IBUPROFEN 100- ibuprofen tablet, coated
IBUPROFEN 250- ibuprofen tablet, coated
JHK Inc dba American Safety & First Aid

IBUPROFEN

Drug Facts

Active ingredient (In Each Tablets)

Ibuprofen, USP 200 mg (NSAID)

*Nonsteroidal Anti-Inflammatory Drug

Purpose

pain reliever/fever reducer

Use

temporarily relieves fever and minor aches and pains associated with:
- common cold
- backache
- headache
- toothache
- arthritis
- muscular aches
- menstrual cramps

Warnings

Allergy Alert

Ibuprofen may cause a severe allergic reaction, especially in people allergic to aspirin. Symptoms may include:
- hives
- asthma (wheezing)
- facial swelling
- shock
- skin reddening
- rash
- blisters
- If an allergic reaction occurs, stop use and seek medical help right away.

Stomach bleeding warning

This product contains a nonsteroidal anti-inflammatory drug (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

Do not use
- if you ever had an allergic reaction to any other pain reliever/fever reducer
- right before or after heart surgery

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

Ask a doctor or pharmacist before use if you are
- taking any other drug containing an NSAID (prescription or nonprescription)
- taking a blood thinning (anticoagulant) drug
- 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
- long term continuous use may increase the risk of heart attack or stroke

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 more than 10 days
- fever gets worse or lasts more than 3 days
- redness or swelling is present in the painful area
- any new symptoms appear

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 DEFINITELY 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 take 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: 1 tablet every 4 to 6 hours while symptoms occur; if pain or fever does not respond to 1 tablet, 2 tablets may be used, but do not exceed 6 tablets in 24 hours unless directed by a doctor.

Children under 12 years: Ask a doctor.

Other Information

- do not use if packet is opened or torn
- Store at 20-25°C (68-77°F) avoid excessive heat above 40°C (104°F)

Inactive Ingredients

crosscarmellose sodium[1], FD& C Blue #2[1], FD& C red #40[1], FD& C Yellow #6[1], hypromellose[1], iron oxide red[1], lactose monohydrate[1], magnesium stearate[1], maltodextrin[1], medium chain triglycerides[1], microcrystalline cellulose[1], polydextrose[1], polyethylene glycol[1], polyvinyl alcohol[1], povidone (K-30)[1], silicon dioxide, sodium starch glycolate[1], starch, stearic acid[1], talc[1], titanium dioxide, triacetin[1],

---

1 **contains one or more of these ingredients**

Questions?

1-866-651-3660 Monday-Friday

8:00am-5:00pm EST

PRINCIPAL DISPLAY PANEL - 100 Tablet Packet Box

This product is not manufactured or distributed by
Pfizer Consumer Healthcare
Owner of the registered trademark Advil®.

