LAPIS LAZULI 8129- lapis lazuli liquid Professional Complementary Health Formulas

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.

ULAP

ACTIVE INGREDIENTS

Lapis lazuli 6X, 8X, 12X, 30X

QUESTIONS

Professional Formulas

PO Box 2034 Lake Oswego, OR 97035

INDICATIONS

Temporarily relieves difficulty in communication or understanding, confusion, or past emotional upset or anxiousness.*

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

WARNINGS

In case of overdose, get medical help or contact a poison control center right away.

Keep out of the reach of children.

If pregnant or breastfeeding, ask a healthcare professional before use.

DIRECTIONS

Place drops under tongue 30 minutes before/after meals. Adults and children 12 years and over: Take 10 drops up to 3 times per day. Consult a physician for use in children under 12 years of age.

OTHER INFORMATION

Tamper resistant. If seal is broken, do not use. After opening, close container tightly and store at room temperature away from heat.

INACTIVE INGREDIENTS

20% ethanol, purified water.

LABEL

Est 1985 Professional Formulas Complementary Health Lapis Lazuli Homeopathic Remedy 2 FL. OZ. (59 mL)



LAPIS LAZULI 8129										
lapis lazuli liquid										
Product Information										
Product Type	HUMAN OTC DRUG	Item Code (Source)		NDC:63083-8129						
Route of Administration	ORAL									
Active Ingredient/Active Moiety										
Ingredient Name			Basis of Strength		Strength					
CALCIUM HEXAFLUOROSILICATE (UNII: 2NVP93XVQ3) (CALCIUM HEXAFLUOROSILICATE - UNII:2NVP93XVQ3)			CALCIUM HEXAFLUOROS ILICATE		6 [hp_X] in 59 mL					
Inactivo Ingradiante										
Inactive Ingredients										
Ingredient Name				Strength						
ALCOHOL (UNII: 3K9958V90M)										
WATER (UNII: 059QF0K00R)										

Packaging							
# Item Code	Package Description	Marketing Start Date	Marketing End Date				
1 NDC:63083- 8129-2	59 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	08/15/1985					
Marketing	Information						
Marketing Marketing Category		Marketing Start Date	Marketing End Date				

Labeler - Professional Complementary Health Formulas (167339027)

Registrant - Natural Pharmaceutical Manufacturing LLC (015624923)

Establishment

Name	Address	ID/FEI	Business Operations
Natural Pharmaceutical Manufacturing LLC		015624923	manufacture(63083-8129)

Revised: 8/2019

Professional Complementary Health Formulas