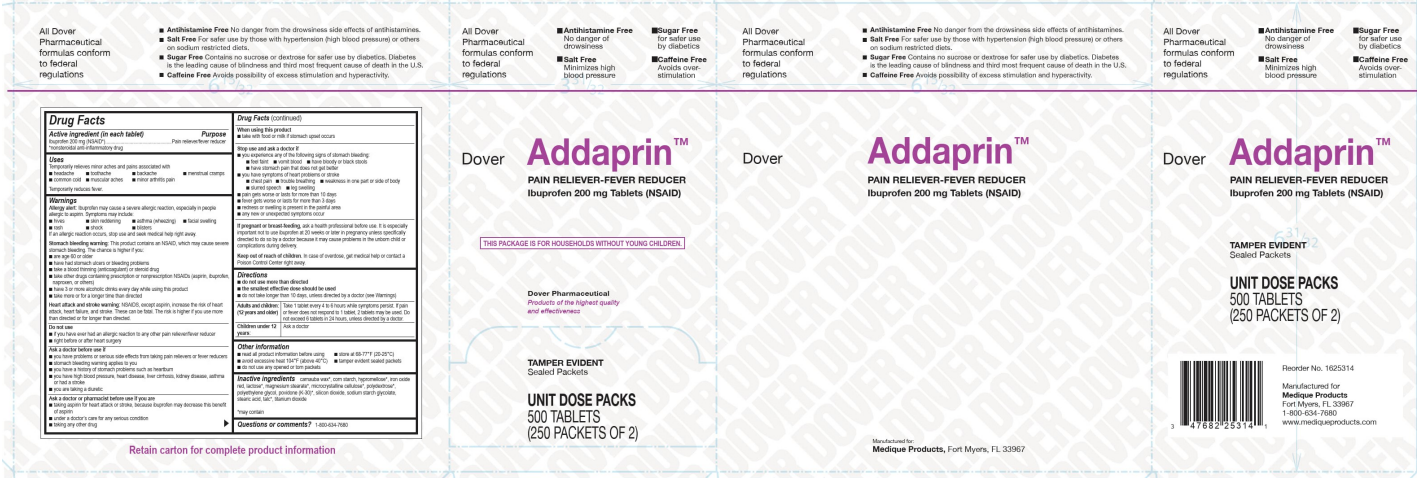
Tamper Evident

Sealed Packets

Unit Dose Packs

500 Tablets

(250 Packets of 2)




## Otis Clapp Ultraprin Label

OC Otis Clapp

Quality & Integrity Since 1840

Ultraprin ™

Pain Reliever-Fever Reducer (NSAID)

