

CONVALLARIA MAJALIS - convallaria majalis liquid
Newton Laboratories, 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.

Convallaria majalis

INDICATIONS & USAGE SECTION

Aching of legs, Irregular pulse, Headache.

DOSAGE & ADMINISTRATION SECTION

Directions: Ages 12 and up, take 6 drops by mouth (ages 0 to 11, give 3 drops) as needed or as directed by a health professional. Sensitive persons begin with 1 drop and gradually increase to full dose.

OTC - ACTIVE INGREDIENT SECTION

Convallaria majalis 15x, 10x, 30c.

OTC - PURPOSE SECTION

Aching of legs, Irregular pulse, Headache.

INACTIVE INGREDIENT SECTION

Inactive Ingredients: USP Purified Water; USP Gluten-free, non-GMO, organic cane alcohol 20%.

QUESTIONS SECTION

www.newtonlabs.net Newton Laboratories, Inc. FDA Est # 1051203 - Conyers, GA 30012
Questions? 1.800.448.7256

WARNINGS SECTION

WARNINGS: Keep out of reach of children. Do not use if tamper-evident seal is broken or missing. If symptoms worsen or persist for more than a few days, consult a doctor. If **pregnant or breast-feeding**, ask a doctor before use.

OTC - PREGNANCY OR BREAST FEEDING SECTION

If **pregnant or breast-feeding**, ask a doctor before use.

OTC - KEEP OUT OF REACH OF CHILDREN SECTION

Keep out of reach of children.

PACKAGE LABEL



NEWTON
homeopathics™ 

30c

Nurturing Naturally Since 1987™

NDC# 55714-6174-1

**Convallaria
majalis**

*Aching of legs; Irregular
pulse; Headache*

1 fl oz (29.57 ml)

To
Direc
Ages 12
as need
personal
Aches
Insects
non-GMO
Contains
Diy at h
Sinc

CONVALLARIA MAJALIS

convallaria majalis liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55714-6174
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Convallaria Majalis (UNII: QHH4HVF5QE) (Convallaria Majalis - UNII:QHH4HVF5QE)	Convallaria Majalis	15 [hp_X] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
Alcohol (UNII: 3K9958V90M)	
Water (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55714-6174-1	30 mL in 1 BOTTLE, GLASS		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		09/01/2011	

Labeler - Newton Laboratories, Inc. (788793610)

Registrant - Newton Laboratories, Inc. (788793610)

Establishment

Name	Address	ID/FEI	Business Operations
Newton Laboratories, Inc.		788793610	MANUFACTURE(55714-6174)