**MEDIQUE DECOREL FORTE- acetaminophen, dextromethorphan hydrobromide, guaifenesin, phenylephrine hydrochloride tablet, film coated**
**MEDI-FIRST COUGH COLD- acetaminophen, dextromethorphan hydrobromide, guaifenesin, phenylephrine hydrochloride tablet, film coated**
**MEDI-FIRST PLUS COUGH COLD- acetaminophen, dextromethorphan hydrobromide, guaifenesin, phenylephrine hydrochloride tablet, film coated**
**Unifirst First Aid Corporation**

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**Medique Decorel Forte**
**MF/MFP Cough & Cold Relief**

***Drug Facts***

***Active ingredients (in each caplet)***

Acetaminophen 325 mg

Dextromethorphan Hydrobromide 10 mg

Guaifenesin 100 mg

Phenylephrine Hydrochloride 5 mg

***Purpose***

Pain reliever/fever reducer

Cough suppressant

Expectorant

Nasal decongestant

***Uses***

Temporarily relieves these symptoms due to the common cold:

- headache
- nasal congestion
- cough
- minor aches and pains
- sore throat
- sinus congestion and pressure
- helps loosen phlegm (mucus) and thin bronchial secretions to drain bronchial tubes and make cough more productive

Temporarily reduces fever.

***Warnings***

**Liver warning:** This product contains acetaminophen. Severe liver damage may occur if you take:

- more than 4,000 mg of acetaminophen in 24 hours
- with other drugs containing acetaminophen
- 3 or more alcoholic drinks every day while using this product

**Allergy alert:** Acetaminophen may cause severe skin reactions. Symptoms may include:

- blisters
- rash
- skin reddening

If a skin reaction occurs, stop use and seek medical help right away.

**Sore throat warning:** If sore throat is severe, persists for more than 2 days, is accompanied or followed by fever, headache, rash, nausea, or vomiting, consult a doctor promptly.

**Do not use**

- with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.

- if you have ever had an allergic reaction to this product or any of its ingredients
- if you are now taking a prescription monoamine oxidase inhibitor (MAOI) (certain drugs for depression, psychiatric or emotional conditions, or Parkinson's disease), or for 2 weeks after stopping the MAOI drug. If you do not know if your prescription drug contains an MAOI, ask a doctor or pharmacist before taking this product.

**Ask a doctor before use if you have**

- liver disease
- heart disease
- high blood pressure
- thyroid disease
- diabetes
- difficulty in urination due to enlargement of the prostate gland
- cough that occurs with too much phlegm (mucus)
- persistent or chronic cough that lasts as occurs with smoking, asthma, chronic bronchitis, or emphysema

**Ask a doctor or pharmacist before use if you are** taking the blood thinning drug warfarin.

**When using this product do not exceed recommended dosage.**

**Stop use and ask a doctor if**

- nervousness, dizziness, or sleeplessness occur
- pain, cough, or nasal congestion gets worse or lasts more than 7 days
- fever gets worse or lasts more than 3 days
- redness or swelling is present
- new symptoms occur
- cough comes back or occurs with rash or headache that lasts.

These could be signs of a serious condition.

**If pregnant or breast-feeding,** ask a health professional before use.

**Keep out of reach of children.**

In case of overdose, get medical help or contact a Poison Control Center right away. Prompt medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

***Directions***

- **do not take more than directed**

**Adults and children: (12 years and over)**

- take 2 caplets every 4 hours
- not to take more than 10 caplets in 24 hours

**Children under 12 years:** ask a doctor

***Other information***

- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
- tamper evident sealed packets
- do not use any opened or torn packets

***Inactive ingredients***

corn starch, crospovidone, FD&C yellow #6 aluminum lake, magnesium stearate, maltodextrin, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, povidone, silicon dioxide, sodium starch glycolate, stearic acid, talc, titanium dioxide

