GLUCOSAMINE CREAM EXTRA STRENGTH- histamine dihydrochloride cream
Q.A. Laboratories

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Glucosamine Cream EXTRA STRENGTH

Drug Facts

Active Ingredient

Histamine Dihydrochloride 0.05%

Purpose

Topical Analgesic

Uses For the temporary relief of minor aches and pains of muscles and joints associated with simple backache, arthritis, strains, bruises and sprains

Warnings For external use only

When using this product

- Avoid contact with eyes
- Do not bandage tightly

Do not use

- On wounds or damaged skin or if you are allergic to ingredients in this product.

Discontinue use and consult a physician if: Condition worsens or if symptoms persists for more than 7 days or clear up and occur again within a few days

Keep out of reach of children.

If swallowed get medical help or contact a poison control center

Directions

For use by adults and children over 12 years:

Apply to affected area not more than 3 to 4 times daily.

Inactive ingredients Deionized water, cetearyl alcohol, emu oil, glycerine, ceteareth-

20, glucosamine sulfate, dimethyl sulfone, caprylic capric triglyceride, phenoxyethanol, cetyl alcohol, methyl paraben, tocopherol acetate, butylene glycol, propylene glycol, chamomilla recutita flower extract, rosmarinus officinalis leaf extract, boswellia serrata extract, propyl paraben

Pain Relief On Contact

Dual Action Moisturizing Cream

Odorless, Greaseless, Stainless.

Made in USA

For re-orders call 800-280-5590

www.drnewtons.com

Distributed by

Dr. Newton's Naturals

26 Thomas Drive

Westbrook, ME 04092

Packaging

DR. NEWTON'S™
NATURALS

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www.drnewtons.com Net Wt. 3 oz. (85g) www.drnewtons.com

GLUCOSAMINE CREAM EXTRA STRENGTH

histamine dihydrochloride cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:52099-0008
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
HISTAMINE DIHYDROCHLORIDE (UNII: 3POA0Q644U) (HISTAMINE - UNII:820484N8I3)	HISTAMINE DIHYDROCHLORIDE	0.05 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
EMU OIL (UNII: 344821WD61)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLYOXYL 20 CETOSTEARYL ETHER (UNII: YRC528SWJY)	
GLUCOSAMINE SULFATE (UNII: 1FW7WLR731)	
DIMETHYL SULFONE (UNII: 9H4PO4Z4FT)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CHAMOMILE (UNII: FGL3685T2X)	
ROSEMARY (UNII: IJ67X351P9)	
INDIAN FRANKINCENSE (UNII: 4PW41QCO2M)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	

Product Characteristics

Color	white	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52099-0008-3	85 g in 1 TUBE; Type 0: Not a Combination Product	04/01/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	04/01/2017	

Labeler - Q.A. Laboratories (065361149)

Establishment

Name	Address	ID/FEI	Business Operations

Q.A. Laboratories		065361149	manufacture(52099-0008)