MOTRIN DUAL ACTION WITH TYLENOL- acetaminophen and ibuprofen tablet, film coated
Johnson & Johnson Consumer Inc.

Motrin® Dual Action With Tylenol®

Drug Facts

Active ingredients (in each tablet)

Acetaminophen 250 mg

Ibuprofen 125 mg (NSAID *)

Purposes

Pain reliever

Pain reliever

Uses

■ temporarily relieves minor aches and pains due to:

■ headache

■ toothache

■ backache

■ menstrual cramps

■ muscular aches

■ minor pain of arthritis

Warnings

Liver damage warning:

This product contains acetaminophen. Severe liver damage may occur if you take:
■ with other drugs containing acetaminophen
■ more than 6 tablets in 24 hours, which is the maximum daily amount for this product
■ 3 or more alcoholic drinks every day while using this product

Allergy alert

Acetaminophen allergy alert: may cause severe skin reactions.
Symptoms may include:
■ skin reddening ■ blisters ■ rash
If skin reaction occurs, stop use and seek medical help right away.

NSAID allergy alert: ibuprofen may cause a severe allergic reaction, especially in people allergic to aspirin.
Symptoms may include:
■ hives ■ facial swelling ■ asthma (wheezing)
■ shock ■ skin reddening ■ rash ■ 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 and 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

■ 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 acetaminophen or any other pain reliever
■ right before or after heart surgery

Ask a doctor before use if

■ you have liver disease
■ stomach bleeding warning applies to you
■ you have problems or serious side effects from taking pain relievers
■ 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

■ under a doctor’s care for any serious condition
■ taking aspirin for heart attack or stroke, because ibuprofen may decrease this benefit of aspirin
■ 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 more than 10 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 at 20 weeks or later in 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. (1-800-222-1222).

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 tablets every 8 hours** while symptoms persist
children under 12 years	■ ask a doctor

■ do not take more than 6 tablets in 24 hours, unless directed by a doctor

Other information

■ read all warnings and directions before use. Keep carton.
■ store at 20-25°C (68-77°F)
■ avoid excessive heat above 40°C (104°F)

Inactive ingredients

colloidal silicon dioxide, corn starch, croscarmellose sodium, crospovidone, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, povidone, pregelatinized

starch, stearic acid, sucralose, talc, titanium dioxide

Questions?

Call **1-877-895-3665** (toll-free) or **215-273-8755** (collect) or visit www.motrin.com

