DOC IN THE BOX ALL DAY SUPPLY- acetaminophen, calcium carbonate, dextromethorphan hbr, guaifenesin, phenylephrine hcl, ibuprofen, loperamide hcl
Doc in the Box LLC

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

DOC IN THE BOX ALL DAY SUPPLY

EXTRA STRENGTH NON-ASPIRIN

Purpose

Pain reliever/fever reducer

EXTRA STRENGTH NON-ASPIRIN

Active ingredient (in each tablet)

Acetaminophen 500 mg

EXTRA STRENGTH NON-ASPIRIN

Uses

For the temporary relief of minor aches and pains associated with

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

For the reduction of fever.

EXTRA STRENGTH NON-ASPIRIN

Warnings

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

- more than 8 tablets in 24 hours, which is the maximum daily amount
- 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:

- skin reddening
- blisters
- rash

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

EXTRA STRENGTH NON-ASPIRIN

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.
- for more than 10 days for pain unless directed by a doctor
- for more than 3 days for fever unless directed by a doctor

EXTRA STRENGTH NON-ASPIRIN

Ask a doctor before use if you have liver disease

EXTRA STRENGTH NON-ASPIRIN

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

EXTRA STRENGTH NON-ASPIRIN

Stop using and ask a doctor if
- symptoms do not improve
- new symptoms occur
- pain or fever persists or gets worse
- redness or swelling is present

EXTRA STRENGTH NON-ASPIRIN

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

EXTRA STRENGTH NON-ASPIRIN

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.

EXTRA STRENGH NON-ASPIRIN

Directions
- **do not use more than directed**

Adults and children: (12 years and older)	Take 2 tablets with water every 4 to 6 hours as needed. Do not take more than 8 tablets in 24 hours.
Children under 12 years:	Do not give this adult strength product to children under 12 years of age; this will provide more than the recommended dose (overdose) and may cause liver damage.

EXTRA STRENGTH NON-ASPIRIN

Other information
- store at room temperature 59° - 86°F (15° - 30°C)
- tamper evident sealed packets
- do not use any opened or torn packets

EXTRA STRENGTH NON-ASPIRIN

Inactive ingredients

corn starch, hypromellose, maltodextrin*, microcrystalline cellulose*, polyethylene glycol, povidone*, pregelatinized starch*, sodium starch glycolate*, stearic acid, titanium dioxide*.
* may contain

EXTRA STRENGTH NON-ASPIRIN

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

ANTACID

Active ingredient (in each tablet)

Calcium Carbonate 420 mg

ANTACID

Purpose

Antacid

ANTACID

Uses

For the relief of the following symptoms associated with
- acid indigestion
- sour stomach
- heartburn
- upset stomach

ANTACID

Warnings

ANTACID

Ask a doctor or health professional before use if you have
- been taking a prescription drug. Antacids may interact with certain prescription drugs
- kidney stones
- a calcium-restricted diet

ANTACID

Stop using this product and ask a doctor if symptoms last more than 2 weeks

ANTACID

Do not exceed recommended dosage.

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

ANTACID

Keep out of the reach of children.

ANTACID

Directions

- **do not use more than directed**
- **Adults and children (12 years and older):** Chew 2 tablets every 2 or 3 hours as symptoms occur or as directed by a physician. Do not take more than 19 tablets in a 24 hour period, or use the maximum dosage of this product for more than 2 weeks, except under the advice and supervision of a physician.
- **Children under 12 years:** Do not give to children under 12 years of age.

ANTACID

Other information

- Phenylketonurics: contains phenylalanine 1.5 mg per tablet
- each tablet contains 168 mg of elemental calcium
- store at room temperature 59° - 86°F (15° - 30°C)
- tamper-evident sealed packets
- do not use any opened or torn packets

ANTACID

Inactive ingredients

aspartame*, croscarmellose sodium*, gum acacia*, magnesium stearate, maltodextrin, mineral oil*, mint flavor, sorbitol*, sucrose*.
* may contain

ANTACID

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

COLD RELIEF

Active ingredient (in each tablet)

Acetaminophen 325 mg

Dextromethorphan Hydrobromide 15 mg

Guaifenesin 200 mg

Phenylephrine Hydrochloride 5 mg

COLD RELIEF

Purpose

Pain reliever/fever reducer

Cough suppressant

Expectorant

Nasal decongestant

COLD RELIEF

Uses

Temporarily relieves these cold symptoms

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

Temporarily reduces fever.

COLD RELIEF

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:

- skin reddening
- blisters
- rash

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.

