

SILICONE DETOX 7513- silicone detox 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.

SILCNE

ACTIVE INGREDIENTS

Silicone 6X, 12X, 30X

QUESTIONS

Professional Formulas

PO Box 2034 Lake Oswego, OR 97035

INDICATIONS

For the temporary relief of fatigue, weakness, mild joint or muscle pain, dry eyes, minor abdominal pain, or occasional headache due to sensitivity to or exposure to silicone.*

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

WARNINGS

Consult a doctor if condition worsens or 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

Silicone Detox

Homeopathic Remedy

2 FL. OZ. (59 mL)



SILICONE DETOX 7513

silicone detox liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63083-7513
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TRIMETHYLSILYL TREATED DIMETHICONOL/TRIMETHYLSILOXYSILICATE CROSSPOLYMER (35/65 W/W; 5000000 PA.S) (UNII: 0A6MDS9SLT) (TRIMETHYLSILYL TREATED DIMETHICONOL/TRIMETHYLSILOXYSILICATE CROSSPOLYMER (35/65 W/W; 5000000 PA.S) - UNII:0A6MDS9SLT)	TRIMETHYLSILYL TREATED DIMETHICONOL/TRIMETHYLSILOXYSILICATE CROSSPOLYMER (35/65 W/W; 5000000 PA.S)	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:63083-7513-2	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(63083-7513)

Revised: 1/2026

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