DENTASTAT- allium sativum, berberis vulgaris, fucus vesiculosus, phytolacca decandra, argentum metallicum, aurum metallicum, calcarea fluorica, calcarea phosphorica, iodium, mercurius solubilis, niccolum metallicum, silicea, zincum metallicum, platinum metallicum liquid Energique, 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:

(in each drop): 9.06% of Argentum Metallicum 12X, Aurum Metallicum 12X, Calcarea Fluorica 12X, Calcarea Phosphorica 12X, Iodium 12X, Mercurius Solubilis 12X, Niccolum Metallicum 12X, Phytolacca Decandra 6X, Platinum Metallicum 16X, Silicea 12X, Zincum Metallicum 12X; 0.10% of Allium Sativum 3X, Berberis Vulgaris 3X, Fucus Vesiculosus 3X.

INDICATIONS:

May temporarily relieve symptoms associated with dental procedures, such as pain and minor inflammation in the mouth or on the gums.**

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

WARNINGS:

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

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

Do not use if tamper evident seal is broken or missing. Store in a cool, dry place.

KEEP OUT OF REACH OF CHILDREN:

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

DIRECTIONS:

Adults and children 5 to 10 drops orally, 3 times daily or as otherwise directed by a health care professional. If symptoms persist for more than 7 days, consult your health care professional. Consult a physician for use in children under 12 years of age.

INDICATIONS:

May temporarily relieve symptoms associated with dental procedures, such as pain and minor inflammation in the mouth or on the gums.**

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

INACTIVE INGREDIENTS:

Demineralized water, 20% Ethanol.

QUESTIONS:

Dist. by Energique, Inc. 201 Apple Blvd Woodbine, IA 51579 800.869.8078

PACKAGE LABEL DISPLAY:

ENERGIQUE

SINCE 1987

HOMEOPATHIC REMEDY

DENTASTAT

1 fl. oz. (30 ml)

WARNINGS: If pregnant or breastfeeding, ask a health professional before use. Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away. Do not use if tamper evident seal is broken or missing. Store in a cool, dry place.

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DENTASTAT™

1 fl. oz. (30 ml) 20% Ethanol

Active Ingredients (in each drop): 9.06% of Arg Met 12X, Aur Met 12X, Calc Fluor 12X, Calc Phos 12X, Iodium 12X, Merc Solub 12X, Niccolum Met 12X, Phytolacca 6X, Platinum Met 16X, Silicea 12X, Zinc Met 12X; 0.10% of Allium Sat 3X, Berber Vulg 3X, Fucus 3X.

Inactive Ingredients:

Demineralized water, 20% Ethanol.

DIRECTIONS: Adults and children 5 to 10 drops orally, 3 times daily or as otherwise directed by a health care professional. If symptoms persist for more than 7 days, consult your health care professional. Consult a physician for use in children under 12 years of age.

LOT: XXXXXXX

DENTASTAT

allium sativum, berberis vulgaris, fucus vesiculosus, phytolacca decandra, argentum metallicum, aurum metallicum, calcarea fluorica, calcarea phosphorica, iodium, mercurius solubilis, niccolum

metallicum, silicea, zincum metallicum, platinum metallicum liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:44911-0555
Route of Administration	ORAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
GARLIC (UNII: V1V998DC17) (GARLIC - UNII:V1V998DC17)	GARLIC	3 [hp_X] in 1 mL	
BERBERIS VULGARIS ROOT BARK (UNII: 1TH8Q20J0U) (BERBERIS VULGARIS ROOT BARK - UNII:1TH8Q20J0U)	BERBERIS VULGARIS ROOT BARK	3 [hp_X] in 1 mL	
FUCUS VESICULOSUS (UNII: 535G2ABX9M) (FUCUS VESICULOSUS - UNII:535G2ABX9M)	FUCUS VESICULOSUS	3 [hp_X] in 1 mL	
PHYTOLACCA AMERICANA ROOT (UNII: 11E6VI8VEG) (PHYTOLACCA AMERICANA ROOT - UNII:11E6VI8VEG)	PHYTOLACCA AMERICANA ROOT	6 [hp_X] in 1 mL	
SILVER (UNII: 3M4G523W1G) (SILVER - UNII:3M4G523W1G)	SILVER	12 [hp_X] in 1 mL	
GOLD (UNII: 79Y1949PYO) (GOLD - UNII:79Y1949PYO)	GOLD	12 [hp_X] in 1 mL	
CALCIUM FLUORIDE (UNII: O3B55K4YKI) (FLUORIDE ION - UNII:Q80VPU4080)	CALCIUM FLUORIDE	12 [hp_X] in 1 mL	
TRIBASIC CALCIUM PHOSPHATE (UNII: 91D9GV0Z28) (CALCIUM CATION - UNII: 2M83C4R6ZB)	CALCIUM CATION	12 [hp_X] in 1 mL	
IODINE (UNII: 9679TC07X4) (IODINE - UNII:9679TC07X4)	IODINE	12 [hp_X] in 1 mL	
MERCURIUS SOLUBILIS (UNII: 324Y4038G2) (MERCURIUS SOLUBILIS - UNII: 324Y4038G2)	MERCURIUS SOLUBILIS	12 [hp_X] in 1 mL	
NICKEL (UNII: 70V03QG267) (NICKEL - UNII:70V03QG267)	NICKEL	12 [hp_X] in 1 mL	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4) (SILICON DIOXIDE - UNII:ETJ7Z6XBU4)	SILICON DIOXIDE	12 [hp_X] in 1 mL	
ZINC (UNII: J41CSQ7QDS) (ZINC - UNII:J41CSQ7QDS)	ZINC	12 [hp_X] in 1 mL	
PLATINUM (UNII: 49DFR088MY) (PLATINUM - UNII:49DFR088MY)	PLATINUM	16 [hp_X] in 1 mL	

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

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:44911- 0555-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	08/12/2020	08/25/2025	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		08/12/2020	08/25/2025
homeopathic		00/12/2020	00,23,2023

Labeler - Energique, Inc. (789886132)

Registrant - Apotheca Company (844330915)

Establishment			
Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(44911-0555), api manufacture(44911-0555), label(44911-0555), pack(44911-0555)

Revised: 10/2022 Energique, Inc.