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

Medi-First®

Cough & Cold Relief

✓ Aches, Fever • Acetaminophen 325 mg

✓ Coughs • Dextromethorphan HBr 10 mg

✓ Chest Congestion • Guaifenesin 100 mg

✓ Nasal Congestion • Phenylephrine HCl 5 mg

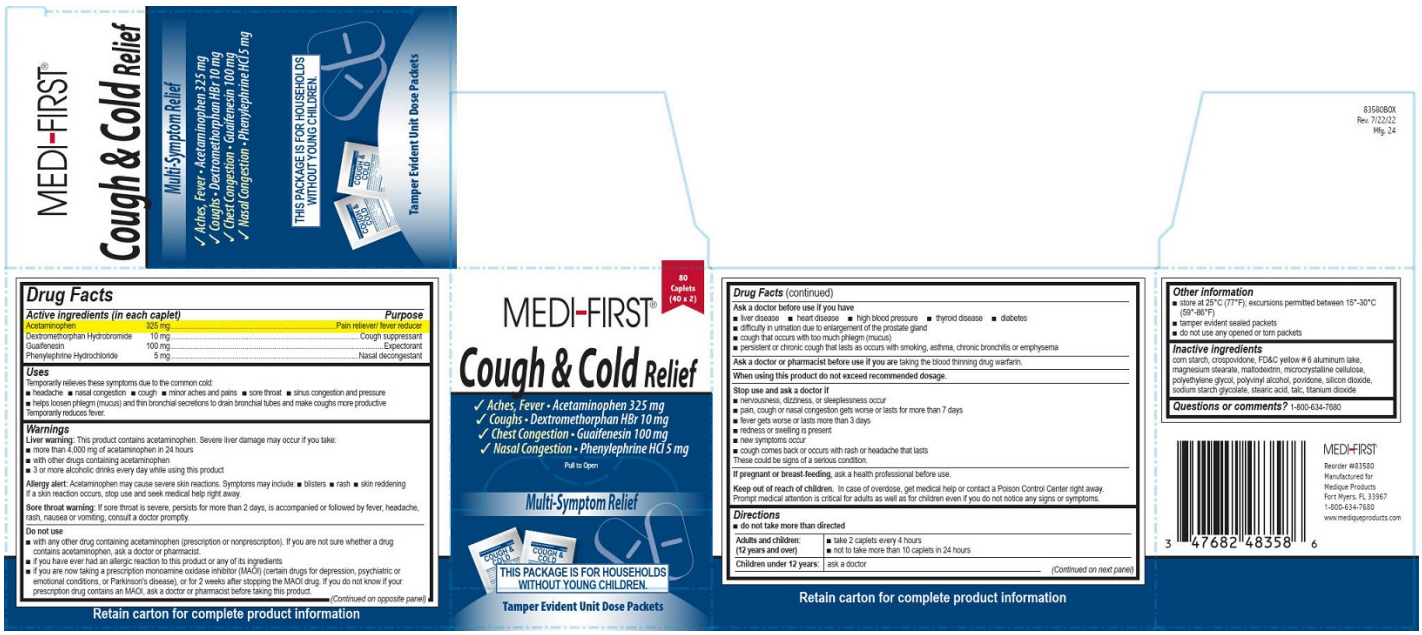
Pull to Open

Multi-Symptom Relief

This Package is for Households Without Young Children.

80 Caplets (40 x 2)

Tamper evident Unit Dose Packets




Medi-First® Plus

Cough & Cold Relief

Aches, Fever • Acetaminophen 325 mg

Coughs • Dextromethorphan HBr 10 mg

Chest Congestion • Guaifenesin 100 mg

Nasal Congestion • Phenylephrine HCl 5 mg

This Package is for Households Without Young Children.

