ARNICA MONTANA 200C HPUS- arnica montana tablet
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.

DRUG FACTS:

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 - Bruising, swelling, pain*

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

USES:

*Temporarily relieves these symptoms:

• Bruising • Swelling • Pain • Stiffness • Muscle Soreness

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

WARNINGS:

Ask a doctor before use if pregnant or breastfeeding.

Stop use or consult a physician if

• Any adverse reactions occur

• Symptoms persist for more than 14-days or worsen

Keep this and all medications out of reach of children.

In case of an emergency or accidental overdose, contact a medical professional or Poison Control immediately.

CONTAINS: Lactose (milk)

• Do not use if seal is broken or shows any signs of tampering.

• Store at 68-77°F (20-25°C)

KEEP OUT OF REACH OF CHILDREN:

In case of an emergency or accidental overdose, contact a medical professional or Poison Control immediately.

DIRECTIONS:

• Do not handle the tablets. Instead, drop tablets directly under the tongue and let dissolve naturally.

Adults and children 12 years and over - Take 3 tablets, 3 times a day

Children under 12 years - Not intended

INACTIVE INGREDIENTS:

Croscarmellose Sodium, Lactose, Magnesium Stearate, Microcrystalline Cellulose, Sucrose.

QUESTIONS:

Formulated and Distributed by

VitaMedica, Inc.

Tampa, Fl 33624

888-367-8605

www.vitamedica.com

customerservice@vitamedica.com

PACKAGE DISPLAY LABEL

VitaMedica

Arnica Montana

200C HPUS

Maximum Potency

Reduces

•Pain*

•Bruising*

•Swelling*

Natural Remedy Stimulates Healing

150 Tablets

Homeopathic Medicine

Drug Facts

Active Ingredient* — ***Purpose****

Arnica Montana 200C HPUS ..... Bruising, swelling, pain

Uses* Temporarily relieves these symptoms:

• Bruising • Swelling • Pain • Stiffness • Muscle Soreness

Warnings

Ask a doctor before use if pregnant or breastfeeding.

Stop use or consult a physician if

• Any adverse reactions occur

• Symptoms persist for more than 14-days or worsen

Keep this and all medications out of reach of children. In case of an emergency or accidental overdose, contact a medical professional or Poison Control Center immediately.

Drug Facts (continued)

Directions

• Do not handle the tablets. Instead, drop tablets directly under the tongue and let dissolve naturally.

Adults and children 12 years and over	Take 3 tablets, 3 times a day
Children under 12 years	Not intended

Other Information

• Do not use if seal is broken or shows any signs of tampering.

• Store at 68-77°F (20-25°C)

Inactive Ingredients

Croscarmellose Sodium, Lactose, Magnesium Stearate, Microcrystalline Cellulose, Sucrose.

Drug Facts (continued)

Questions or Comments?

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CONTAINS: Lactose (milk)

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ARNICA MONTANA 200C HPUS

arnica montana tablet

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:80004-009
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ARNICA MONTANA WHOLE (UNII: O80TY208ZW) (ARNICA MONTANA - UNII:O80TY208ZW)	ARNICA MONTANA WHOLE	200 [hp_C]

Inactive Ingredients

Ingredient Name	Strength
CROSCARMELLOSE (UNII: 029TFK992N)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
SUCROSE (UNII: C151H8M554)	

Product Characteristics

Color	white	**Score**	no score
Shape	ROUND	**Size**	6mm
Flavor		**Imprint Code**	V
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80004-009-20	150 in 1 BOTTLE; Type 0: Not a Combination Product	01/23/2026	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		01/23/2026	

Labeler - VITAMEDICA, INC. (119401422)

Registrant - Apotheca Company (844330915)

Establishment

Name	Address	ID/FEI	Business Operations

Apotheca Company		844330915	manufacture(80004-009) , api manufacture(80004-009) , label(80004-009) , pack(80004-009)

Revised: 1/2026 VITAMEDICA, INC.