KIDNEY STONE DROPS 2039- kidney stone drops 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.

C39

ACTIVE INGREDIENTS

Equisetum arvense 2X
Sarsaparilla 3X
Berberis vulgaris 6X
Crataegus oxyacantha 6X
Digitalis purpurea 6X
Kali carbonicum 6X
Lycopodium clavatum 6X
Nitricum acidum 6X
Mercurius corrosivus 12X
Oxalicum acidum 30X
Uricum acidum 30X

QUESTIONS

Professional Formulas

PO Box 2034 Lake Oswego, OR 97035

INDICATIONS

For the temporary relief of mild pain in the back or abdomen.*

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

WARNINGS

Severe or persistent symptoms may be a sign of a serious condition. Consult a doctor promptly if symptoms are severe, sharp, or worsen during urination. Keep out of the reach of children. In case of overdose, get medical help or contact a poison control center right away. If pregnant or breastfeeding, ask a healthcare professional before use.

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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

Kidney Stone Drops

Homeopathic Remedy

2 FL. OZ. (59 mL)

KIDNEY STONE DROPS 2039

kidney stone drops liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63083-2039

Route of Administration	ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
EQUISETUM ARVENSE TOP (UNII: 1DP6Y6B65Z) (EQUISETUM ARVENSE TOP - UNII:1DP6Y6B65Z)	EQUISETUM ARVENSE TOP	2 [hp_X] in 59 mL
SMILAX ORNATA ROOT (UNII: 2H1576D5WG) (SMILAX ORNATA ROOT - UNII:2H1576D5WG)	SMILAX ORNATA ROOT	3 [hp_X] in 59 mL
BERBERIS VULGARIS ROOT BARK (UNII: 1TH8Q20J0U) (BERBERIS VULGARIS ROOT BARK - UNII:1TH8Q20J0U)	BERBERIS VULGARIS ROOT BARK	6 [hp_X] in 59 mL
HAWTHORN LEAF WITH FLOWER (UNII: 6OM09RPY36) (HAWTHORN LEAF WITH FLOWER - UNII:6OM09RPY36)	HAWTHORN LEAF WITH FLOWER	6 [hp_X] in 59 mL
DIGITALIS (UNII: F1T8QT9U8B) (DIGITALIS - UNII:F1T8QT9U8B)	DIGITALIS	6 [hp_X] in 59 mL
POTASSIUM CARBONATE (UNII: BQN1B9B9HA) (CARBONATE ION - UNII:7UJQ5OPE7D)	POTASSIUM CARBONATE	6 [hp_X] in 59 mL
LYCOPODIUM CLAVATUM SPORE (UNII: C88X29Y479) (LYCOPODIUM CLAVATUM SPORE - UNII:C88X29Y479)	LYCOPODIUM CLAVATUM SPORE	6 [hp_X] in 59 mL
NITRIC ACID (UNII: 411VRN1TV4) (NITRIC ACID - UNII:411VRN1TV4)	NITRIC ACID	6 [hp_X] in 59 mL
MERCURIC CHLORIDE (UNII: 53GH7MZT1R) (MERCURIC CATION - UNII:ED30FJ8Y42)	MERCURIC CHLORIDE	12 [hp_X] in 59 mL
OXALIC ACID (UNII: 9E7R5L6H31) (OXALIC ACID - UNII:9E7R5L6H31)	OXALIC ACID	30 [hp_X] in 59 mL
URIC ACID (UNII: 268B43MJ25) (URIC ACID - UNII:268B43MJ25)	URIC ACID	30 [hp_X] in 59 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:63083-2039-2	59 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	08/15/1985	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		08/15/1984	

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(63083-2039)