COLD RELIEF

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.

COLD RELIEF

- liver disease
- heart disease
- high blood pressure
- thyroid disease
- diabetes

- trouble urinating due to an enlarged 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

COLD RELIEF

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

COLD RELIEF

When using this product
- do not use more than directed

COLD RELIEF

Stop use and ask a doctor if
- new symptoms occur
- redness or swelling is present
- pain or nasal congestion gets worse or lasts for more than 7 days
- fever gets worse or lasts for more than 3 days
- you get nervous, dizzy or sleepless
- cough comes back or occurs with rash or headache that lasts.

These could be signs of a serious condition.

COLD RELIEF

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

COLD RELIEF

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.

COLD RELIEF

Directions

Adults and children: (12 years and older)	Take 2 tablets with water every 6-8 hours as needed. Do not take more than 8 tablets in 24 hours.
Children under 12 years:	Do not give to children under 12 years of age.

COLD RELIEF

Other information
- store at room temperature 59° - 86°F (15° - 30°C)
- avoid excessive heat and humidity
- tamper evident sealed packets

- do not use any opened or torn packets

COLD RELIEF

Inactive ingredients maltodextrin, microcrystalline cellulose, povidone, sodium starch glycolate, starch, stearic acid

COLD RELIEF

Questions?

1-800-634-7680

IBUPROFEN

Active ingredient (in each tablet)

Ibuprofen 200 mg (NSAID)*

*non-steroidal anti-inflammatory drug

IBUPROFEN

Purpose

Pain reliever/fever reducer

IBUPROFEN

Uses

Temporarily relieves minor aches and pains associated with

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

Temporarily reduces fever.

IBUPROFEN

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
- taking 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

IBUPROFEN

Do not use

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

IBUPROFEN

Ask a doctor before use if

- 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, or kidney disease
- you are taking a diuretic
- you have problems or serious side effects from taking pain relievers or fever reducers
- you have asthma

IBUPROFEN

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
- taking any other drug
- under a doctor’s care for any serious condition

IBUPROFEN

When using this product

- the risk of heart attack or stroke may increase if you use more than directed or longer than directed
- take with food or milk if stomach upset occurs

IBUPROFEN

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

- 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

IBUPROFEN

If pregnant or breast-feeding, ask a health professional before use. It is especially important not to use ibuprofen during the last 3 months of pregnancy unless specifically directed to do so by a doctor because it may cause problems in the unborn child or complications during delivery.

IBUPROFEN

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

IBUPROFEN

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 for 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:	Do not give to children under 12 years of age.

IBUPROFEN

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 packet

IBUPROFEN

Inactive ingredients

carnauba wax*, corn starch, hypromellose*, iron oxide red, lactose*, magnesium stearate*, microcrystalline cellulose*, polydextrose*, polyethylene glycol, polyvinyl alcohol*, povidone (K-30)*, silicon dioxide, sodium starch glycolate, stearic acid, talc*, titanium dioxide

* may contain

IBUPROFEN

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

DIAMODE

Active ingredient (in each caplet)

Loperamide Hydrochloride 2 mg

DIAMODE

Purpose

Antidiarrheal

DIAMODE

Uses

Controls the symptoms of diarrhea, including Traveler's diarrhea

DIAMODE

Warnings

Allergy alert: Do not use if you have ever had a rash or other allergic reaction to Loperamide HCl

DIAMODE

Do not use if you have bloody or black stool

DIAMODE

Ask a doctor before use if you have

- a fever
- mucus in stool
- a history of liver disease

DIAMODE

Ask a doctor or pharmacist before use if you are taking antibiotics

DIAMODE

When using this product

- tiredness, drowsiness or dizziness may occur
- be careful when driving or operating machinery

DIAMODE

Stop use and ask a doctor if

- symptoms get worse
- diarrhea lasts more than 2 days
- upi get abdominal swelling or bulging. These may be signs of a serious condition.

DIAMODE

If you are pregnant or breast-feeding, ask a health professional before use

DIAMODE

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

DIAMODE

Directions

- do not use more than directed
- drink plenty of clear fluids to help prevent dehydration caused by diarrhea

Adults and children: (12 years and older)	Take 2 caplets after the first loose stool followed by 1 caplet after each subsequent loose stool but no more than 4 caplets in 24 hours.
Children under 12 years:	Do not give to children under 12 years of age.

