

**ASPIRIN ADULT LOW STRENGTH- aspirin tablet, delayed release**  
**Valu Merchandisers Company (Best Choice)**

*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.*

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**DRUG FACTS**

**Active ingredient (in each tablet)**

**Aspirin 81 mg (NSAID)\***

\*nonsteroidal anti- inflammatory drug

**Purpose**

**Pain reliever**

**Uses**

- temporarily relieves minor aches and pains
- other therapy as recommended by your doctor. **Because of its delayed action, this product will not provide fast relief of headaches, fever, or other symptoms needing immediate relief.**

**Warnings**

**Reye's syndrome:** Children and teenagers who have or are recovering from chicken pox or flu-like symptoms should not use this product. When using this product, if changes in behavior with nausea and vomiting occur, consult a doctor because these symptoms could be an early sign of **Reye's syndrome**, a rare but serious illness.

**Allergy alert:** Aspirin may cause a severe allergic reaction which may include:

- hives
- facial swelling
- asthma(wheezing)
- shock

**Stomach bleeding warning:** This product contains a nonsteroidal anti-inflammatory drug (NSAID), which may cause 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 have ever had an allergic reaction to any other pain reliever/fever reducer

**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 asthma

**Ask a doctor or pharmacist before use if you are**

taking a prescription drug for:

- anticoagulation (thinning of the blood)
- gout
- diabetes
- arthritis

**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
- allergic reaction occurs
- ringing in the ears or a loss of hearing occurs
- pain gets worse or lasts more than 10 days
- fever gets worse or lasts more than 3 days
- any new symptoms appear
- redness or swelling is present in the painful area

**If pregnant or breast-feeding,**

ask a health professional before use. **It is especially important not to use aspirin 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 exceed recommended dosage**
- drink a full glass of water with each dose
- adults and children 12 years of age and over: take 4 to 8 tablets every 4 hours not to exceed 48 tablets in 24 hours, unless directed by a doctor
- children under 12 years of age: consult a doctor

**Other information**

- store at controlled room temperature 15°-30°C (59°-86°F)
- do not use if imprinted safety seal under cap is broken or missing

**Inactive ingredients**

\*acetylated monoglycerides, \*anhydrous lactose, \*carnauba wax, colloidal silicon dioxide, \*corn starch, \*croscarmellose sodium, D&C Yellow #10 Aluminum Lake, FD&C Yellow #6 Aluminum Lake, hypromellose, \*hypromellose phthalate, \*iron oxide Yellow (iron oxide ochre), methacrylic acid copolymer, microcrystalline cellulose, \*mineral oil, \*polyethylene

glycol (PEG)-400, \*polysorbate 80, povidone, pregelatinized starch, \*propylene glycol, \*simethicone, silicon dioxide, sodium bicarbonate, sodium hydroxide, sodium lauryl sulfate, starch, stearic acid, talc, titanium dioxide, triacetin, and triethyl citrate. \*May also contain.

**Questions or comments?**

**Call toll free 1-877-753-3935**

**Principal Display Panel**

Compare to the active ingredient in ASPIRIN REGIMEN BAYER® 81 mg\*\*

**SEE NEW WARNINGS INFORMATION**

Low Dose 81 mg

ASPIRIN

adult low strength

Pain reliever (NSAID)\*

**KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION**

**\*\*This product is not manufactured or distributed by Bayer Corporation Consumer care division, owner of the registered trademark Aspirin Regimen Bayer® 81 mg**

**Product Label**

4526Q1C

V51e

**Drug Facts** (continued)

Older ■ have had stomach ulcers or bleeding problems ■ take a blood thinning (anticoagulant) or steroid drug ■ take other drugs containing prescription or nonprescription (NSAID) aspirin, ibuprofen, naproxen, or others) ■ have 3 or more alcoholic drinks every day

**Do not use** ■ if you have ever had an allergic reaction to any other pain reliever/fever reducer while using this product ■ take more or for a longer time than directed

**Ask a doctor before use if** ■ you have a history of stomach problems, such as heartburn ■ you are taking a diuretic ■ you have asthma, cirrhosis, or kidney disease ■ you have high blood pressure, heart disease, liver

**Ask a doctor or pharmacist before use if you are taking a prescription drug for:** ■ anticoagulation (thinning of the blood) ■ gout ■ diabetes ■ arthritis

**Stop use and ask a doctor if** ■ you experience any of the following signs of stomach bleeding: ■ vomit blood ■ have bloody or black stools ■ have stomach pain that does not get better or lasts more than 3 days ■ ringing in the ears or a loss of hearing occurs ■ pain gets worse or lasts more than 3 days ■ redness or swelling is present in the painful area

symptoms appear

PROUDLY DISTRIBUTED BY:  
VALU MERCHANDISERS, CO.  
KANSAAS CITY, MO 64111  
www.aspirin.com

**BEST CHOICE**

PLD-F  
F44BC120 0 ■ 700.58 ■ 59137 ■ 5

Lot No.:

Exp. Date:

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION

**Drug Facts** (continued)

**Active Ingredient (in each tablet)**  
Aspirin 81 mg (NSAID)\*  
\*nonsteroidal anti-inflammatory drug  
Pain reliever

**Uses** ■ temporarily relieves minor aches and pains ■ other therapy as recommended by your doctor. ■ Because of its delayed action, this product will not provide fast relief of headaches, fever, or other symptoms needing immediate relief.