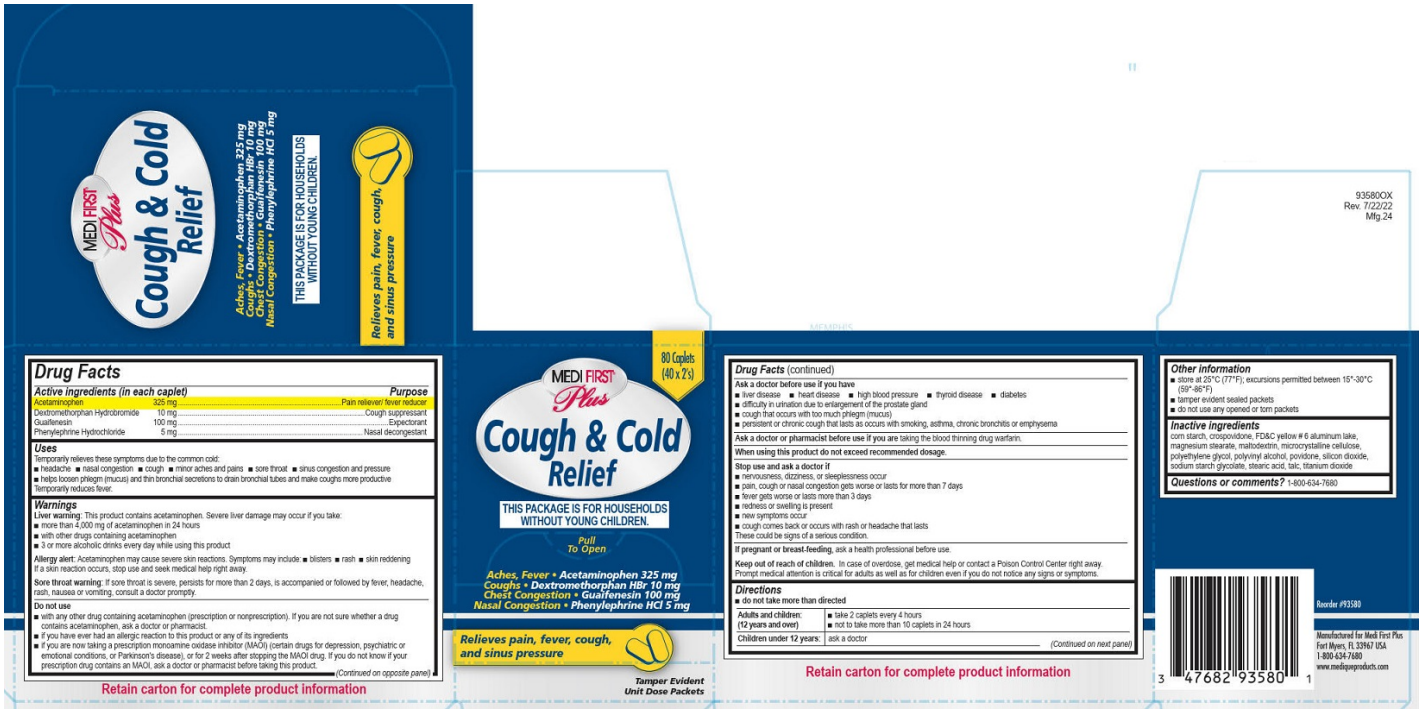
Pull To Open

80 Caplets

(40 x 2's)

