

MYTOX- borax, colchicum autumnale, drosera (rotundifolia), arsenicum album, ubidecarenonum, agaricus muscarius, bufo rana spray
Nutritional Specialties, Inc.

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.

DRUG FACTS:

ACTIVE INGREDIENTS:

Borax 4X, Colchicum Autumnale 4X, Drosera (Rotundifolia) 6X, Arsenicum Album 8X, Ubidecarenonum (CoQ 10) 8X, Agaricus Muscarius 12X, Bufo Rana 12X.

PURPOSE:

Aids in temporary relief of symptoms associated with fungal conditions, such as night sweats, headache, fatigue, muscle aches, joint pain, and itchy, red or scaly skin†

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

WARNINGS:

Professional Use Only

If pregnant or breast-feeding, ask a health professional before use.

In case of overdose, get medical help or contact a Poison Control Center right away.

If condition worsens, seek medical attention.

KEEP OUT OF REACH OF CHILDREN

Do not use if tamper evident seal is broken or missing.

Store in a cool place after opening

KEEP OUT OF REACH OF CHILDREN:

In case of overdose, get medical help or contact a Poison Control Center right away.

DIRECTIONS:

Adults: 2 to 3 sprays orally, three times daily. Children under twelve one half adult dosage. Do not take within 15 minutes of consuming food, beverage or brushing teeth. Consult a physician for use in children under 12 years of age.

INDICATIONS:

Aids in temporary relief of symptoms associated with fungal conditions, such as night sweats, headache, fatigue, muscle aches, joint pain, and itchy, red or scaly skin†

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INACTIVE INGREDIENTS:

Alcohol USP 20%, Purified Water USP.

QUESTIONS:

**MANUFACTURED EXCLUSIVELY FOR
NUTRITIONAL SPECIALTIES, INC.**

PO BOX 97227

PITTSBURGH, PA 15229

www.phpltd.com

PACKAGE LABEL DISPLAY:

Professional

Health Products

HOMEOPATHIC

NDC 83027-0092-1

MYTOX

2 FL. OZ (60 ml)

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NDC 83027-0092-1

MYTOX

X21 (reorder code)

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MYTOX

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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83027-0092
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 1 mL
COLCHICUM AUTUMNALE BULB (UNII: 993QHL78E6) (COLCHICUM AUTUMNALE BULB - UNII:993QHL78E6)	COLCHICUM AUTUMNALE BULB	4 [hp_X] in 1 mL
DROSELA ROTUNDIFOLIA WHOLE (UNII: QR44N9XPJQ) (DROSELA ROTUNDIFOLIA - UNII:QR44N9XPJQ)	DROSELA ROTUNDIFOLIA WHOLE	6 [hp_X] in 1 mL
ARSENIC TRIOXIDE (UNII: S7V92P67HO) (ARSENIC CATION (3+) - UNII:C96613F5AV)	ARSENIC TRIOXIDE	8 [hp_X] in 1 mL
UBIDECARENONE (UNII: EJ27X76M46) (UBIDECARENONE - UNII:EJ27X76M46)	UBIDECARENONE	8 [hp_X] in 1 mL
AMANITA MUSCARIA FRUITING BODY (UNII: DIF093I037) (AMANITA MUSCARIA FRUITING BODY - UNII:DIF093I037)	AMANITA MUSCARIA FRUITING BODY	12 [hp_X] in 1 mL
BUFO BUFO CUTANEOUS GLAND (UNII: Q59QU6N72Q) (BUFO BUFO CUTANEOUS GLAND - UNII:Q59QU6N72Q)	BUFO BUFO CUTANEOUS GLAND	12 [hp_X] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83027-0092-1	60 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	08/02/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		08/02/2023	

Labeler - Nutritional Specialties, Inc. (032744609)

