ARNICA MONTANA - arnica montana tablet Schwabe North America

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 Montana 6X

Active Ingredient

Arnica montana 6X

Inactive Ingredient

CELLULOSE

LACTOSE MONOHYDRATE

MAGNESIUM STEARATE

SODIUM CROSCARMELLOSE

Dosage & Administration

Dosage:

Adults or children over 2 years: Take 3 tablets 4-6 times a day.

Allow to dissolve under tongue.

Indications

Temporarily relieves minor muscle or joint aches and pain; strains, sprains, bruises, backache or over-exertion.

Purpose

For the temporary relief of minor muscle or joint aches and pain; strains, sprains, bruises, backache or over-exertion.

Warning

Do not take this product for pain for more than 10 days (adult) or 5 days (children) unless directed by a doctor.

When Using

When using this product if pain persists or gets worse, if new symptoms occur, or if redness or swelling is present, consult a doctor because these could be signs of a serious condition.

Pregnancy and Breast Feeding

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children.

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Overdosage

In case of overdose, seek medical help or contact a Poison Control Center immediately.



ARNICA MONTANA

arnica montana tablet

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:53499-3533

Route of Administration SUBLINGUAL

Active Ingredient/Active Moiety

l	Ingredient Name	Basis of Strength	Strength
ı	ARNICA MONTANA (UNII: O80TY208ZW) (ARNICA MONTANA - UNII:O80TY208ZW)	ARNICA MONTANA	6 [hp_X]

Inactive Ingredients

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Ingredient Name	Strength	
CELLULO SE, MICRO CRYSTALLINE (UNII: OP1R32D61U)		
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)		
MAGNESIUM STEARATE (UNII: 70097M6I30)		
CROSCARMELLOSE SODIUM (UNII: M28 OL1HH48)		

Product Characteristics

Color	white (white tablet)	Score	no score
Shape	ROUND (B&T)	Size	6 mm
Flavor		Imprint Code	B;T
Contains			

Packaging

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#	t Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53499-3533-	100 in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	0 1/0 1/20 0 6	09/30/2019
2	NDC:53499-3533-	1 in 1 CARTON	0 1/0 1/20 0 6	10/31/2025
2	1	250 in 1 BOTTLE, GLASS; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		0 1/0 1/20 0 6	10/31/2025

Labeler - Schwabe North America (831153908)

Establishment					
Name	Address	ID/FEI	Business Operations		
Schwabe Mexico, S.A. de C.V.		812805901	manufacture(53499-3533)		

Revised: 10/2020 Schwabe North America