

**ARNICA GEL 15 ARNICA MONTANA 1X HPUS- arnica gel gel
VITAMEDICA, 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.

Arnica Gel + 15

ACTIVE INGREDIENT:

Arnica Montana 200C HPUS.**

**The letters 'HPUS' indicate that the component in this product is officially monographed in the Homeopathic Pharmacopoeia of the United States. This product has been manufactured at an FDA-regulated facility.

PURPOSE:

Arnica Montana - Disoloration from bruising; relieves join and muscle pain, soreness & stiffness; relieves swelling from *

*Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated.

USES:

*Temporarily relieves these symptoms:

- Bruising • Swelling • Pain • Stiff • Muscle Soreness

*Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated.

WARNINGS:

For extrenal use only. Avoid contract with eyes and mucous membranes.

Do not use on open wounds or broken skin

Stop use or consult a physician if

- A rash or irritation occurs
- Symptoms persist for more than 7 days or worsen.

Keep this and all medications out of reach of children.

If sawllowed, seek medical attention or contact a Poison control center immediately.

KEEP OUT OF REACH OF CHILDREN:

If swallowed, seek medical attention or contact a Poison Control Center immediately.

DIRECTIONS:

Apply a thin layer of gel to target area. Reapply 3 times daily or as needed.

Other Information

Other Information

- Do not use if tamper evident seal is broken.
- Store at 68-77°F (20-25°C)

INACTIVE INGREDIENTS:

Water, Aloe Vera Leaf Juice Extract, Benzyl Alcohol, Ethylhexylglycerin, Tocopherol, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Disodium EDTA, Sodium Hydroxide, Ethanol

QUESTIONS:

Formulated and Distributed by

VitaMedica, Inc.

Tampa, FL 33624

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www.vitamedica.com

888-367-8605

customerservice@vitamedica.com

Label

NDC 80004-400-40

Pain Relief*

VitaMedica

Arnica Gel + 15

Extra Strength Pain Relief

1/4" NO PRINT AREA

Quiet zone. All copy must be 1/16" below the eye-mark.

PAIN RELIEF*

VitaMedica

Arnica

Center
Front Panel

Extra Strength Pain Relief

Clinical Strength

- Bruising & swelling*
- Muscle soreness & pain*
- Joint pain & stiffness*

All-Natural Pain Relief

With Soothing Aloe & Arnica
+15 Homeopathics

Homeopathic Medicine
NET WT 3.0 oz (85 g)

Center
Art Work

2-25/64"

4-25/32" Actual Tube Circumference

Drug Facts

Active Ingredients** Arnica Montana 1X HPUS, Calendula Officinalis 1X HPUS, Hypericum Perforatum 1X HPUS, Symphytum Officinale 6X HPUS, Bellis Perennis 12X HPUS, Ledum Palustre 12X HPUS, Sulphuricum Acidum 12X HPUS, Calcarea Fluorica 30X HPUS, Arnica Montana 30C HPUS, Ferrum Phosphoricum 30X HPUS, Rhus Tox 30X HPUS, Hypericum Perigratum 30C HPUS, Ruta Graveolens 30X HPUS, Natrum Sulphuricum 30C HPUS, Calcarea Carbonica 30C HPUS, Millefolium 12X HPUS, Hamamelis 1X HPUS, Belladonna 30X HPUS

Warnings*
• discoloration from bruising; relieves joint and pain, soreness & stiffness; relieves swelling from

• Swelling
• Stiffness
trauma due to: • Min

Warnings
For external use only. Avoid contact with eyes and mucous membranes.

- Do not use on open wounds or broken skin.
- Stop use and ask a doctor if**
- A rash or irritation occurs
 - Symptoms persist for more than 7 days or worsen

KEEP OUT OF REACH OF CHILDREN. If swallowed, seek medical attention or contact a Poison Control Center immediately.

Directions
Apply a thin layer of gel to target area. Reapply 3 times daily or as needed.

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Inactive Ingredients
Water, Aloe Vera Leaf Juice Extract, Benzyl Alcohol, Ethylhexylglycerin, Tocopherol, Acrylates/C10-30 Alkyl Acrylate, EDTA, Sodium Hydroxide, Ethanol.