DIAMODE

Other information

- store at room temperature 68° - 77°F (20° - 25°C)
- tamper-evident sealed packets
- do not use any opened or torn packet

DIAMODE

Inactive ingredients anhydrous lactose, croscarmellose sodium, crospovidone, D&C Yellow #10, FD&C Blue #1, hydrogenated vegetable oil, magnesium stearate, powdered cellulose, pregelatinized starch

DIAMODE

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

EXTRA STRENGTH NON-ASPIRIN

EXTRA STRENGTH

NON-ASPIRIN

2 Tablets

Mfd. for ***MEDIQUE PRODUCTS • Fort Myers, FL 33967***

ANTACID

ANTACID

2 Tablets

Mfd for MEDIQUE PRODUCTS, Fort Myers, FL 33967

COLD RELIEF

COLD RELIEF

2 Tablets

Mfd for MEDIQUE PRODUCTS, Fort Myers, FL 33967

IBUPROFEN

IBUPROFEN

2 Tablets

Mfd. for: ***MEDIQUE PRODUCTS*** *• Fort Myers, FL 33967*

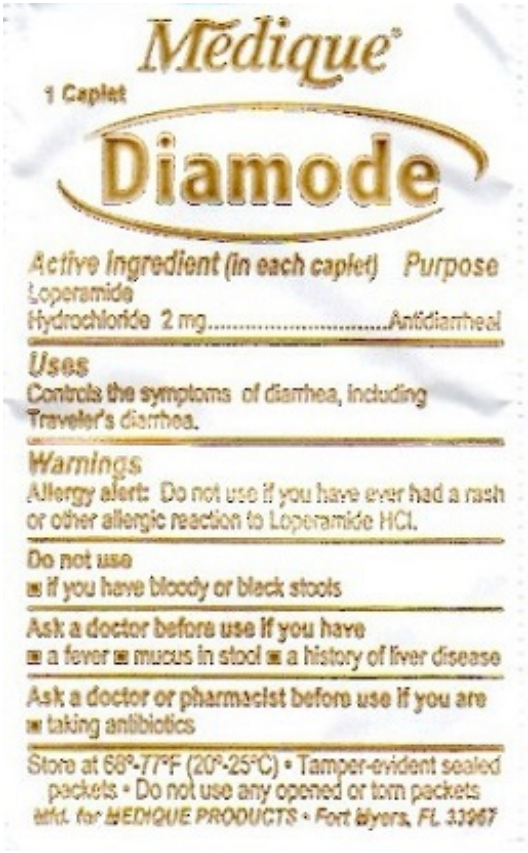
DIAMODE

Medique

Diamode

1 Caplet

Mfd. for MEDIQUE PRODUCTS, Fort Myers, FL 33967



OUTER KIT CARTON

DOC IN THE BOX

ALL DAY SUPPLY...and More!

5 Different OTC Medications

17 Individual Packets

Headache, Toothache, Diarrhea, Cough/Cold, Upset Stomach, Heartburn, Muscle Aches

ALL PRODUCTS ARE PACKAGED IN TAMPER EVIDENT PACKETS. DO NOT USE IF ANY PACKETS ARE OPEN OR TORN.

KIT CONTENTS	***Drug Facts*** ***Active ingredients (in each dosage unit***	***Purposes***
NON-ASPIRIN TABLETS (3 Packets, 2 Tablets Each)	Acetaminophen 500 mg	Pain reliever/Fever reducer
ANTACID TABLETS (4 Packets, 2 Chewable Tablets Each)	Calcium Carbonate 420 mg	Antacid
COLD RELIEF TABLETS (3 Packets 2 Tablets	Acetaminophen 325 mg Dextromethorphan hydrobromide 15 mg	Pain reliever/Fever reducer Cough suppressant

Each)	Guaifenesin 200 mg Phenylephrine hydrochloride 5 mg	Cough suppressant Expectorant Nasal decongestant
IBUPROFEN TABLETS (3 Packets 2 Tablets Each)	Ibuprofen 200 mg (NSAID)	Pain reliever/Fever reducer
DIAMODE TABLETS (4Packets 1 Caplet Each)	Loperamide hydrochloride 2 mg	Antidiarrheal

SEE ENCLOSED PACKAGE INSERT FOR COMPLETE DRUG FACTS

Distributed by: DocintheBox, LLC, Florence, NJ 08518 • www.docinthebox.com

DOC IN THE BOX ALL DAY SUPPLY

acetaminophen, calcium carbonate, dextromethorphan hbr, guaifenesin, phenylephrine hcl, ibuprofen, loperamide hcl kit

