MYTOX 6021- mytox 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.

X21

ACTIVE INGREDIENTS

Borax 4X
Colchicum autumnale 4X
Agaricus muscarius 6X
Arsenicum album 6X
Drosera rotundifolia 6X
Bufo rana 8X
Coenzyme Q10 12X

QUESTIONS

Professional Formulas

PO Box 2034 Lake Oswego, OR 97035

INDICATIONS

For the temporary relief of red or itchy skin, cough, joint or muscle pain or stiffness, shortness of breath, sneezing, occasional headache, dizziness, or fatigue.*

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

WARNINGS

Persistent symptoms may be a sign of a serious condition. If symptoms persist or are accompanied by a fever, rash, or persistent headache, consult a doctor. 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.

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

Mytox

Homeopathic Remedy

2 FL. OZ. (59 mL)

MYTOX 6021

mytox liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63083-6021
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM BORATE (UNII: 91MBZ8H3QO) (BORATE ION - UNII:44OAE30D22)	SODIUM BORATE	4 [hp_X] in 59 mL
COLCHICUM AUTUMNALE BULB (UNII: 993QHL78E6) (COLCHICUM AUTUMNALE BULB - UNII:993QHL78E6)	COLCHICUM AUTUMNALE BULB	4 [hp_X] in 59 mL
AMANITA MUSCARIA FRUITING BODY (UNII: DIF093I037) (AMANITA MUSCARIA FRUITING BODY - UNII:DIF093I037)	AMANITA MUSCARIA FRUITING BODY	6 [hp_X] in 59 mL
ARSENIC TRIOXIDE (UNII: S7V92P67HO) (ARSENIC CATION (3+) - UNII:C96613F5AV)	ARSENIC TRIOXIDE	6 [hp_X] in 59 mL
DROSERA ROTUNDIFOLIA WHOLE (UNII: QR44N9XPJQ) (DROSERA ROTUNDIFOLIA WHOLE - UNII:QR44N9XPJQ)	DROSERA ROTUNDIFOLIA WHOLE	6 [hp_X] in 59 mL
BUFO BUFO CUTANEOUS GLAND (UNII: Q59QU6N72Q) (BUFO BUFO CUTANEOUS GLAND - UNII:Q59QU6N72Q)	BUFO BUFO CUTANEOUS GLAND	8 [hp_X] in 59 mL
UBIDECARENONE (UNII: EJ27X76M46) (UBIDECARENONE - UNII:EJ27X76M46)	UBIDECARENONE	12 [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:63083-6021-2	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(63083-6021)