

ACALYPHA INDICA- acalypha indica pellet
HOMEOLAB USA 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.

HOMEOPATHIC MEDICINE NDC 60512-6440-1

ACTIVE INGREDIENT HPUS

ACALYPHA INDICA 1X

(Indian nettle)

DRY COUGH

USE

For self-limiting condition listed above or as directed by a health professional.

WARNINGS

Do not use if pellet-dispenser seal is broken.

Stop use and ask a doctor if symptoms persist more than 3 days or worsen.

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

Keep out of reach of children.

DIRECTIONS

Adults: Allow 3 or 4 pellets to dissolve in the mouth 3 times a day until symptoms are relieved or as directed by a health professional.

OTHER INFORMATION

Store at room temperature.

INACTIVE INGREDIENTS

Lactose, sucrose.

QUESTIONS?

1-800-404-4666

The letters 'HPUS' indicate that the component in this product is officially monographed in the Homeopathic Pharmacopoeia of the United States.

These claims have not been reviewed by the Food and Drug Administration. They are based on traditional homeopathic practice.□

80 Pellets

Pellet dispenser

Mfd for: HOMEOLAB USA INC., 3025 De L'Assomption, Montreal, QC, H1N 2H2, CANADA
Product of Canada

LABEL

HOMEOPATHIC MEDICINE

ACALYPHA
INDICA

Indian Nettle

NDC 60512-6440-1

DRY COUGH *

1x

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Rev. 10/13

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3025 De L'Assomption, Montreal QC H1N 2H2 CANADA

1-800-404-4666 / www.homeolab.com

PRODUCT OF CANADA



HOMEOLAB
USA

Break seal, turn & twist.



ACALYPHA INDICA			
acalypha indica pellet			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:60512-6440
Route of Administration	ORAL		

Active Ingredient/Active Moiety				
	Ingredient Name	Basis of Strength	Strength	
	ACALYPHA INDICA (UNII: Y34S6V2PSH) (ACALYPHA INDICA - UNII:Y34S6V2PSH)	ACALYPHA INDICA	1 [hp_X]	

Inactive Ingredients	
Ingredient Name	Strength
LACTOSE (UNII: J2B2A4N98G)	
SUCROSE (UNII: C151H8M554)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:60512-6440-1	80 in 1 TUBE		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		10/11/1995	

Labeler - HOMEOLAB USA INC. (202032533)

Registrant - HOMEOLAB USA INC. (202032533)

Establishment			
Name	Address	ID/FEI	Business Operations
HOMEOLAB USA INC.		202032533	manufacture(60512-6440)