REGULAR STRENGTH PAIN RELIEF- acetaminophen capsule, gelatin coated SPIRIT PHARMACEUTICALS LLC

VALUMEDS™ REGULAR STRENGTH PAIN RELIEF

Drug Facts

Active ingredient (in each softgel)

Acetaminophen 325 mg

Purpose

Pain reliever/fever reducer

Uses

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

Warnings

Liver warning

This product contains acetaminophen. The maximum daily dose of this product is 10 softgels (3,250 mg) in 24 hours for adults or 5 softgels (1,625 mg) in 24 hours for children. Severe liver damage may occur if

- adult takes more than 4,000 mg of acetaminophen in 24 hours
- child takes more than 5 doses in 24 hours, which is the maximum daily amount
- taken with other drugs containing acetaminophen
- adult has 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.

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 are allergic to acetaminophen or any of the inactive ingredients in this product
- do not give this product to children for the pain of arthritis unless directed by a doctor

Ask a doctor before use if the user has liver disease

Ask a doctor or pharmacist before use if the user is taking the blood thinning drug warfarin

Stop use and ask a doctor if

- pain gets worse or lasts more than 10 days in adults
- pain gets worse or lasts more than 5 days in children under 12 years
- fever gets worse or lasts more than 3 days
- new symptoms occur
- redness or swelling is present

These could be signs of a serious condition.

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

Keep out of reach of children.

Overdose warning

In case of overdose, get medical help or contact a Poison Control Center right away. (1-800-222-1222) Quick 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 (see overdose warning)

adults and children 12 years and over	 take 2 softgels every 4 to 6 hours while symptoms last do not take more than 10 softgels in 24 hours do not use for more than 10 days unless directed by a doctor 		
children 6-11 years	 take 1 softgel every 4 to 6 hours while symptoms last do not take more than 5 softgels in 24 hours do not use for more than 5 days unless directed by a doctor 		
children under 6 years ask a doctor			

Other information

store between 20-25°C (68-77°F)

Inactive ingredients

FD&C red # 40, gelatin, glycerin, polyethylene glycol 400, povidone, propylene glycol, purified water, sorbitol sorbitan solution, titanium dioxide

Questions or comments?

Call **1-888-333-9792**

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RONKONKOMA, NY 11779

PRINCIPAL DISPLAY PANEL - 10 Softgel Bottle Carton

VALUMEDS ™

COMPARE to the active ingredient in REGULAR STRENGTH TYLENOL®*

REGULAR STRENGTH

PAIN RELIEF

Acetaminophen 325 mg

PAIN RELIEVER/FEVER REDUCER

10 softgels

Actual Size



REGULAR STRENGTH PAIN RELIEF

acetaminophen capsule, gelatin coated

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68210-0120
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength

Inactive Ingredients		
Ingredient Name	Strength	
FD&C RED NO. 40 (UNII: WZB9127XOA)		
GELATIN, UNSPECIFIED (UNII: 2G86QN327L)		
GLYCERIN (UNII: PDC6A3C0OX)		
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)		
POVIDONE K30 (UNII: U725QWY32X)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
WATER (UNII: 059QF0KO0R)		
SORBITOL (UNII: 506T60A25R)		
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)		

Product Characteristics			
Color	red	Score	no score
Shape	CAPSULE (OBLONG)	Size	21mm
Flavor		Imprint Code	800
Contains			

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68210- 0120-1	1 in 1 CARTON	04/10/2018	
1		10 in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M013	04/10/2018	
3 1 3			

Labeler - SPIRIT PHARMACEUTICALS LLC (179621011)

Revised: 12/2023 SPIRIT PHARMACEUTICALS LLC