

HEPATA-CHORD- homeopathic liquid liquid

Energetix Corporation

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.

Hepata-Chord

Active ingredients 59.1 mL contains 3.22% of: Aceticum ac 18X; Acetylsalicylicum ac 18X; Alumina 18X; Benzinum 12X; Berberis vulg 4X; Carduus mar 4X; Chelidonium maj 4X; Chelone 12X; Chloramphenicolum 18X; Chlorpromazinum 18X; Cholesterinum 12X; Cortisone aceticum 18X; Folliculinum 18X; Gambogia 12X; Hepar suis 9X, 12X, 6C; Hydrocotyle 5X; Hypothalamus 12X, 6C; Lappa 4X; Nat sulphuricum 12X, 30X; Petroleum 18X; Phenacetinum 18X; Phos 12X; Silicea 12X; Taraxacum 4X; Thyroidinum 9X, 12X, 6C.

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

Uses

Temporary relief of skin rash, right-side abdominal discomfort, headache

Warnings

- In case of overdose, get medical help or contact a Poison Control Center right away.
- **If pregnant or breast feeding**, ask a health professional before use.
- **Keep out of reach of children.**

Directions

- Take 30 drops orally twice daily or as directed by a healthcare professional.
- Consult a physician for use in children under 12 years of age or if symptoms worsen or persist.

Other information

- Store at room temperature out of direct sunlight.
- Do not use if neck wrap is broken or missing.
- Shake well before use.

Inactive ingredients

Alcohol, Vegetable Glycerin, Purified Water.

Distributed by Energetix Corp.

Dahlonega, GA 30533

Questions? Comments?

800.990.7085

www.energetix.com

energetix®

Hepata-Chord

Homeopathic Remedy

Skin rash, right-side abdominal discomfort, headache

2 fl oz (59.1 mL) / 15% Alcohol

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Drug Facts (continued)
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Purpose Temporary relief on skin rash, right-side abdominal discomfort, headache

HEPATA-CHORD			
homeopathic liquid liquid			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:64578-0176
Route of Administration	ORAL		
Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ACETIC ACID (UNII: Q40Q9N063P) (ACETIC ACID - UNII:Q40Q9N063P)	ACETIC ACID	18 [hp_X] in 59.1 mL	
ASPIRIN (UNII: R16CO5Y76E) (ASPIRIN - UNII:R16CO5Y76E)	ASPIRIN	18 [hp_X] in 59.1 mL	
ALUMINUM OXIDE (UNII: LMI26O6933) (ALUMINUM OXIDE - UNII:LMI26O6933)	ALUMINUM OXIDE	18 [hp_X] in 59.1 mL	
BENZENE (UNII: J64922108F) (BENZENE - UNII:J64922108F)	BENZENE	12 [hp_X] in 59.1 mL	
BERBERIS VULGARIS ROOT BARK (UNII: 1TH8Q20J0U) (BERBERIS VULGARIS ROOT BARK - UNII:1TH8Q20J0U)	BERBERIS VULGARIS ROOT BARK	4 [hp_X] in 59.1 mL	
MILK THISTLE (UNII: U946SH95EE) (MILK THISTLE - UNII:U946SH95EE)	MILK THISTLE	4 [hp_X] in 59.1 mL	
CHELIDONIUM MAJUS (UNII: 7E889U5RNN) (CHELIDONIUM MAJUS - UNII:7E889U5RNN)	CHELIDONIUM MAJUS	4 [hp_X] in 59.1 mL	
CHELONE GLABRA (UNII: 6G3UN932VF) (CHELONE GLABRA -	CHELONE GLABRA	12 [hp_X]	

UNII:6G3UN932VF)	CHLORAMPHENICOL	in 59.1 mL
CHLORAMPHENICOL (UNII: 66974FR9Q1) (CHLORAMPHENICOL - UNII:66974FR9Q1)	CHLORAMPHENICOL	18 [hp_X] in 59.1 mL
CHLORPROMAZINE (UNII: U42B7VYA4P) (CHLORPROMAZINE - UNII:U42B7VYA4P)	CHLORPROMAZINE	18 [hp_X] in 59.1 mL
CHOLESTEROL (UNII: 97C5T2UQ7J) (CHOLESTEROL - UNII:97C5T2UQ7J)	CHOLESTEROL	12 [hp_X] in 59.1 mL
CORTISONE ACETATE (UNII: 883WKN7W8X) (CORTISONE - UNII:V27W9254FZ)	CORTISONE ACETATE	18 [hp_X] in 59.1 mL
ESTRONE (UNII: 2DI9HA706A) (ESTRONE - UNII:2DI9HA706A)	ESTRONE	18 [hp_X] in 59.1 mL
GAMBOGE (UNII: 7556HJ7587) (GAMBOGE - UNII:7556HJ7587)	GAMBOGE	12 [hp_X] in 59.1 mL
PORK LIVER (UNII: 6EC706HI7F) (PORK LIVER - UNII:6EC706HI7F)	PORK LIVER	9 [hp_X] in 59.1 mL
CENTELLA ASIATICA (UNII: 7M867G6T1U) (CENTELLA ASIATICA - UNII:7M867G6T1U)	CENTELLA ASIATICA	5 [hp_X] in 59.1 mL
BOS TAURUS HYPOTHALAMUS (UNII: S6G2NLH4Y7) (BOS TAURUS HYPOTHALAMUS - UNII:S6G2NLH4Y7)	BOS TAURUS HYPOTHALAMUS	12 [hp_X] in 59.1 mL
ARCTIUM LAPPA ROOT (UNII: 597E9BI3Z3) (ARCTIUM LAPPA ROOT - UNII:597E9BI3Z3)	ARCTIUM LAPPA ROOT	4 [hp_X] in 59.1 mL
SODIUM SULFATE (UNII: 0YPR65R21J) (SODIUM SULFATE ANHYDROUS - UNII:36KCS0R750)	SODIUM SULFATE	12 [hp_X] in 59.1 mL
KEROSENE (UNII: 1C89KKC04E) (KEROSENE - UNII:1C89KKC04E)	KEROSENE	18 [hp_X] in 59.1 mL
PHENACETIN (UNII: ER0CTH01H9) (PHENACETIN - UNII:ER0CTH01H9)	PHENACETIN	18 [hp_X] in 59.1 mL
PHOSPHORUS (UNII: 27YLU75U4W) (PHOSPHORUS - UNII:27YLU75U4W)	PHOSPHORUS	12 [hp_X] in 59.1 mL
SILICON DIOXIDE (UNII: ETJ7Z6XBU4) (SILICON DIOXIDE - UNII:ETJ7Z6XBU4)	SILICON DIOXIDE	12 [hp_X] in 59.1 mL
TARAXACUM OFFICINALE (UNII: 39981FM375) (TARAXACUM OFFICINALE - UNII:39981FM375)	TARAXACUM OFFICINALE	4 [hp_X] in 59.1 mL
THYROID, UNSPECIFIED (UNII: 0B4FDL9I6P) (THYROID, UNSPECIFIED - UNII:0B4FDL9I6P)	THYROID, UNSPECIFIED	9 [hp_X] in 59.1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALCOHOL (UNII: 3K9958V90M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:64578-0176-1	59.1 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	03/24/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved			

unapproved
homeopathic

03/24/2023

Labeler - Energetix Corporation (969572502)

Revised: 12/2024

Energetix Corporation