

STO NOSODE 4008- sto nosode 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.

N8

ACTIVE INGREDIENTS

all the following at 30X, 60X, 100X:

Chlamydia
Gardnerella
Lymphogranuloma
Medorrhinum
Proteus vulgaris
Pseudomonas
Syphilinum
Trichomonas vaginalis

QUESTIONS

Professional Formulas

PO Box 2034 Lake Oswego, OR 97035

INDICATIONS

For the temporary relief of mild skin rash, itchy skin, changes in urination, chills, joint pain or body aches, nausea, 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 sores, blood in urine, or painful urination, 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 to 15 drops once weekly or monthly. If mild symptoms are present, 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

STO Nosode

Homeopathic Remedy

2 FL. OZ. (59 mL)



STO NOSODE 4008

sto nosode liquid

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:63083-4008

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CHLAMYDIA TRACHOMATIS (UNII: T6NI39QU44) (CHLAMYDIA TRACHOMATIS - UNII:T6NI39QU44)	CHLAMYDIA TRACHOMATIS	30 [hp_X] in 59 mL
GARDNERELLA VAGINALIS (UNII: 238RQ26259) (GARDNERELLA VAGINALIS - UNII:238RQ26259)	GARDNERELLA VAGINALIS	30 [hp_X] in 59 mL
GONORRHEAL URETHRAL SECRETION HUMAN (UNII: 9BZG9E3I8F) (GONORRHEAL URETHRAL SECRETION HUMAN - UNII:9BZG9E3I8F)	GONORRHEAL URETHRAL SECRETION HUMAN	30 [hp_X] in 59 mL
PROTEUS VULGARIS (UNII: 11T9HCO30O) (PROTEUS VULGARIS - UNII:11T9HCO30O)	PROTEUS VULGARIS	30 [hp_X] in 59 mL
PSEUDOMONAS AERUGINOSA (UNII: Y793W5V55N) (PSEUDOMONAS AERUGINOSA - UNII:Y793W5V55N)	PSEUDOMONAS AERUGINOSA	30 [hp_X] in 59 mL
TREPONEMIC SKIN CANKER HUMAN (UNII: 4ZWP7FW8W) (TREPONEMIC SKIN CANKER HUMAN - UNII:4ZWP7FW8W)	TREPONEMIC SKIN CANKER HUMAN	30 [hp_X] in 59 mL
TRICHOMONAS VAGINALIS (UNII: Q25X3G314L) (TRICHOMONAS VAGINALIS - UNII:Q25X3G314L)	TRICHOMONAS VAGINALIS	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:63083-4008-2	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(63083-4008)