IBUPROFEN CA- ibuprofen tablet, coated
TARGET CORPORATION

600R -TARGET IBUPROFEN 200 MG CAPLETS NDC 11673-990

ACTIVE INGREDIENT(S)

Ibuprofen 200 mg (NSAID)*

* nonstreoidal anti-inflammatory drug

SILICON DIOXIDE

CROSCARMELLOSE SODIUM

FERRIC OXIDE RED

MAGNESIUM STEARATE

CELLULOSE, MICROCRYSTALLINE

TALC

TITANIUM DIOXIDE

STARCH, PREGELATINIZED CORN

POLYVINYL ALCOHOL

POLYETHYLENE GLYCOL

PURPOSE

Pain reliever / fever reducer

USE(S)

temporarily relieves minor aches and pains due to :
backache
headache
menstrual cramps
minor pain of arthritis
muscular aches
the common cold
toothache
temporarily reduces fever

DIRECTIONS

do not take more than directed
the smallest effective dose should be used

adults and children 12 years and older: take 1 caplet every 4 to 6 hours while symptoms persist

if pain or fever does not respond to 1caplet, 2 caplets may be used
do not exceed 6 caplets in 24 hours, unless directed by a doctor.

Children under 12 years: ask a doctor

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)

WARNINGS

Allergy alert: Ibuprofen may cause a severe allergic reaction, especially in people allergic to aspirin.

Symptoms may include:

asthma (wheezing)
blisters
facial swelling
hives
rash
shock
skin reddening

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 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: NSAID's except aspirin increases the risk of heart attack, heart failure, and stroke. These can be fatal. The risk is higher if you use more than directed or longer than directed.

NDC 11673-990-42

ibuprofen

caplets, 200 mg

pain reliever/fever reducer

(NSAID)

up&up

24 CAPLETS**

(**CAPSULE-SHAPED TABLETS)

TAMPER EVIDENT: DO NOT USE THIS PRODUCT IF THE IMPRINTED FOIL SEAL OVER THE MOUTH OF THE BOTTLE IS CUT, TORN, BROKEN OR MISSING

READ AND KEEP CARTON FOR COMPLETE WARNINGS AND INFORMATION

Uses ■ temporarily relieves minor aches and pains due to: ◆ backache ◆ headache ◆ menstrual cramps ◆ minor pain of arthritis ◆ muscular aches ◆ the common cold ◆ toothache ■ temporarily reduces fever

Warnings Allergy alert: Ibuprofen may cause a severe allergic reaction, especially in people allergic to aspirin. Symptoms may include: ■ asthma (wheezing) ■ blisters ■ facial swelling ■ hives ■ rash ■ shock ■ skin reddening. 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 non prescription 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. 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 (1-800-222-1222).

Directions ■ do not take more than directed ■ the smallest effective dose should be used ■ adults and children 12 years and over: ■ take 1 caplet every 4 to 6 hours while symptoms persist ■ if pain or fever does not respond to 1 caplet, 2 caplets may be used ■ do not exceed 6 caplets in 24 hours, unless directed by a doctor ■ children under 12 years: ■ ask a doctor

Other information ■ read all warnings and directions before use ■ store at 20-25°C (68-77°F)

Questions? Call 1-800-910-6874

094 01 0425 R00 C-001227-01-033

Distributed by Target Corporation, Minneapolis, MN 55403

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500R0719

Lot No.:

Exp Date:

Varnish Omit Area

IBUPROFEN CA

ibuprofen tablet, coated

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11673-990
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**
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
STARCH, CORN (UNII: O8232NY3SJ)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics			
Color	brown	**Score**	no score
Shape	CAPSULE ((CAPLET))	**Size**	15mm
Flavor		**Imprint Code**	117
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:11673-990-42	24 in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2020	
2	NDC:11673-990-20	200 in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA091239	01/01/2020	

Labeler - TARGET CORPORATION (006961700)

Registrant - TIME CAP LABORATORIES, INC (037005209)

Establishment

Name	Address	ID/FEI	Business Operations
MARKSANS PHARMA LTD		925822975	manufacture(11673-990)

Revised: 5/2020

TARGET CORPORATION