Relieves pain, fever, cough, and sinus pressure

Tamper Evident Unit Dose Packets




Medique®

Decorel Forte

✓ Aches, Fever • Acetaminophen 325 mg

✓ Coughs • Dextromethorphan HBr 10 mg

✓ Chest Congestion • Guaifenesin 100 mg

✓ Nasal Congestion • Phenylephrine HCl 5 mg

Severe Cold & Cough Relief

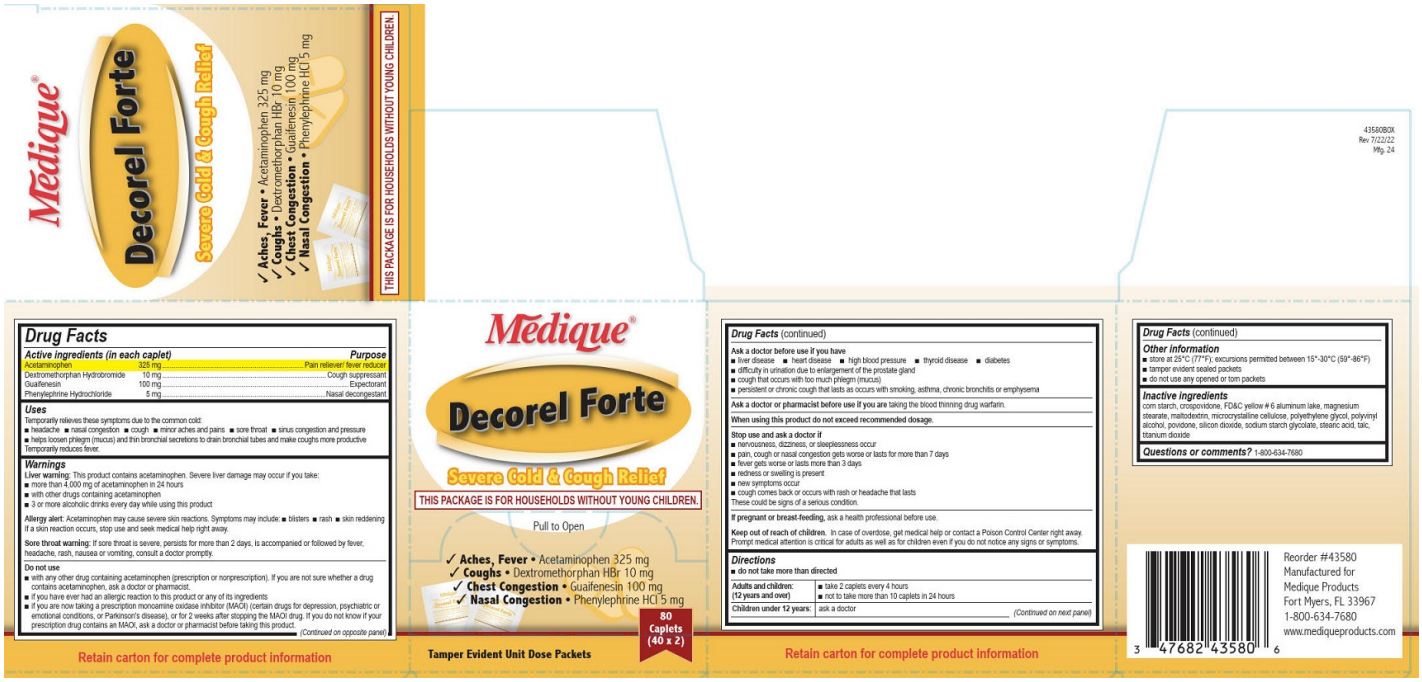
This Package is for Households Without Young Children.

Pull to Open

80 Caplets

(40 x 2)

Tamper Evident Unit Dose Packets




# MEDIQUE DECOREL FORTE

acetaminophen, dextromethorphan hydrobromide, guaifenesin, phenylephrine hydrochloride tablet, film coated

## Product Information

| | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:47682-435 |
| **Route of Administration** | ORAL | | |

## Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|---|---|
| **PHENYLEPHRINE HYDROCHLORIDE** (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV) | PHENYLEPHRINE HYDROCHLORIDE | 5 mg |
| **ACETAMINOPHEN** (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D) | ACETAMINOPHEN | 325 mg |
| **DEXTROMETHORPHAN HYDROBROMIDE** (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS) | DEXTROMETHORPHAN HYDROBROMIDE | 10 mg |
| **GUAIFENESIN** (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ) | GUAIFENESIN | 100 mg |

