

BHI ARTHRITIS- arnica montana root, berberis vulgaris root bark, bryonia alba whole, causticum, citrullus colocynthis fruit pulp, solanum dulcamara top, ferrosulfuric phosphate, ledum palustre twig, lycopodium clavatum spore, ranunculus bulbosus, rhododendron aureum leaf, toxicodendron pubescens leaf , and sulfur tablet
MediNatura Inc

Disclaimer: This homeopathic product has not been evaluated by the Food and Drug Administration for safety or efficacy. FDA is not aware of scientific evidence to support homeopathy as effective.

BHI Arthritis Tablet

ACTIVE INGREDIENTS

Each tablet contains: *Arnica montana, radix 6X, *Berberis vulgaris 8X, *Bryonia alba 4X, 8X, 12X, 30X, Causticum 8X, *Colocynthis 5X, *Dulcamara 6X, Ferrum phosphoricum 12X, *Ledum palustre 6X, *Lycopodium clavatum 10X, *Ranunculus bulbosus 6X, *Rhododendron chrysanthum 8X, *Rhus toxicodendron 4X, 8X, 12X, 30X, *Sulphur 10X 15.8 mg each.

***Natural Ingredients**

INACTIVE INGREDIENTS

Inactive Ingredients: **Dextrose, ** Lactose, Magnesium Stearate, **Maltodextrin
**contains one or more of these ingredients.

PURPOSE

Arthritis Pain Relief Tablets

Relieves:

- **Arthritis Pain**
- **Joint Stiffness**

WARNINGS

If pregnant or breast-feeding, ask a health professional before use. **Keep out of reach of children.** If symptoms persist or worsen, a health professional should be consulted.**Do not** use if known sensitivity to Arthritis or any of its ingredients exists.

USES

For the temporary relief of minor arthritis pain, joint stiffness.

Directions

At first sign of symptoms: Adults: 1 tablet every 1/2 to 1 hour until symptoms lessen, then continue with standard dosage.

Standard dosage: Adults: Take 1-2 tablets every 4 to 6 hours. Do not exceed 12 tablets in 24 hours.

For children under 18, consult your health professional.

Allow tablets to dissolve completely in the mouth, do not swallow.

KEEP OUT OF REACH OF CHILDREN

Keep out of reach of children.



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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:62795-1010
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ARNICA MONTANA ROOT (UNII: MUE8Y11327) (ARNICA MONTANA ROOT - UNII:MUE8Y11327)	ARNICA MONTANA ROOT	6 [hp_X]
BERBERIS VULGARIS ROOT BARK (UNII: 1TH8Q20J0U) (BERBERIS VULGARIS ROOT BARK - UNII:1TH8Q20J0U)	BERBERIS VULGARIS ROOT BARK	8 [hp_X]
BRYONIA ALBA WHOLE (UNII: 56K0VVT47P) (BRYONIA ALBA WHOLE - UNII:56K0VVT47P)	BRYONIA ALBA WHOLE	4 [hp_X]
CAUSTICUM (UNII: DD5FO1WKFU) (CAUSTICUM - UNII:DD5FO1WKFU)	CAUSTICUM	4 [hp_X]
CITRULLUS COLOCYNTHIS FRUIT PULP (UNII: 23H32AOH17) (CITRULLUS COLOCYNTHIS FRUIT PULP - UNII:23H32AOH17)	CITRULLUS COLOCYNTHIS FRUIT PULP	5 [hp_X]
SOLANUM DULCAMARA TOP (UNII: KPS1B1162N) (SOLANUM DULCAMARA TOP - UNII:KPS1B1162N)	SOLANUM DULCAMARA TOP	6 [hp_X]
FERROSFERRIC PHOSPHATE (UNII: 91GQH8I5F7) (FERROSFERRIC PHOSPHATE - UNII:91GQH8I5F7)	FERROSFERRIC PHOSPHATE	12 [hp_X]
LEDUM PALUSTRE TWIG (UNII: 877L01IZ0P) (LEDUM PALUSTRE TWIG - UNII:877L01IZ0P)	LEDUM PALUSTRE TWIG	6 [hp_X]
LYCOPODIUM CLAVATUM SPORE (UNII: C88X29Y479) (LYCOPODIUM CLAVATUM SPORE - UNII:C88X29Y479)	LYCOPODIUM CLAVATUM SPORE	10 [hp_X]
RANUNCULUS BULBOSUS (UNII: AEQ8NXJ0MB) (RANUNCULUS BULBOSUS - UNII:AEQ8NXJ0MB)	RANUNCULUS BULBOSUS	8 [hp_X]
RHODODENDRON AUREUM LEAF (UNII: IV92NQJ73U) (RHODODENDRON AUREUM LEAF - UNII:IV92NQJ73U)	RHODODENDRON AUREUM LEAF	6 [hp_X]
TOXICODENDRON PUBESCENS LEAF (UNII: 6IO182RP7A) (TOXICODENDRON PUBESCENS LEAF - UNII:6IO182RP7A)	TOXICODENDRON PUBESCENS LEAF	4 [hp_X]
SULFUR (UNII: 70FD1KFU70) (SULFUR - UNII:70FD1KFU70)	SULFUR	10 [hp_X]

Inactive Ingredients

Ingredient Name	Strength
LACTOSE (UNII: J2B2A4N98G)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
DEXTROSE (UNII: IY9XDZ35W2)	

Product Characteristics

Color	white	Score	no score
Shape	ROUND	Size	9mm
Flavor		Imprint Code	Leafman
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62795-1010-3	1 in 1 CARTON	01/01/2015	
1		100 in 1 BOTTLE; Type 0: Not a Combination Product		
1	NDC:62795-	100 in 1 BOTTLE; Type 0: Not a Combination	05/22/2022	

1010-2

Product

03/23/2023

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		01/01/2015	

Labeler - MediNatura Inc (079324099)

Establishment

Name	Address	ID/FEI	Business Operations
MediNatura Inc		102783016	manufacture(62795-1010)

Revised: 12/2024

MediNatura Inc