DNA INSULIN DROPS 2019- dna insulin 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.

C19

ACTIVE INGREDIENTS

Chromium muriaticum 6X
Lycopodium clavatum 6X
Vanadium metallicum 6X
Pancreatinum 6X, 12X, 30X, 100X
DNA insulin 12X, 60X, 100X, 1000X
Phosphoricum acidum 30X
Uranium nitricum 30X
Radium bromatum 60X

QUESTIONS

Professional Formulas

PO Box 2034 Lake Oswego, OR 97035

INDICATIONS

For the temporary relief of increased hunger or thirst, frequent urination, fatigue, nausea, or vomiting.*

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

WARNINGS

Consult a doctor if condition worsens or if symptoms persist. 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

DNA Insulin Drops

Homeopathic Remedy

2 FL. OZ. (59 mL)

DNA INSULIN DROPS 2019

dna insulin drops liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63083-2019
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CHROMIUM (UNII: 0R0008Q3JB) (CHROMIUM - UNII:0R0008Q3JB)	CHROMIUM	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
VANADIUM (UNII: 00J9J9XKDE) (VANADIUM - UNII:00J9J9XKDE)	VANADIUM	6 [hp_X] in 59 mL
PANCRELIPASE AMYLASE (UNII: YOJ58O116E) (PANCRELIPASE AMYLASE - UNII:YOJ58O116E)	PANCRELIPASE AMYLASE	6 [hp_X] in 59 mL
INSULIN GLULISINE (UNII: 7XIY785AZD) (INSULIN GLULISINE - UNII:7XIY785AZD)	INSULIN GLULISINE	12 [hp_X] in 59 mL
PHOSPHORIC ACID (UNII: E4GA8884NN) (PHOSPHORIC ACID - UNII:E4GA8884NN)	PHOSPHORIC ACID	30 [hp_X] in 59 mL
URANYL NITRATE HEXAHYDRATE (UNII: 3V057702FY) (URANIUM CATION (6+) - UNII:5PI36AS4G7)	URANYL NITRATE HEXAHYDRATE	30 [hp_X] in 59 mL
RADIUM BROMIDE (UNII: R74O7T8569) (RADIUM CATION - UNII:05456MVL7T)	RADIUM BROMIDE	60 [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:63083-2019-2	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(63083-2019)