## Inactive Ingredients

| Ingredient Name | Strength |
|---|---|
| **CROSPOVIDONE** (UNII: 2S7830E561) | |
| **FD&C YELLOW NO. 6** (UNII: H77VEI93A8) | |
| **MALTODEXTRIN** (UNII: 7CVR7L4A2D) | |
| **SILICON DIOXIDE** (UNII: ETJ7Z6XBU4) | |
| **POVIDONE** (UNII: FZ989GH94E) | |
| **STARCH, CORN** (UNII: O8232NY3SJ) | |
| **MAGNESIUM STEARATE** (UNII: 70097M6I30) | |
| **MICROCRYSTALLINE CELLULOSE** (UNII: OP1R32D61U) | |
| **POLYETHYLENE GLYCOL, UNSPECIFIED** (UNII: 3WQ0SDW1A) | |

| | |
|---|---|
| **SODIUM STARCH GLYCOLATE TYPE A POTATO** (UNII: 5856J3G2A2) | |
| **TALC** (UNII: 7SEV7J4R1U) | |
| **TITANIUM DIOXIDE** (UNII: 15FIX9V2JP) | |
| **POLYVINYL ALCOHOL, UNSPECIFIED** (UNII: 532B59J990) | |
| **STEARIC ACID** (UNII: 4ELV7Z65AP) | |

### Product Characteristics

| | | | |
|---|---|---|---|
| **Color** | orange | **Score** | no score |
| **Shape** | CAPSULE (Caplet) | **Size** | 17mm |
| **Flavor** | | **Imprint Code** | 44;546 |
| **Contains** | | | |

### Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:47682-435-80 | 40 in 1 BOX | 09/26/2022 | |
| **1** | NDC:47682-435-99 | 2 in 1 PACKET; Type 0: Not a Combination Product | | |
| **2** | NDC:47682-435-36 | 75 in 1 BOX | 12/12/2022 | |
| **2** | NDC:47682-435-99 | 2 in 1 PACKET; Type 0: Not a Combination Product | | |
| **3** | NDC:47682-435-99 | 2 in 1 PACKET; Type 0: Not a Combination Product | 09/26/2022 | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC Monograph Drug | M012 | 09/26/2022 | |

# MEDI-FIRST COUGH COLD

acetaminophen, dextromethorphan hydrobromide, guaifenesin, phenylephrine hydrochloride tablet, film coated

### Product Information

| | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:47682-835 |
| **Route of Administration** | ORAL | | |

### Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|---|---|
| **PHENYLEPHRINE HYDROCHLORIDE** (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV) | PHENYLEPHRINE HYDROCHLORIDE | 5 mg |
| **ACETAMINOPHEN** (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D) | ACETAMINOPHEN | 325 mg |

| | | |
|---|---|---|
| **DEXTROMETHORPHAN HYDROBROMIDE** (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS) | DEXTROMETHORPHAN HYDROBROMIDE | 10 mg |
| **GUAIFENESIN** (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ) | GUAIFENESIN | 100 mg |

## Inactive Ingredients

| Ingredient Name | Strength |
|---|---|
| **CROSPOVIDONE** (UNII: 2S7830E561) | |
| **FD&C YELLOW NO. 6** (UNII: H77VEI93A8) | |
| **MALTODEXTRIN** (UNII: 7CVR7L4A2D) | |
| **SILICON DIOXIDE** (UNII: ETJ7Z6XBU4) | |
| **POVIDONE** (UNII: FZ989GH94E) | |
| **STARCH, CORN** (UNII: O8232NY3SJ) | |
| **MAGNESIUM STEARATE** (UNII: 70097M6I30) | |
| **MICROCRYSTALLINE CELLULOSE** (UNII: OP1R32D61U) | |
| **POLYETHYLENE GLYCOL, UNSPECIFIED** (UNII: 3WQ0SDW1A) | |
| **SODIUM STARCH GLYCOLATE TYPE A POTATO** (UNII: 5856J3G2A2) | |
| **TALC** (UNII: 7SEV7J4R1U) | |
| **TITANIUM DIOXIDE** (UNII: 15FIX9V2JP) | |
| **STEARIC ACID** (UNII: 4ELV7Z65AP) | |
| **POLYVINYL ALCOHOL, UNSPECIFIED** (UNII: 532B59J990) | |

