ARTHRITIS PAIN RELIEF- histamine dihydrochloride .025% cream
Walgreens

Walgreens - Arthritis Pain Relief Cream

Histamine Dihydrochloride 0.025%

Topical Analgesic

For temporary relief of minor aches and pains of muscles and joints associated with arthritis, simple backache, strains & bruises.

For external use only

Do not use on wounds or damaged skin or if you are allergic to ingredients in the products

When using this product

Avoid contact with eyes. If product gets into eyes, rinse thoroughly with water.

Do not bandage tightly or use a heating pad.

Stop use and ask a doctor if

Rash appears.

Condition worsens, if symptoms persist for more than 7 days, or if symptoms clear up and occur again within a few days.

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

Keep out of the reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

For Use by Adult and Children over 12 years of age.

Apply a thin layer to the pain site and message until thoroughly absorbed into skin.

Apply no more than 3 to 4 times daily.

Children 12 years or younger consult a physician

Store between 68°C and 77° F (20° and 25° C)

Acrylates/Acrylamide Copolymer)and) Mineral Oil (and) Polysorbate 85, Cetyl Alcohol, Chondroitin Sulfate, Dimethicone, disodium EDTA, Emu Oil, Glucosamine HCl, Glycerin, Helianthus Annus (Sunflower) Extract, Isopropyl Palmitate, Methyl Sulfonyl Methane (MSM), Phenoxyethanol, Propylene Glycol, Stearic Acid, Tocopheryl Actate, Triethanolamine, Water

ARTHRITIS PAIN RELIEF

histamine dihydrochloride .025% cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0363-3035
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
HISTAMINE DIHYDROCHLORIDE (UNII: 3POA0Q644U) (HISTAMINE -	HISTAMINE	0.025 g

UNII:820484N8I3)	DIHYDROCHLORIDE	in 100 g

Inactive Ingredients

Ingredient Name	Strength
GLUCOSAMINE HYDROCHLORIDE (UNII: 750W5330FY)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
EMU OIL (UNII: 344821WD61)	
HELIANTHUS ANNUUS FLOWERING TOP (UNII: BKJ0J3D1BP)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
TROLAMINE (UNII: 9O3K93S3TK)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL PALMITATE (UNII: 8CRQ2TH63M)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
CHONDROITIN SULFATE (PORCINE) (UNII: V5E8ELO4W9)	
MINERAL OIL (UNII: T5L8T28FGP)	
DIMETHYL SULFONE (UNII: 9H4PO4Z4FT)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-3035-25	1 in 1 CARTON	11/01/2022	
1		113 g in 1 JAR; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	11/01/2022	

Labeler - Walgreens (008965063)

Revised: 12/2024 Walgreens