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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888-367-8605

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Label

NDC 80004-400-40

Pain Relief*

VitaMedica

Arnica Gel + 15

Extra Strength Pain Relief

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.