## Product Characteristics

| | | | |
|---|---|---|---|
| **Color** | orange | **Score** | no score |
| **Shape** | CAPSULE (Caplet) | **Size** | 17mm |
| **Flavor** | | **Imprint Code** | 44;546 |
| **Contains** | | | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:47682-835-80 | 40 in 1 BOX | 09/26/2022 | |
| **1** | NDC:47682-835-99 | 2 in 1 PACKET; Type 0: Not a Combination Product | | |
| **2** | NDC:47682-835-36 | 75 in 1 BOX | 12/12/2022 | |
| **2** | NDC:47682-835-99 | 2 in 1 PACKET; Type 0: Not a Combination Product | | |
| **3** | NDC:47682-835-99 | 2 in 1 PACKET; Type 0: Not a Combination Product | 09/26/2022 | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC Monograph Drug | M012 | 09/26/2022 | |

# MEDI-FIRST PLUS COUGH COLD

acetaminophen, dextromethorphan hydrobromide, guaifenesin, phenylephrine hydrochloride
tablet, film coated

## Product Information

| | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:47682-836 |
| **Route of Administration** | ORAL | | |

## Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|---|---|
| **GUAIFENESIN** (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ) | GUAIFENESIN | 100 mg |
| **PHENYLEPHRINE HYDROCHLORIDE** (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV) | PHENYLEPHRINE HYDROCHLORIDE | 5 mg |
| **ACETAMINOPHEN** (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D) | ACETAMINOPHEN | 325 mg |
| **DEXTROMETHORPHAN HYDROBROMIDE** (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS) | DEXTROMETHORPHAN HYDROBROMIDE | 10 mg |

## Inactive Ingredients

| Ingredient Name | Strength |
|---|---|
| **POLYVINYL ALCOHOL, UNSPECIFIED** (UNII: 532B59J990) | |
| **STEARIC ACID** (UNII: 4ELV7Z65AP) | |
| **CROSPOVIDONE** (UNII: 2S7830E561) | |
| **FD&C YELLOW NO. 6** (UNII: H77VEI93A8) | |
| **MALTODEXTRIN** (UNII: 7CVR7L4A2D) | |
| **SILICON DIOXIDE** (UNII: ETJ7Z6XBU4) | |
| **POVIDONE** (UNII: FZ989GH94E) | |
| **STARCH, CORN** (UNII: O8232NY3SJ) | |
| **MAGNESIUM STEARATE** (UNII: 70097M6I30) | |
| **MICROCRYSTALLINE CELLULOSE** (UNII: OP1R32D61U) | |
| **POLYETHYLENE GLYCOL, UNSPECIFIED** (UNII: 3WQ0SDW1A) | |
| **SODIUM STARCH GLYCOLATE TYPE A POTATO** (UNII: 5856J3G2A2) | |
| **TALC** (UNII: 7SEV7J4R1U) | |
| **TITANIUM DIOXIDE** (UNII: 15FIX9V2JP) | |

## Product Characteristics

| | | | |
|---|---|---|---|
| **Color** | orange | **Score** | no score |
| **Shape** | CAPSULE (Caplet) | **Size** | 17mm |
| **Flavor** | | **Imprint Code** | 44;546 |
| **Contains** | | | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| 1 | NDC:47682-836- | 40 in 1 BOX | 09/26/2022 | |

| | | | | |
|---|---|---|---|---|
| **1** | 80 | 40 in 1 BOX | 09/26/2022 | |
| **1** | NDC:47682-836-99 | 2 in 1 PACKET; Type 0: Not a Combination Product | | |
| **2** | NDC:47682-836-36 | 75 in 1 BOX | 12/12/2022 | |
| **2** | NDC:47682-836-99 | 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 |
|---|---|---|---|
| OTC Monograph Drug | M012 | 09/26/2022 | |

## Labeler - Unifirst First Aid Corporation (832947092)

Revised: 9/2024 Unifirst First Aid Corporation