SHARK CARTILAGE CORD- shark cartilage 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 INGREDIENT:

(in each drop): 49.95% of Shark Cartilage 12C, 30C; 0.10% of Shark Cartilage 6C.

INDICATIONS:

May temporarily relieve minor joint pain and stiffness.**

**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, 1 time 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:

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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 SHARK CARTILAGE CORD 2 fl. oz. (60 ml)

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LOT: XXXXXX MFD: MM/YY



SHARK CARTILAGE CORD shark cartilage liquid								
Product Information								
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:44911-0260					
Route of Administration	ORAL							

		Ingredient Name		Basis of Strength	Stronath
	IARK CARTILAG	E (UNII: D2YCN1I522) (SHARK CARTILAGE -		SHARK CARTILAG	6 [hp_C] in 1 mL
In	active Ingr	edients			
		Ingredient Name	Strength		
w	ATER (UNII: 059	QF0KO0R)			
AL	COHOL (UNII: 3	K9958V90M)			
Pa	ackaging				
#	Item Code	Package Description	Ma	arketing Starl Date	t Marketing En Date
1	NDC:44911- 0260-1	60 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	09/0	04/2015	05/11/2027
Μ	larketing	Information			
	Marketing Category	Application Number or Monograph Citation	Ma	rketing Start Date	Marketing End Date
un	approved meopathic		09/04	/2015	05/11/2027

Labeler - Energique, Inc. (789886132)

Registrant - Apotheca Company (844330915)

Establishment

Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(44911-0260) , api manufacture(44911-0260) , label(44911-0260) , pack(44911-0260)

Revised: 4/2023

Energique, Inc.