ARNICA MONTANA- arnica montana pellet Rxhomeo Private Limited d.b.a. Rxhomeo, 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.

ACTIVE INGREDIENT

ARNICA MONTANA HPUS 3X and higher

USES

USES: Temporary Relief - Bruises, Aches, Pains*

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

INDICATIONS

Condition listed above or as directed by the physician

DOSAGE

Adults- Take 4 or 6 Pellets by mouth, three times daily or as suggested by physician. Children 2 years and older- take 1/2 the adult dose.

WARNINGS

This product is to be used for self-limiting conditions

If symptoms do not improve in 4 days, or worsen, discontinue use and seek assistance of health professional

As with any drug, if you are preganant, or nursing a baby, seek professional advice before taking this product

Keep this and all medication out of reach of children

Do not use if capseal is broken or missing.

Close the cap tightly after use.

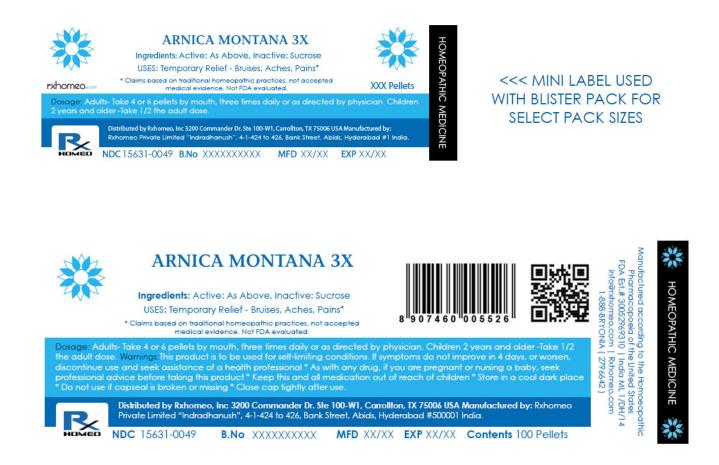
INACTIVE INGREDIENTS

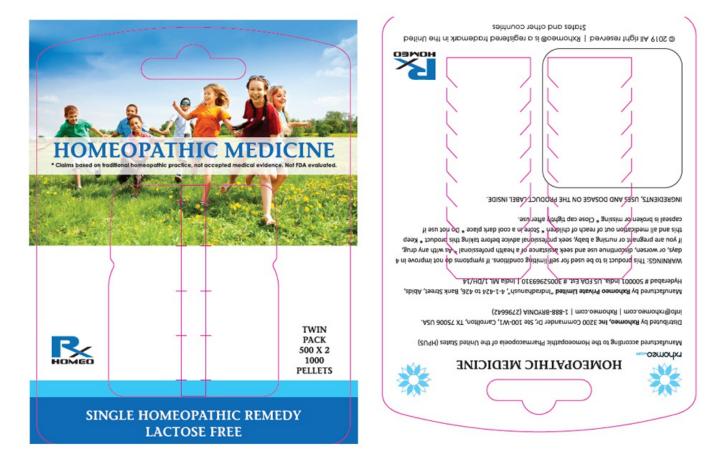
Sucrose

STORAGE

QUESTIONS OR COMMENTS

www.Rxhomeo.com | 1.888.2796642 | info@rxhomeo.com Rxhomeo, Inc 3200 Commander Dr, Ste 100-W1, Carrollton, TX 75006 USA







HOMEOPATHIC MEDICINE

i.e. regular size or mini size. 3. Secondary Packing > If the bottle/container is small, we will use a secondary packing i.e a

Carton box or a Blister pack.

ARNICA MONTANA

arnica montana pellet

Product Information

Product Type		HUMAN OTC DRUG	Item Code (Source)		NDC:15631-0049			
Route of Administration			ORAL					
A	ctive Ingred	ient/Active	Moiety					
Ingredient Name Basis of Strength Stren								
AF	RNICA MONTANA	A (UNII: 080TY20)8ZW) (ARNICA MONTANA -	UNII:080TY208ZW)	ARNICA MON	ITANA	3 [hp_X]	
In	active Ingre							
		Ing	gredient Name		Strength			
5ι	JCROSE (UNII: C	151H8M554)						
Pa	ackaging							
#	ltem Code	Pa	ckage Description		ting Start ate		ting End ate	
1	NDC:15631- 0049-0	100 in 1 PACKA Product	GE; Type 0: Not a Combin	ation 01/01/2018	01/01/2018			
2	NDC:15631- 0049-1	200 in 1 PACKA Product	GE; Type 0: Not a Combin	ation 01/01/2018	01/01/2018			
3	NDC:15631- 0049-2	400 in 1 PACKA Product	GE; Type 0: Not a Combin	ation 01/01/2018	3			
4	NDC:15631- 0049-3	Product	GE; Type 0: Not a Combin	01/01/2010	3			
5	NDC:15631- 0049-4	Product	AGE; Type 0: Not a Combi	01/01/2010	3			
6	NDC:15631- 0049-5	Product	KAGE; Type 0: Not a Coml	01/01/2010	3			
7	NDC:15631- 0049-6	Product	GE; Type 0: Not a Combin	00/25/202	1			
8	NDC:15631- 0049-7	1000 in 1 PACK Product	AGE; Type 0: Not a Combi	nation 08/23/2023	1			
M	larketing	Informat	ion					
	Marketing Category		tion Number or Mono Citation		ting Start Date		ting End ate	
	approved			09/07/203				

Labeler - Rxhomeo Private Limited d.b.a. Rxhomeo, Inc (650833994)

Establishment										
Address	ID/FEI	Business Operations								
	650833994	manufacture(15631-0049) ,	label(15631-0049)							
	Address	•	Address ID/FEI Business Ope 650833994 manufacture(15631-0049) ,							