Product #1102

Ibuprofen

Pain Reliever/Fever Reducer

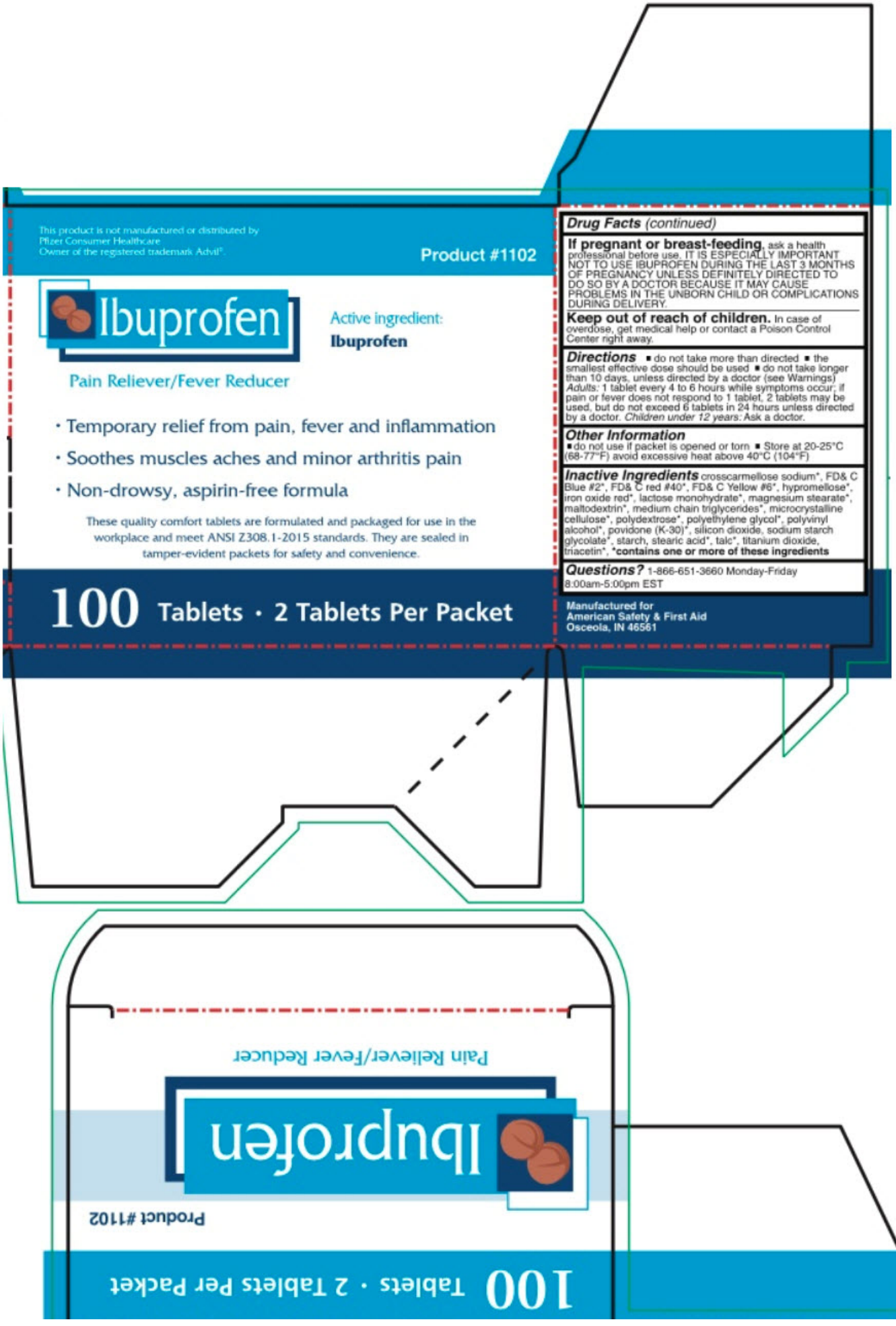
Active ingredient:
Ibuprofen

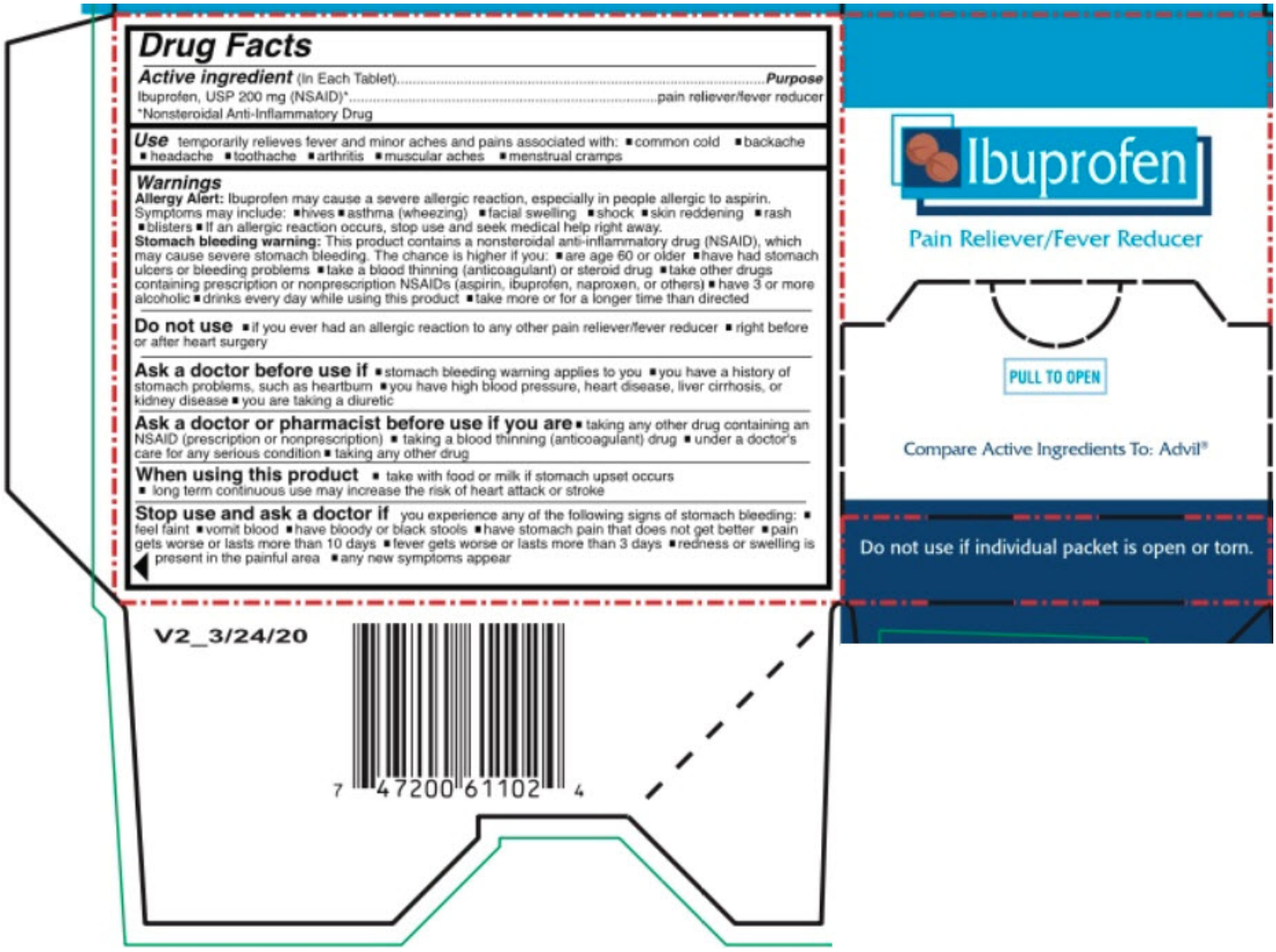
- Temporary relief from pain, fever and inflammation
- Soothes muscles aches and minor arthritis pain
- Non-drowsy, aspirin-free formula

These quality comfort tablets are formulated and packaged for use in the

workplace and meet ANSI Z308.1-2015 standards. They are sealed in tamper-evident packets for safety and convenience.

100 Tablets • 2 Tablets Per Packet


This product is not manufactured or distributed by
Pfizer Consumer Healthcare
Owner of the registered trademark Advil®.
Product #1102
Ibuprofen
Active ingredient:
Ibuprofen
Pain Reliever/Fever Reducer
• Temporary relief from pain, fever and inflammation
• Soothes muscles aches and minor arthritis pain
• Non-drowsy, aspirin-free formula
These quality comfort tablets are formulated and packaged for use in the workplace and meet ANSI Z308.1-2015 standards. They are sealed in tamper-evident packets for safety and convenience.
100 Tablets • 2 Tablets Per Packet
Drug Facts (continued)
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 DEFINITELY 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 take 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: 1 tablet every 4 to 6 hours while symptoms occur; if pain or fever does not respond to 1 tablet, 2 tablets may be used, but do not exceed 6 tablets in 24 hours unless directed by a doctor. Children under 12 years: Ask a doctor.
Other Information
▪ do not use if packet is opened or torn ▪ Store at 20-25°C (68-77°F) avoid excessive heat above 40°C (104°F)
Inactive Ingredients crosscarmellose sodium*, FD& C Blue #2*, FD& C red #40*, FD& C Yellow #6*, hypromellose*, iron oxide red*, lactose monohydrate*, magnesium stearate*, maltodextrin*, medium chain triglycerides*, microcrystalline cellulose*, polydextrose*, polyethylene glycol*, polyvinyl alcohol*, povidone (K-30)*, silicon dioxide, sodium starch glycolate*, starch, stearic acid*, talc*, titanium dioxide, triacetin*, *contains one or more of these ingredients
Questions? 1-866-651-3660 Monday-Friday 8:00am-5:00pm EST
Manufactured for
American Safety & First Aid
Osceola, IN 46561
Pain Reliever/Fever Reducer
Ibuprofen
Product #1102
100 Tablets • 2 Tablets Per Packet