**Warnings**  
Hayes's syndrome: Children and teenagers who have or are recovering from chicken pox or flu-like symptoms should not use this product. When using this product, if changes in behavior with nausea and vomiting occur, consult a doctor because these symptoms could be an early sign of Hayes's syndrome, a rare but serious illness.

Allergic aspirin may cause a severe allergic reaction which may include: ■ hives ■ facial swelling ■ asthma (wheezing) ■ shock

**Stomach bleeding warning:** This product contains a nonsteroidal anti-inflammatory drug (NSAID), which may cause stomach bleeding. The chance is higher if you: ■ are age 60 or older ■ take other drugs containing prescription or nonprescription (NSAID) aspirin, ibuprofen, naproxen, or others) ■ have 3 or more alcoholic drinks every day

**Drug Facts** (continued)

methacrylic acid copolymer, microcrystalline cellulose, \*mineral oil, \*polyethylene glycol (PEG)-400, \*polysorbate 80, povidone, pregelatinized starch, \*propylene glycol, \*simethicone, silicon dioxide, sodium bicarbonate, sodium hydroxide, sodium lauryl sulfate, starch, stearic acid, talc, titanium dioxide, triacetin, and triethyl citrate. \*May also contain.

**Questions or comments?** Call toll free 1-877-753-3935

\*This product is not manufactured or distributed by Bayer Corporation Consumer Care Division, owner of the registered trademark Aspirin Regimen Bayer® 81 mg



**Drug Facts** (continued)

If pregnant or breast-feeding, ask a health professional before use. It is especially important not to use aspirin 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 exceed recommended dosage ■ drink a full glass of water with each dose ■ adults and children 12 years of age and over: take 4 to 8 tablets every 4 hours not to exceed 48 tablets in 24 hours, unless directed by a doctor ■ children under 12 years of age: consult a doctor

**Other information** ■ store at controlled room temperature 15°-30°C (59°-86°F) ■ do not use if imprinted safety seal under cap is broken or missing

**Inactive ingredients** aluminum lake, hypromellose, hypromellose phthalate, iron oxide yellow (iron oxide ochre), silicon dioxide, corn starch, croscarmellose sodium, D&C Yellow #10 Aluminum Lake, FD&C Yellow #6

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION

01/27/2010

Enteric coated Aspirin 81 mg

<b>ASPIRIN ADULT LOW STRENGTH</b>			
aspirin tablet, delayed release			
<b>Product Information</b>			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63941-440
Route of Administration	ORAL		
<b>Active Ingredient/Active Moiety</b>			
	<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
	ASPIRIN (UNII: R16CO5Y76E) (ASPIRIN - UNII:R16CO5Y76E)	ASPIRIN	81 mg

## Inactive Ingredients

Ingredient Name	Strength
DIACETYLATED MONO GLYCERIDES (UNII: 5Z17386USF)	
ANHYDROUS LACTOSE (UNII: 3SY5LH9PMK)	
CARNAUBA WAX (UNII: R12CBM0EIZ)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
STARCH, CORN (UNII: O8232NY3SJ)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
HYPROMELLOSE PHTHALATE (24 % PHTHALATE, 55 CST) (UNII: 87Y6436BKR)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
METHACRYLIC ACID - METHYL METHACRYLATE COPOLYMER (1:1) (UNII: 74G4R6TH13)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
MINERAL OIL (UNII: T5L8T28FGP)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
POVIDONES (UNII: FZ989GH94E)	
STARCH, CORN (UNII: O8232NY3SJ)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
STARCH, CORN (UNII: O8232NY3SJ)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
TRIACETIN (UNII: XHX3C3X673)	
TRIETHYL CITRATE (UNII: 8Z96QXD6UM)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
ALUMINUM OXIDE (UNII: LM26O6933)	

## Product Characteristics

Color	YELLOW	Score	no score
Shape	ROUND	Size	10mm
Flavor		Imprint Code	E;HEART;81
Contains			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63941-440-12	1 in 1 CARTON		
1		120 in 1 BOTTLE		

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH FINAL	part343	07/26/2010	

**Labeler** - Valu Merchandisers Company (Best Choice) (868703513)

**Registrant** - P and L Development of New York Corporation (800014821)

Revised: 11/2012

Valu Merchandisers Company (Best Choice)