ARNICA MONTANA- arnica montana tablet
Schwabe North America, 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 Montana 30X

Active Ingredient

Arnica montana 30X

Inactive Ingredients

CELLULOSE

LACTOSE MONOHYDRATE

MAGNESIUM STEARATE

SODIUM CROSCARMELLOSE

Dosage & Admininstration

Adults or children over 2 years: Take 3 tablets 1 to 3 times per day.

Allow to dissolve under tongue.

Indications & Usage

Temporarily relieves minor muscle or joint aches and pain: strains, sprains, bruises, backached, overexertion.

Purpose

Temporarily relieves minor muscle or joint aches and pain: strains, sprains, bruises, backached, overexertion.

Warning

Do not use this product for pain for more than 10 days for adults and 5 days for 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 serious conditions.

Pregnancy or breast feeding

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

Keep out of reach of children

Keep out of reach of children.

Overdose

In case of overdoese, seek medical help or contact a Posion Control Center immediately.

ARNICA MONTANA

arnica montana tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:53499-3572
Route of Administration	SUBLINGUAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ARNICA MONTANA (UNII: O80TY208ZW) (ARNICA MONTANA - UNII:O80TY208ZW)	ARNICA MONTANA	30 [hp_X]

Inactive Ingredients	
Ingredient Name	**Strength**
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	

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

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:53499-3572-2	100 in 1 BOTTLE, GLASS; Type 1: Convenience Kit of Co-Package	01/01/2006	03/31/2019
2	NDC:53499-3572-3	250 in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	01/01/2006	12/30/2025

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		01/01/2006	12/30/2025

Labeler - Schwabe North America, Inc. (831153908)

Establishment

Name	Address	ID/FEI	Business Operations
Schwabe Mexico, S.A. de C.V.		812805901	manufacture(53499-3572)