PRINCIPAL DISPLAY PANEL - 250 Tablet Packet Box

Product #1104

Ibuprofen

Pain Reliever/Fever Reducer

Active ingredient:
Ibuprofen

- Temporary relief from pain, fever and inflammation
- Soothes muscles aches and minor arthritis pain
- Non-drowsy, aspirin-free formula

These quality comfort tablets are formulated and packaged for use in the workplace and meet ANSI Z308.1-2015 standards. They are sealed in tamper-evident packets for safety and convenience.

250 Tablets • 2 Tablets Per Packet


This product is not manufactured or distributed by
Pfizer Consumer Healthcare
Owner of the registered trademark Advil®.
Product #1104
Ibuprofen
Active ingredient:
Ibuprofen
Pain Reliever/Fever Reducer
• Temporary relief from pain, fever and inflammation
• Soothes muscles aches and minor arthritis pain
• Non-drowsy, aspirin-free formula
These quality comfort tablets are formulated and packaged for use in the workplace and meet ANSI Z308.1-2015 standards. They are sealed in tamper-evident packets for safety and convenience.
250 Tablets • 2 Tablets Per Packet
Ibuprofen
Pain Reliever/Fever Reducer
• Temporary relief from pain, fever and inflammation
• Soothes muscle aches and minor arthritis pain
• Non-drowsy, aspirin-free formula
Manufactured for
American Safety & First Aid
Osceola, IN 46561
Pain Reliever/Fever Reducer
Ibuprofen

IBUPROFEN 100

ibuprofen tablet, coated

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73598-1102
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
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
FD&C BLUE NO. 2 (UNII: L06K8R7DQK)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYDEXTROSE (UNII: VH2XOU12IE)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
POVIDONE K30 (UNII: U725QWY32X)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM STARCH GLYCOLATE TYPE A (UNII: H8AV0SQX4D)	
STARCH, CORN (UNII: O8232NY3SJ)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
TRIACETIN (UNII: XHX3C3X673)	

Product Characteristics

Color	BROWN	**Score**	2 pieces
Shape	ROUND	**Size**	10mm
Flavor		**Imprint Code**	G2
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73598-1102-1	50 in 1 BOX	02/02/2000	
1		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	ANDA072096	02/02/2000	

IBUPROFEN 250

ibuprofen tablet, coated

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73598-1104
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
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
FD&C BLUE NO. 2 (UNII: L06K8R7DQK)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYDEXTROSE (UNII: VH2XOU12IE)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
POVIDONE K30 (UNII: U725QWY32X)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM STARCH GLYCOLATE TYPE A (UNII: H8AV0SQX4D)	
STARCH, CORN (UNII: O8232NY3SJ)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
TRIACETIN (UNII: XHX3C3X673)	

Product Characteristics

Color	BROWN	**Score**	2 pieces
Shape	ROUND	**Size**	10mm

Flavor		**Imprint Code**	G2
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73598-1104-1	125 in 1 BOX	02/02/2000	
1		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	ANDA072096	02/02/2000	

Labeler - JHK Inc dba American Safety & First Aid (867236309)

Revised: 4/2024 JHK Inc dba American Safety & First Aid