## SPA/POOL DETOX 6027- spa/pool 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.

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**X27** 

## **ACTIVE INGREDIENTS**

Sodium carbonate 12X
Pool water 12X, 30X, 60X
Spa water 12X, 30X, 60X
Algaecide 30X
Brominating tablets 30X
Bromium 30X
Clarifier 30X
Chlorinum 30X
Spa oxidizer 30X
Spa stain/scale 30X
Sodium bisulfate 30X
Cryptosporidium 30X, 60X, 100X
Escherichia coli 30X, 60X, 100X

## **QUESTIONS**

Professional Formulas

PO Box 2034 Lake Oswego, OR 97035

#### INDICATIONS

For the temporary relief of skin irritations, red or watery eyes, minor earache, stomach cramps, diarrhea, or vomiting due to sensitivity to or exposure to pool or spa chemicals or contaminants.\*

\*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

Spa/Pool Detox

Homeopathic Remedy

2 FL. OZ. (59 mL)



## **SPA/POOL DETOX 6027**

spa/pool detox liquid

#### **Product Information**

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63083-6027
Route of Administration	ORAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
<b>SODIUM CARBONATE</b> (UNII: 45P3261C7T) (CARBONATE ION - UNII: 7UJQ50PE7D)	SODIUM CARBONATE	12 [hp_X] in 59 mL	
CHLORINE (UNII: 4R7X1O2820) (CHLORINE - UNII:4R7X1O2820)	CHLORINE	12 [hp_X] in 59 mL	
CUPRIC CATION (UNII: 8CBV67279L) (CUPRIC CATION - UNII:8CBV67279L)	CUPRIC CATION	30 [hp_X] in 59 mL	
BROMINE (UNII: SBV4XY874G) (BROMINE - UNII:SBV4XY874G)	BROMINE	30 [hp_X] in 59 mL	
<b>AMMONIUM CATION</b> (UNII: 54S68520I4) (AMMONIUM CATION - UNII:54S68520I4)	AMMONIUM CATION	30 [hp_X] in 59 mL	
SODIUM CATION (UNII: LYR4M0NH37) (SODIUM CATION - UNII:LYR4M0NH37)	SODIUM CATION	30 [hp_X] in 59 mL	
ETIDRONIC ACID (UNII: M2F465ROXU) (ETIDRONIC ACID - UNII:M2F465ROXU)	ETIDRONIC ACID	30 [hp_X] in 59 mL	
CRYPTOSPORIDIUM HOMINIS (UNII: DKE8M34J72) (CRYPTOSPORIDIUM HOMINIS - UNII:DKE8M34J72)	CRYPTOS PORIDIUM HOMINIS	30 [hp_X] in 59 mL	
ESCHERICHIA COLI (UNII: 514B9K0L10) (ESCHERICHIA COLI - UNII:514B9K0L10)	ESCHERICHIA COLI	30 [hp_X] in 59 mL	

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

ı	Packaging			
1	# Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:63083- 6027-2	59 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	08/15/1985	

Marketing Information			
Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
	08/15/1984		
	Application Number or Monograph	Application Number or Monograph Marketing Start Citation Date	

# **Labeler -** Professional Complementary Health Formulas (167339027)

## **Registrant -** Natural Pharmaceutical Manufacturing LLC (015624923)

Establishment				
Name	Address	ID/FEI	<b>Business Operations</b>	
Natural Pharmaceutical Manufacturing LLC		015624923	manufacture(63083-6027)	

Revised: 8/2019

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