

## **PAIN RELIEVING ARTHRITIS- histamine dihydrochloride cream**

**Yash Pharmaceuticals**

*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.*

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### **Pain Relieving Arthritis Cream- Bulk**

#### **Drug Facts**

##### ***Active Ingredient***

Histamine dihydrochloride ..... 0.025%

##### ***Purpose***

External Analgesic

##### ***Uses***

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

##### ***Warnings***

- **For external use only.**
- **Do not use** on wounds or damaged skin or if you are allergic to ingredients in the product.

##### **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 reach of children.**

If swallowed, get medical help or contact a Poison Control Center right away.

##### ***Directions***

- **For Use by Adults and Children over 12 years.**
- Apply a thin layer to pain site and massage until thoroughly absorbed into skin. Apply no more than 3 to 4 times daily.
- **Children 12 years or younger consult a physician.**

***Other information***

- Store between 40°F and 86°F ( 4°C and 30C).
- Tamper Evident Feature: do not use if outer shrink wrap on jar is torn, broken or missing.

***Inactive ingredients***

arylamide, butylated hydroxyl toluene, cetostearyl alcohol , cetomacrogol 1000, disodium EDTA, glucosamine sulfate, glycerin, isohexadecane, polysorbate 80, propylene glycol, sodium acrylodimethyl taurate copolymer, titanium dioxide, transquitol P, vitamin E acetate, white petroleum jelly, water

**Questions or Comments? Call 1855-314-1850**

Distributed by: Velocity Pharma LLC  
Farmingdale, NY, 11735

**Pain Relieving Arthritis Cream**

**Net Wt. 4 oz (119g)-- Bulk of 144 in a case**

## arthritis pain relief cream ( histamine dihydrochloride 0.025% )

### Active Ingredient

Histamine dihydrochloride 0.025%..... Topical Analgesic

BATCH NO.	HDC701	QUANTITY	1x12x4x3x119 g= 144 jar
MFG. DATE	06/2017	SHIPPER NO.	21 out of 23
EXP. DATE	05/2019	GROSS WT.	25.395 kg.
		NET WT.	23.680 kg.

**WARNING:** KEEP OUT OF REACH OF CHILDREN AND PETS. FOR EXTERNAL USE ONLY.

STORE AT CONTROLLED TEMPERATURE OF 59°F to 86° F (15 to 30 °C )

CONFORMANCE WITH THE F.D & C ACT AND REGULATIONS THEREUNDER

PROTECT FROM DIRECT SUNLIGHT/MOISTURE/FREEZING

### MANUFACTURED BY:

YASH PHARMACEUTICALS  
 KHASRA NO. 19-M, VILLAGE :RAIPUR  
 PARGANA: BHAGWANPUR, DIST: HARIDWAR  
 ROORKEE, UTTARAKHAND, INDIA 247667

Lic#UK/Drugs/39/UA/2016

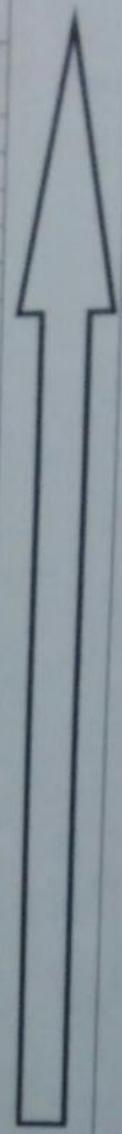
LABELLER CODE 71301

### SHIPPED TO:

VELOCITY PHARMA LLC  
 226/B SHERWOOD AVE,  
 FARMINGDALE , NY-11735

NDC No 76168-300-38

Barcode No.



## PAIN RELIEVING ARTHRITIS

histamine dihydrochloride cream

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:71301-004
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
HISTAMINE DIHYDRO CHLORIDE (UNII: 3POA0Q644U) (HISTAMINE - UNII:820484N8I3)	HISTAMINE DIHYDROCHLORIDE	0.025 g in 100 g

**Inactive Ingredients**

Ingredient Name	Strength
ACRYLAMIDE (UNII: 20R035KLCI)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128MIS)	
DISODIUM HYDROGEN CITRATE (UNII: 6FO62KCQ7A)	
EDETATE SODIUM (UNII: MP1J8420LU)	
GLUCOSAMINE SULFATE (UNII: 1FW7WLR731)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOHEXADECANE (UNII: 918X10UF1E)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
SODIUM ACRYLATE/SODIUM ACRYLOYLDIMETHYLTAURATE COPOLYMER (4000000 MW) (UNII: 1DXE3F3OZX)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
DIETHYLENE GLYCOL MONOETHYL ETHER (UNII: A1A1I8X02B)	
PETROLATUM (UNII: 4T6H12BN9U)	
WATER (UNII: 059QF0K00R)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71301-004-24	144 in 1 PACKAGE	06/06/2017	
1		119 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 not final	part348	05/26/2017	

**Labeler** - Yash Pharmaceuticals (871409551)**Registrant** - Velocity Pharma LLC (962198409)**Establishment**

Name	Address	ID/FEI	Business Operations
Yash Pharmaceuticals		871409551	manufacture(71301-004)