

ACTIVON ULTRA STRENGTH ARTHRITIS- menthol and histamine dihydrochloride stick
Family First Pharmaceuticals, Inc.

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.

ACTIVON®
Topical Analgesic
ULTRA STRENGTH ARTHRITIS

Drug Facts

<i>Active Ingredient</i>	<i>Purpose</i>
Histamine Dihydrochloride 0.028%	Topical Analgesic
Menthol 4.574%	Topical Analgesic

Uses

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

Warnings

For external use only.

Do not use

- otherwise than as directed
- if you are allergic to any ingredient in this product
- on a child 12 years of age with arthritis-like conditions
- with a heating pad

When using this product

- avoid contact with eyes, wounds, mucous membranes, broken or irritated skin
- do not share this product with anyone
- do not bandage tightly

Stop use and ask a doctor if

- condition worsens, or if symptoms persist for more than 7 days or clear up and occur again within a few days
- skin redness or excessive irritation of the skin develops

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.

Directions

- adults and children 12 years of age and older: apply to affected area not more than 3 to 4 times daily.
- children under 12 years of age: ask a doctor

Other information

Keep away from heat. Store between 15° and 30° C (59° and 86° F).

Inactive Ingredients

diazoldinyl urea, ethyl alcohol, iodopropynyl butylcarbamate, menthyl lactate, propylene glycol, sodium stearate, steareth-21, tetrasodium EDTA, triethenolamine, water

Questions ?

call **1-800-379-8870**, Weekdays 9AM to 5PM EST or visit us online at **activ-on.com**

Dist. by Family First Pharmaceuticals, Inc., Reno, NV 89502

PRINCIPAL DISPLAY PANEL - 57 g Carton

NEW

STRONGEST

ACTIVON[®] ARTHRITIS

ACTIVON[®]

Topical Analgesic

ULTRA

STRENGTH

ARTHRITIS

Powerful Pain Relief

for Arthritis & Joint & Muscle Pain

No-Mess

NDC 51068-501-01

NET WT 2 OZ (57 g)



ACTIVON ULTRA STRENGTH ARTHRITIS

menthol and histamine dihydrochloride stick

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:51068-501
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Menthol, unspecified form (UNII: L7T10EIP3A) (Menthol, unspecified form - UNII:L7T10EIP3A)	Menthol, unspecified form	0.04574 g in 1 g
Histamine Dihydrochloride (UNII: 3POA0Q644U) (Histamine - UNII:820484N8I3)	Histamine Dihydrochloride	0.00028 g in 1 g

Inactive Ingredients

Ingredient Name	Strength
diazolidinyl urea (UNII: H5RIZ3MPW4)	
alcohol (UNII: 3K9958V90M)	
iodopropynyl butylcarbamate (UNII: 603P14DHEB)	
menthyl lactate, (-) (UNII: 2BF9E65L7I)	
propylene glycol (UNII: 6DC9Q167V3)	
sodium stearate (UNII: QU7E2XA9TG)	
steareth-21 (UNII: 53J3F32P58)	
Edetate Sodium (UNII: MP1J8420LU)	
trolamine (UNII: 9O3K93S3TK)	
water (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51068-501-01	1 in 1 CARTON		
1		57 g in 1 CANISTER; 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	09/01/2011	

Labeler - Family First Pharmaceuticals, Inc. (832435809)**Establishment**

Name	Address	ID/FEI	Business Operations
RNA Pharma, LLC		079103999	MANUFACTURE(51068-501)

Revised: 4/2015

Family First Pharmaceuticals, Inc.