

**T RELIEF ARTHRITIS- arnica montana, bryonia alba root, sus scrofa cartilage, solanum dulcamara top, sus scrofa embryo,sus scrofa umbilical cord, ledum palustre twig,sus scrofa umbilical cord, ledum palustre twig,sus scrofa placenta, rhododendron aureum leaf,toxicodendron pubescens leaf, sanguinaria canadensis root, sulfur, and comfrey root 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.

T Relief Arthritis Tablet

KEEP OUT OF REACH OF CHILDREN

Keep out of reach of children

WARNINGS

Warnings: If pregnant or breastfeeding, ask health professional. **Keep out of reach of children.** If symptoms persist or worsen, a health professional should be consulted. **Do not use** if known sensitivity to T-Relief™ Arthritis or any of its ingredients exists.

ACTIVE INGREDIENTS

Each tablet contains: *Dulcamara
2X 5mg, *Arnica montana 3X 10mg, *Sanguinaria
canadensis 3X 0.5mg,*Bryonia alba 3X 0.3mg, *Cartilago
suis 4X 0.3mg,*Rhus toxicodendron 4X 5mg, *Sulphur
6X 5mg,*Ledum palustre 6X 4mg,*Bryonia alba 6X
0.3mg, *Embryo suis 6X 0.3mg,*Funiculus umbilicalis
suis 6X 0.3mg, *Placenta totalis suis 6X 0.3mg,
*Symphytum officinale 7X 0.02mg, *Rhododendron
chrysanthum 8X 0.3mg, *Bryonia alba 12X 0.3mg.
*Natural Ingredient

Uses

Relieves

- Arthritis pain
- Joint pain
- Joint stiffness

INACTIVE INGREDIENTS

Inactive Ingredients: Dextrose, Maltodextrin Magnesium stearate.

PURPOSE

Uses: For the temporary relief of minor: Arthritis pain, Joint pain, Joint stiffness

Directions

For max absorption, dissolve under tongue. Can be chewed and swallowed. Adults 2 tablets every 4 hours. Do not exceed 12 tablets per 24 hours. Children under 18 years, consult your health professional.



Add image transcription here...

T RELIEF ARTHRITIS

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twig,sus scrofa placenta, rhododendron aureum leaf,toxicodendron pubescens leaf, sanguinaria canadensis root, sulfur, and comfrey root tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:62795-1033
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ARNICA MONTANA (UNII: O80TY208ZW) (ARNICA MONTANA - UNII:O80TY208ZW)	ARNICA MONTANA	3 [hp_X]
BRYONIA ALBA ROOT (UNII: T7J046YI2B) (BRYONIA ALBA ROOT - UNII:T7J046YI2B)	BRYONIA ALBA ROOT	3 [hp_X]
SUS SCROFA CARTILAGE (UNII: 73ECW5WG2F) (SUS SCROFA CARTILAGE - UNII:73ECW5WG2F)	SUS SCROFA CARTILAGE	4 [hp_X]
SOLANUM DULCAMARA TOP (UNII: KPS1B1162N) (SOLANUM DULCAMARA TOP - UNII:KPS1B1162N)	SOLANUM DULCAMARA TOP	2 [hp_X]
SUS SCROFA EMBRYO (UNII: 9928MC12VO) (SUS SCROFA EMBRYO - UNII:9928MC12VO)	SUS SCROFA EMBRYO	6 [hp_X]
SUS SCROFA UMBILICAL CORD (UNII: 118OYG6W3H) (SUS SCROFA UMBILICAL CORD - UNII:118OYG6W3H)	SUS SCROFA UMBILICAL CORD	6 [hp_X]
LEDUM PALUSTRE TWIG (UNII: 877L01IZ0P) (LEDUM PALUSTRE TWG - UNII:877L01IZ0P)	LEDUM PALUSTRE TWG	6 [hp_X]
SUS SCROFA PLACENTA (UNII: C8CV8867O8) (SUS SCROFA PLACENTA - UNII:C8CV8867O8)	SUS SCROFA PLACENTA	6 [hp_X]
RHODODENDRON AUREUM LEAF (UNII: IV92NQJ73U) (RHODODENDRON AUREUM LEAF - UNII:IV92NQJ73U)	RHODODENDRON AUREUM LEAF	8 [hp_X]
TOXICODENDRON PUBESCENS LEAF (UNII: 6IO182RP7A) (TOXICODENDRON PUBESCENS LEAF - UNII:6IO182RP7A)	TOXICODENDRON PUBESCENS LEAF	4 [hp_X]
SANGUINARIA CANADENSIS ROOT (UNII: N9288CD508) (SANGUINARIA CANADENSIS ROOT - UNII:N9288CD508)	SANGUINARIA CANADENSIS ROOT	3 [hp_X]
SULFUR (UNII: 70FD1KFU70) (SULFUR - UNII:70FD1KFU70)	SULFUR	6 [hp_X]
COMFREY ROOT (UNII: M9VVZ08EKQ) (COMFREY ROOT - UNII:M9VVZ08EKQ)	COMFREY ROOT	7 [hp_X]

Inactive Ingredients

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

Product Characteristics

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

Packaging

		Marketing Start	Marketing End
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#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62795-1033-2	1 in 1 CARTON	03/01/2016	
1		100 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:62795-1033-4	100 in 1 BOTTLE; Type 0: Not a Combination Product	11/16/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		03/01/2016	

Labeler - MediNatura Inc (079324099)

Registrant - MediNatura Inc (079324099)

Establishment

Name	Address	ID/FEI	Business Operations
MediNatura Inc		102783016	manufacture(62795-1033)

Revised: 8/2023

MediNatura Inc