Questions?
888-367-8605 | www.vitamedita.com | info@vitamedita.com

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NDC-80004-005-54
Formulated & Distributed by VitaMedica®, Henderson, NV 89014



ARNICA GEL 15 ARNICA MONTANA 1X HPUS

arnica gel gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:80004-400
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HYPERICUM PERFORATUM (UNII: XK4IUX8MNB) (HYPERICUM PERFORATUM - UNII:XK4IUX8MNB)	HYPERICUM PERFORATUM	1 [hp_X] in 1 g
SYMPHYTUM OFFICINALE WHOLE (UNII: H8FJJ6KX5Y) (SYMPHYTUM OFFICINALE WHOLE - UNII:H8FJJ6KX5Y)	SYMPHYTUM OFFICINALE WHOLE	6 [hp_X] in 1 g
ARNICA MONTANA WHOLE (UNII: O80TY208ZW) (ARNICA MONTANA WHOLE - UNII:O80TY208ZW)	ARNICA MONTANA WHOLE	30 [hp_C] in 1 g
RHUS TYPHINA WHOLE (UNII: Y465D8AABT) (RHUS TYPHINA WHOLE - UNII:Y465D8AABT)	RHUS TYPHINA WHOLE	30 [hp_X] in 1 g

HAMAMELIS VIRGINIANA BARK (UNII: IH3063S9MY) (HAMAMELIS VIRGINIANA BARK - UNII:IH3063S9MY)	HAMAMELIS VIRGINIANA BARK	1 [hp_X] in 1 g
CALENDULA OFFICINALIS FLOWERING TOP (UNII: 18E7415PXQ) (CALENDULA OFFICINALIS FLOWERING TOP - UNII:18E7415PXQ)	CALENDULA OFFICINALIS FLOWERING TOP	1 [hp_X] in 1 g
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ) (ARNICA MONTANA FLOWER - UNII:OZ0E5Y15PZ)	ARNICA MONTANA FLOWER	1 [hp_X] in 1 g
LEDUM PALUSTRE WHOLE (UNII: 1N8KG72C5M) (LEDUM PALUSTRE WHOLE - UNII:1N8KG72C5M)	LEDUM PALUSTRE WHOLE	12 [hp_X] in 1 g
ACHILLEA MILLEFOLIUM FLOWER (UNII: YQR8R0SQEA) (ACHILLEA MILLEFOLIUM FLOWER - UNII:YQR8R0SQEA)	ACHILLEA MILLEFOLIUM FLOWER	12 [hp_X] in 1 g
BELLADONNA LEAF (UNII: 6GZW20TIOI) (BELLADONNA LEAF - UNII:6GZW20TIOI)	BELLADONNA LEAF	30 [hp_X] in 1 g
BELLIS PERENNIS FLOWER (UNII: 26I94X9A1K) (BELLIS PERENNIS FLOWER - UNII:26I94X9A1K)	BELLIS PERENNIS FLOWER	12 [hp_X] in 1 g
RUTA GRAVEOLENS WHOLE (UNII: 181JI0338P) (RUTA GRAVEOLENS WHOLE - UNII:181JI0338P)	RUTA GRAVEOLENS WHOLE	30 [hp_X] in 1 g

Inactive Ingredients

Ingredient Name	Strength
CALCIUM DISODIUM EDTA (UNII: 8U5D034955)	
2-ETHYLHEXYL GLYCIDYL ETHER (UNII: LU1UZ98B89)	
ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER (60000 MPA.S) (UNII: 8Z5ZAL5H3V)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
2-(ETHYLTHIO)ETHANOL (UNII: 6091RQ1378)	
.ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)	
WATER (UNII: 059QF0KO0R)	
ALOE VERA LEAF JUICE (UNII: RUE8E6T4NB)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	

Product Characteristics

Color	white (Gel)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80004-400-40	1 in 1 CARTON	03/01/2024	
1		85 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		03/01/2024	

Labeler - VITAMEDICA, INC. (119401422)

Revised: 3/2024

VITAMEDICA, INC.