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72082-001

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72082-001-05	1 in 1 BOX; Type 1: Convenience Kit of Co-Package	01/01/2019	

Quantity of Parts		
Part #	**Package Quantity**	**Total Product Quantity**
Part 1	2 PACKET	3 in 2
Part 2	2 PACKET	4 in 2
Part 3	2 PACKET	3 in 2
Part 4	2 PACKET	3 in 2
Part 5	4 PACKET	4

Part 1 of 5

MEDI-FIRST NON-ASPIRIN EXTRA STRENGTH

acetaminophen tablet, film coated

Product Information	
Item Code (Source)	NDC:47682-126
Route of Administration	ORAL

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	500 mg

Inactive Ingredients	
Ingredient Name	**Strength**
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
STARCH, CORN (UNII: O8232NY3SJ)	
POVIDONE (UNII: FZ989GH94E)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

Product Characteristics			
Color	white (white)	**Score**	no score
Shape	ROUND (Round)	**Size**	12mm
Flavor		**Imprint Code**	FR;33
Contains			

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

1	NDC:47682-126-99	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 not final	part343	01/01/2019	

Part 2 of 5

MEDI-FIRST ANTACID

calcium carbonate tablet, chewable

Product Information

Item Code (Source)	NDC:47682-820
Route of Administration	ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CALCIUM CARBONATE (UNII: H0G9379FGK) (CALCIUM CATION - UNII:2M83C4R6ZB, CARBONATE ION - UNII:7UJQ5OPE7D)	CALCIUM CARBONATE	420 mg

Inactive Ingredients

Ingredient Name	Strength
SUCROSE (UNII: C151H8M554)	
ASPARTAME (UNII: Z0H242BBR1)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
ACACIA (UNII: 5C5403N26O)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
SORBITOL (UNII: 506T60A25R)	
MINERAL OIL (UNII: T5L8T28FGP)	

Product Characteristics

Color	white	**Score**	no score
Shape	ROUND	**Size**	11mm
Flavor		**Imprint Code**	AZ;036
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47682-820-99	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 final	part331	01/01/2019	

Part 3 of 5

MEDI-FIRST COLD RELIEF

acetaminophen, dextromethorphan hydrobromide, guaifenesin, phenylephrine hydrochloride tablet

Product Information

Item Code (Source)	NDC:47682-139
Route of Administration	ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	15 mg
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	200 mg

Inactive Ingredients

Ingredient Name	Strength
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
POVIDONE (UNII: FZ989GH94E)	
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)	
STARCH, CORN (UNII: O8232NY3SJ)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

Product Characteristics

Color	white (white)	**Score**	no score
Shape	ROUND (round)	**Size**	12mm
Flavor		**Imprint Code**	FR;12
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47682-139-99	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 final	part341	01/01/2019	

Part 4 of 5

MEDI-FIRST IBUPROFEN

ibuprofen tablet, coated

Product Information

Item Code (Source)	NDC:47682-708
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
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
POVIDONE K30 (UNII: U725QWY32X)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CARNAUBA WAX (UNII: R12CBM0EIZ)	
STARCH, CORN (UNII: O8232NY3SJ)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
POLYDEXTROSE (UNII: VH2XOU12IE)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
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:47682-708-99	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	01/01/2019	

Part 5 of 5

MEDIQUE DIAMODE

loperamide hydrochloride tablet

Product Information	
Item Code (Source)	NDC:47682-200
Route of Administration	ORAL

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
LOPERAMIDE HYDROCHLORIDE (UNII: 77TI35393C) (LOPERAMIDE - UNII:6X9OC3H4II)	LOPERAMIDE HYDROCHLORIDE	2 mg

Inactive Ingredients	
Ingredient Name	Strength
STARCH, CORN (UNII: O8232NY3SJ)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
ANHYDROUS LACTOSE (UNII: 3SY5LH9PMK)	
CROSPOVIDONE (UNII: 2S7830E561)	
CORN OIL (UNII: 8470G57WFM)	
POWDERED CELLULOSE (UNII: SMD1X3XO9M)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	

Product Characteristics			
Color	green	Score	2 pieces
Shape	OVAL	Size	10mm
Flavor		Imprint Code	123

Contains	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47682-200-46	1 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	ANDA074091	01/01/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part343	01/01/2019	

Labeler - Doc in the Box LLC (081033259)

Establishment

Name	Address	ID/FEI	Business Operations
Doc in the Box LLC		081033259	repack(72082-001)

Revised: 1/2019 Doc in the Box LLC