Ibuprofen Tablets USP 200 mg

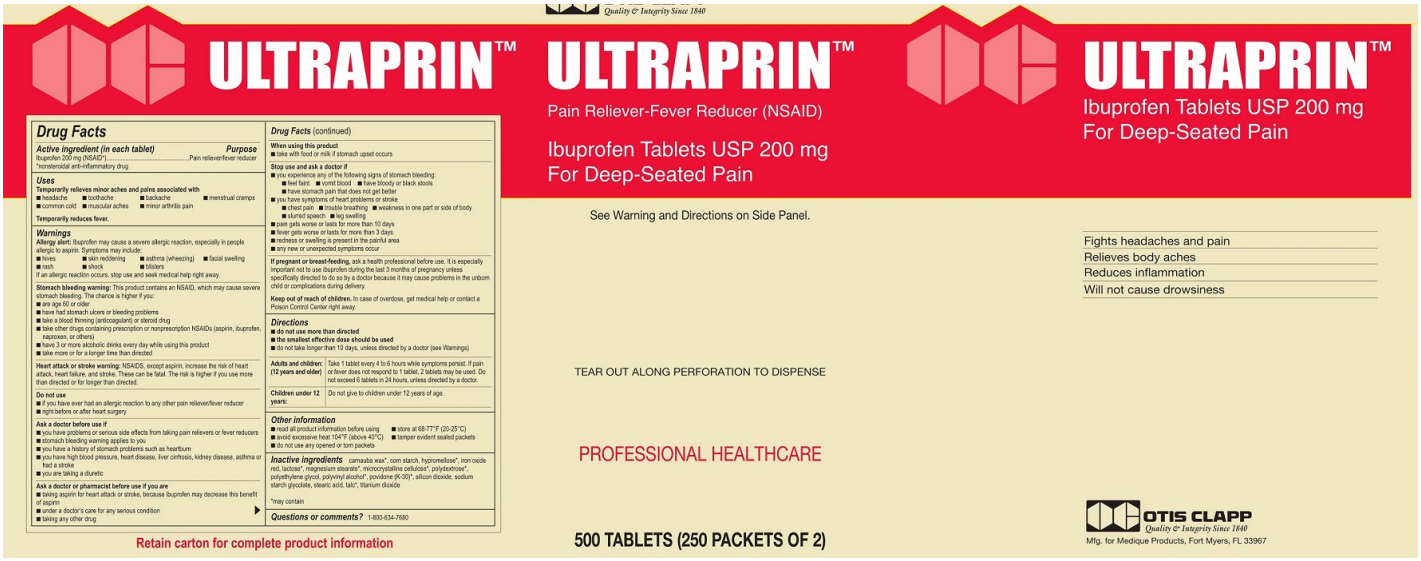
For Deep Seated Pain

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 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) | |
| **POVIDONE K30** (UNII: U725QWY32X) | |

| | |
|---|---|
| **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: 3WQ0SDW1A) | |

## 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 |
|---|---|---|---|---|
| **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 | | |
| **6** | NDC:47682-718-99 | 2 in 1 PACKET; Type 0: Not a Combination Product | 01/26/2017 | |

## 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 | | |

## Active Ingredient/Active Moiety

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

## Inactive Ingredients

| Ingredient Name | Strength |
|---|---|
| **FERRIC OXIDE RED** (UNII: 1K09F3G675) | |
| **POLYETHYLENE GLYCOL, UNSPECIFIED** (UNII: 3WQ0SDW1A) | |
| **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** | | | |

## 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 |
| **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: 3WQ0SDW1A) | |

## 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 |
|---|---|---|---|---|
| **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 | |

| | | | | |
|---|---|---|---|---|
| **4** | | 2 in 1 PACKET; Type 0: Not a Combination Product | | |
| **5** | NDC:47682-700-99 | 2 in 1 PACKET; Type 0: Not a Combination Product | 01/26/2017 | |

## Marketing Information

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

# DOVER ADDAPRIN

ibuprofen tablet, coated

## Product Information

| | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:47682-714 |
| **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) | |
| **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: 3WQ0SDW1A) | |
| **TALC** (UNII: 7SEV7J4R1U) | |

## 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 | Marketing End |
|---|---|---|---|---|

| # | Item Code | Package Description | Date | Date |
|---|---|---|---|---|
| 1 | NDC:47682-714-13 | 250 in 1 BOX | 01/26/2017 | |
| 1 | NDC:47682-714-99 | 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

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

## Inactive Ingredients

| Ingredient Name | Strength |
|---|---|
| **POLYETHYLENE GLYCOL, UNSPECIFIED** (UNII: 3WQ0SDW1A) | |
| **TALC** (UNII: 7SEV7J4R1U) | |
| **POLYVINYL ALCOHOL, UNSPECIFIED** (UNII: 532B59J990) | |
| **SILICON DIOXIDE** (UNII: ETJ7Z6XBU4) | |
| **TITANIUM DIOXIDE** (UNII: 15FIX9V2JP) | |
| **STEARIC ACID** (UNII: 4ELV7Z65AP) | |
| **SODIUM STARCH GLYCOLATE TYPE A POTATO** (UNII: 5856J3G2A2) | |
| **POVIDONE K30** (UNII: U725QWY32X) | |
| **STARCH, CORN** (UNII: O8232NY3SJ) | |
| **FERRIC OXIDE RED** (UNII: 1K09F3G675) | |

## 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 |
|---|---|---|---|---|
| **1** | NDC:47682-702-13 | 250 in 1 BOX | 02/01/2017 | 04/03/2017 |
| **1** | NDC:47682-702-99 | 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 | 11/17/2014 | 04/03/2017 |

## Labeler - Unifirst First Aid Corporation (832947092)

Revised: 9/2024

Unifirst First Aid Corporation