PRINCIPAL DISPLAY PANEL

NDC 50580-208-12

NEW

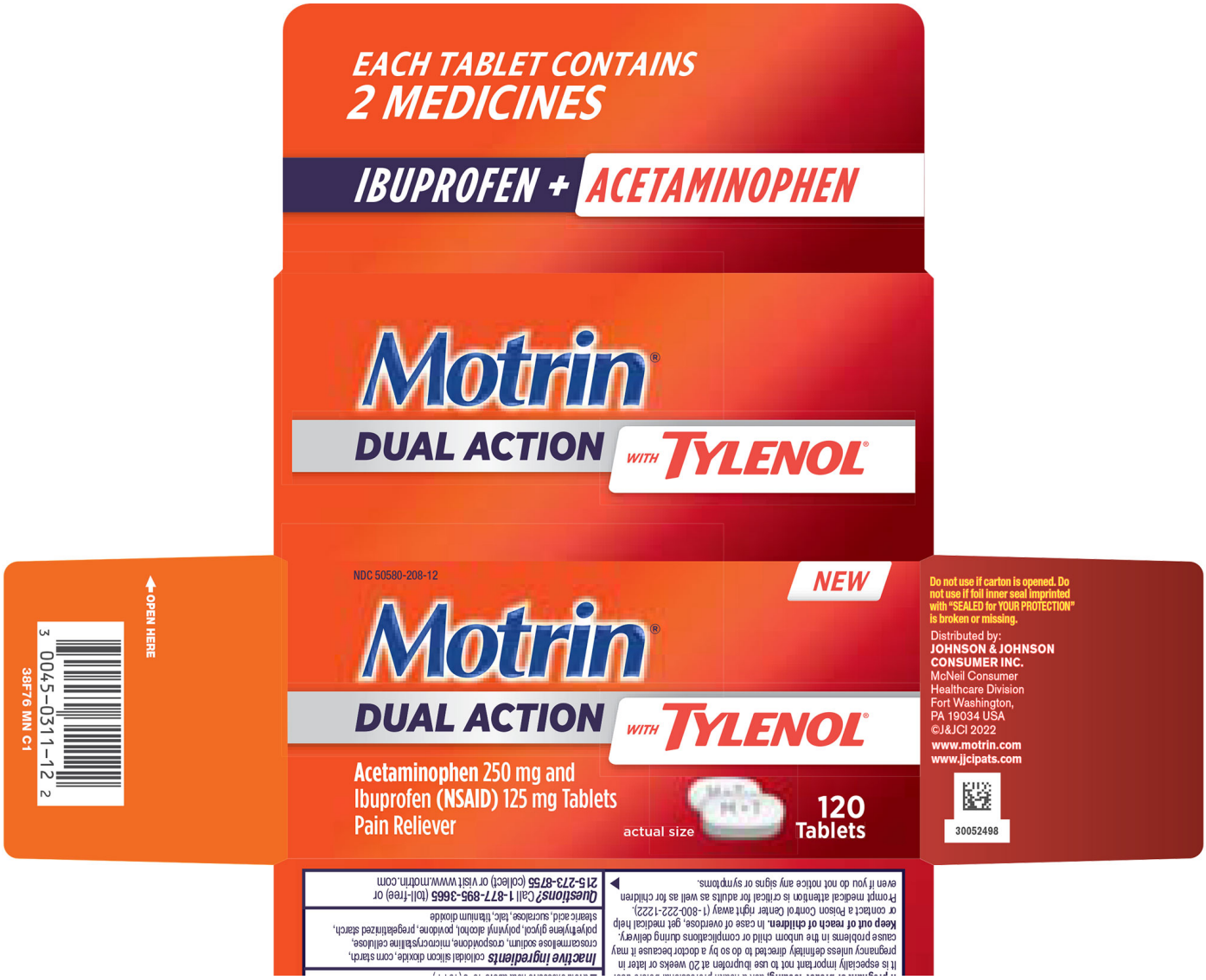
Motrin

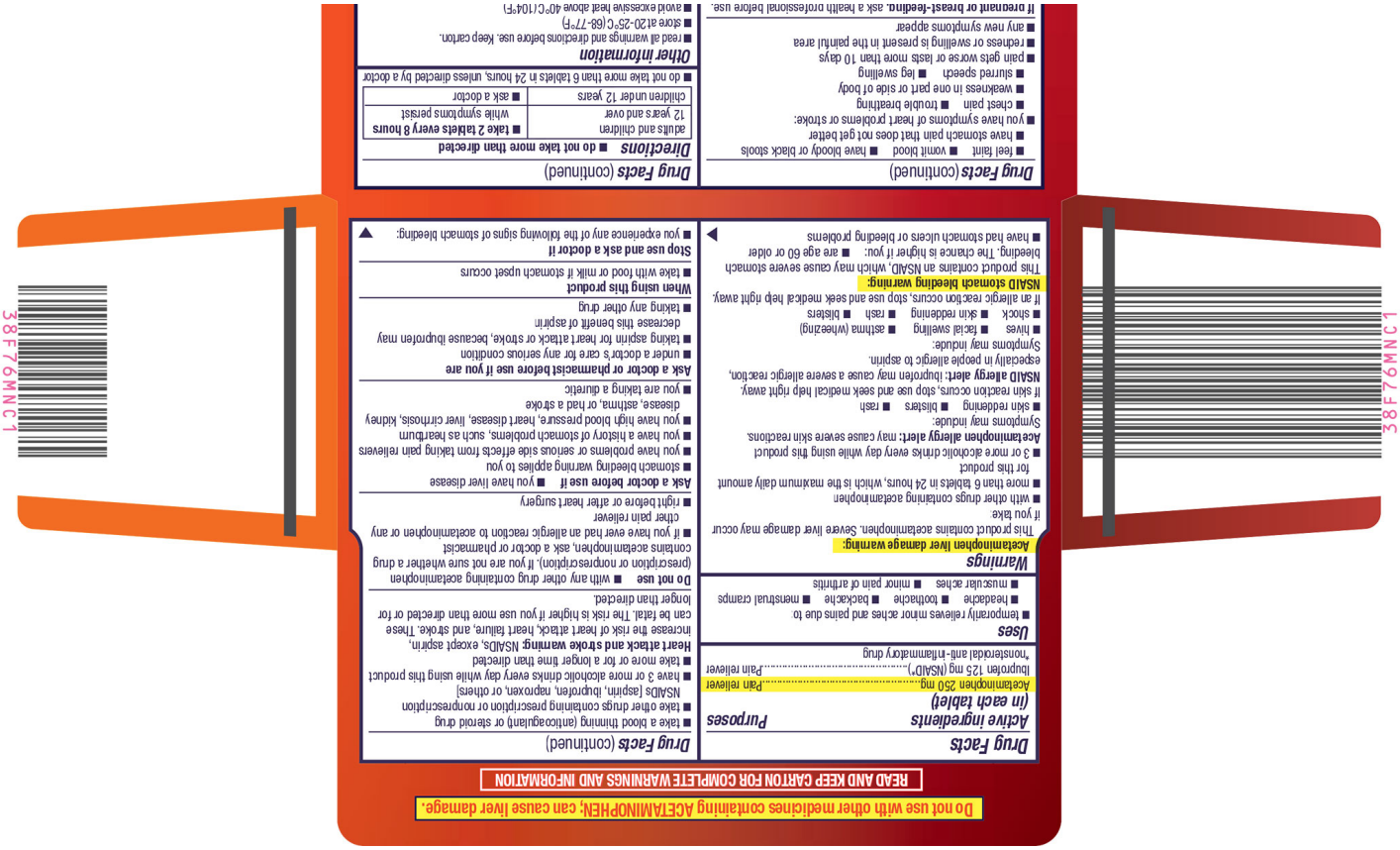
DUAL ACTION WITH TYLENOL

Acetaminophen 250 mg and
Ibuprofen (NSAID) 125 mg Tablets
Pain Reliever

actual size

120 Tablets





MOTRIN DUAL ACTION WITH TYLENOL

acetaminophen and ibuprofen tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50580-208
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
IBUPROFEN (UNII: WK2XYI10QM) (IBUPROFEN - UNII:WK2XYI10QM)	IBUPROFEN	125 mg
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	250 mg

Inactive Ingredients

Ingredient Name	Strength
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
CROSPOVIDONE, UNSPECIFIED (UNII: 2S7830E561)	
TALC (UNII: 7SEV7J4R1U)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	

STARCH, CORN (UNII: O8232NY3SJ)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

Product Characteristics

Color	white	**Score**	no score
Shape	OVAL	**Size**	14mm
Flavor		**Imprint Code**	M;T
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50580-208-02	1 in 1 CARTON	03/20/2023	
1		20 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:50580-208-08	1 in 1 CARTON	03/20/2023	
2		80 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:50580-208-12	1 in 1 CARTON	03/20/2023	
3		120 in 1 BOTTLE; Type 0: Not a Combination Product		
4	NDC:50580-208-16	2 in 1 CARTON	04/01/2024	
4		1 in 1 CARTON		
4		80 in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA214836	03/20/2023	

Labeler - Johnson & Johnson Consumer Inc. (878046358)

Revised: 3/2024 Johnson & Johnson Consumer Inc.