

ARNICA MONTANA- arnica montana tablet
Hyland's 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

BRUISES OR MUSCLE SORENESS

ARNICA MONTANA 30X

BRUISING & MUSCLE SORENESS

Made according to the Homeopathic Pharmacopoeia of the United States since 1903.

Warnings

Do not use if cap band is missing or broken.

If you are pregnant or nursing, consult a licensed health care professional before using this product.

If symptoms persist for 7 days or worsen, contact a licensed practitioner.

Keep this and all medicines out of the reach of children.

To be used according to label indications and/or standard homeopathic indications.

Directions

Adults: Dissolve 4 tablets under tongue 4 times a day.

Children: consult a healthcare professional

Inactive ingredients

Acacia Gum and Lactose

Questions?

800-624-9659

PRINCIPAL DISPLAY PANEL - 250 Tablet Bottle Label

SINCE 1903

Hyland's®

HOMEOPATHIC

NDC 54973-2904-4

**ARNICA
MONTANA
30X**

**BRUISES OR MUSCLE
SORENESS***

250 TABLETS

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

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Standard Homeopathic Company
Los Angeles, CA 90061
Questions? 800-624-9659

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3 54973 29044 0
LOT #: XXXXXX
REV. 2

ARNICA MONTANA

arnica montana tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:54973-2904
Route of Administration	ORAL		

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		
ACACIA (UNII: 5C5403N26O)				
LACTOSE (UNII: J2B2A4N98G)				
Product Characteristics				
Color	white (White to Off-White)	Score	no score	
Shape	ROUND	Size	5mm	
Flavor		Imprint Code		
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54973-2904-4	250 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/01/1955	
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
unapproved homeopathic			01/01/1955	

Labeler - Hyland's Inc. (008316655)

Establishment

Name	Address	ID/FEI	Business Operations
Hyland's Inc.		008316655	manufacture(54973-2904) , pack(54973-2904)

Revised: 12/2022

Hyland's Inc.