

**LOBELIA COMP.- lobelia comp. liquid**  
**Uriel Pharmacy, 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.*

**Lobelia comp.**

Directions: FOR ORAL USE ONLY.

Take 3-4 times daily. Ages 12 and older: 10 drops.  
Ages 2-11: 5 drops. Under age 2: Consult a doctor.

Active Ingredients: Quercus Dec (Oak) 4X, Veronica Dec (Speedwell) 4X, Lobelia 6X, Plumbum met. (Lead) 12X

Inactive Ingredients: Distilled water, 20% Organic cane alcohol  
Prepared using rhythmical processes

Use: Temporary relief of headache.

KEEP OUT OF REACH OF CHILDREN.

Warnings: Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated. Do not use if allergic to any ingredient. Consult a doctor before use for serious conditions or if conditions worsen or persist. If pregnant or nursing, consult a doctor before use. Do not use if safety seal is broken or missing.

Questions? Call 866.642.2858

Made with care by Uriel, East Troy, WI 53120  
shopuriel.com Lot:



**LOBELIA COMP.**

lobelia comp. liquid

**Product Information**

**Product Type**

HUMAN OTC DRUG

**Item Code (Source)**

NDC:48951-6090

Route of Administration ORAL

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>VERONICA OFFICINALIS FLOWERING TOP</b> (UNII: 9IH82J936J) (VERONICA OFFICINALIS FLOWERING TOP - UNII:9IH82J936J)	VERONICA OFFICINALIS FLOWERING TOP	4 [hp_X] in 1 mL
<b>QUERCUS ROBUR WHOLE</b> (UNII: R7QMG0BT2W) (QUERCUS ROBUR WHOLE - UNII:R7QMG0BT2W)	QUERCUS ROBUR WHOLE	4 [hp_X] in 1 mL
<b>LEAD</b> (UNII: 2P299V784P) (LEAD - UNII:2P299V784P)	LEAD	12 [hp_X] in 1 mL
<b>LOBELIA SPICATA LEAF</b> (UNII: 1G4GK01F67) (LOBELIA SPICATA LEAF - UNII:1G4GK01F67)	LOBELIA SPICATA LEAF	6 [hp_X] in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>ALCOHOL</b> (UNII: 3K9958V90M)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-6090-3	60 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	09/01/2009	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		09/01/2009	

**Labeler** - Uriel Pharmacy, Inc. (043471163)

### Establishment

Name	Address	ID/FEI	Business Operations
Uriel Pharmacy, Inc.		043471163	manufacture(48951-6090)

Revised: 11/2024

Uriel Pharmacy, Inc.