TROPICAL 1022- tropical 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.

ATH

ACTIVE INGREDIENTS

Adrenal 6X
ACTH 6X, 30X
Histaminum hydrochloricum 12X
Liver 6X, 12X
and all the following at 6X, 12X, 60X, 100X:
Acacia
Bermuda grass
Cocklebur
English plantain
Eucalyptus
Kentucky bluegrass
Mountain cedar
Olive pollen
Pigweed mix
Queen palm
Ragweed
Saltbush
White mulberry

QUESTIONS

Professional Formulas

PO Box 2034 Lake Oswego, OR 97035

INDICATIONS

For the temporary relief of runny nose, sneezing, itching of the nose or throat, and itchy, watery eyes due to sensitivity to common allergens in the tropics and Hawaii.*

*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 to 15 drops up to 3 times per day. For desensitization, begin with 1 to 5 drops daily, increasing to the standard dose gradually to avoid symptom expression; after 1 to 3 months at the standard dose, decrease gradually to a maintenance dose of 10 to 15 drops weekly. 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

Tropical

Homeopathic Remedy

2 FL. OZ. (59 mL)

TROPICAL 1022

tropical liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63083-1022
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SUS SCROFA ADRENAL GLAND (UNII: 398IYQ16YV) (SUS SCROFA ADRENAL GLAND - UNII:398IYQ16YV)	SUS SCROFA ADRENAL GLAND	6 [hp_X] in 59 mL
CORTICOTROPIN (UNII: K0U68Q2TXA) (CORTICOTROPIN - UNII:K0U68Q2TXA)	CORTICOTROPIN	6 [hp_X] in 59 mL
HISTAMINE DIHYDROCHLORIDE (UNII: 3POA0Q644U) (HISTAMINE - UNII:820484N8I3)	HISTAMINE DIHYDROCHLORIDE	12 [hp_X] in 59 mL
BEEF LIVER (UNII: W8N8R55022) (BEEF LIVER - UNII:W8N8R55022)	BEEF LIVER	6 [hp_X] in 59 mL
ACACIA (UNII: 5C5403N26O) (ACACIA - UNII:5C5403N26O)	ACACIA	6 [hp_X] in 59 mL
CYNODON DACTYLON WHOLE (UNII: 2Q8MR21NHK) (CYNODON DACTYLON WHOLE - UNII:2Q8MR21NHK)	CYNODON DACTYLON WHOLE	6 [hp_X] in 59 mL
XANTHIUM STRUMARIUM VAR. CANADENSE POLLEN (UNII: 0ZK6G3W3BI) (XANTHIUM STRUMARIUM VAR. CANADENSE POLLEN - UNII:0ZK6G3W3BI)	XANTHIUM STRUMARIUM VAR. CANADENSE POLLEN	6 [hp_X] in 59 mL
PLANTAGO LANCEOLATA LEAF (UNII: 2YWL9J7EE8) (PLANTAGO LANCEOLATA LEAF - UNII:2YWL9J7EE8)	PLANTAGO LANCEOLATA LEAF	6 [hp_X] in 59 mL
EUCALYPTUS GUM (UNII: 72T9EZC2VX) (EUCALYPTUS GUM - UNII:72T9EZC2VX)	EUCALYPTUS GUM	6 [hp_X] in 59 mL
POA PRATENSIS TOP (UNII: 7EA48700V9) (POA PRATENSIS TOP - UNII:7EA48700V9)	POA PRATENSIS TOP	6 [hp_X] in 59 mL
JUNIPERUS SCOPULORUM POLLEN (UNII: 0G82TT8ZFY) (JUNIPERUS SCOPULORUM POLLEN - UNII:0G82TT8ZFY)	JUNIPERUS SCOPULORUM POLLEN	6 [hp_X] in 59 mL
OLEA EUROPAEA FLOWER (UNII: 498M34P1VZ) (OLEA EUROPAEA FLOWER - UNII:498M34P1VZ)	OLEA EUROPAEA FLOWER	6 [hp_X] in 59 mL
AMARANTHUS HYBRIDUS LEAF (UNII: 07L86FJ69J) (AMARANTHUS HYBRIDUS LEAF - UNII:07L86FJ69J)	AMARANTHUS HYBRIDUS LEAF	6 [hp_X] in 59 mL
SYAGRUS ROMANZOFFIANA WHOLE (UNII: 3I21564JAW) (SYAGRUS ROMANZOFFIANA WHOLE - UNII:3I21564JAW)	SYAGRUS ROMANZOFFIANA WHOLE	6 [hp_X] in 59 mL
AMBROSIA ARTEMISIIFOLIA WHOLE (UNII: 9W34L2CQ9A) (AMBROSIA ARTEMISIIFOLIA WHOLE - UNII:9W34L2CQ9A)	AMBROSIA ARTEMISIIFOLIA WHOLE	6 [hp_X] in 59 mL
ATRIPLEX POLYCARPA WHOLE (UNII: 12IGX5TK1H) (ATRIPLEX POLYCARPA WHOLE - UNII:12IGX5TK1H)	ATRIPLEX POLYCARPA WHOLE	6 [hp_X] in 59 mL
WHITE MULBERRY (UNII: MN25R0HH5A) (WHITE MULBERRY - UNII:MN25R0HH5A)	WHITE MULBERRY	6 [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:63083-1022-2	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 (015624923)

Establishment

Name	Address	ID/FEI	Business Operations
Natural Pharmaceutical Manufacturing		015624923	manufacture(63083-1022)