

ACTIVON ULTRA STRENGTH ARTHRITIS- menthol, unspecified form 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® Ultra Strength Arthritis

Drug Facts

Active Ingredient

Menthol 5.138%

Purpose

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 under 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

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

Questions ?

call 1-800-379-8870, Weekdays 9AM to 5PM EST

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

PRINCIPAL DISPLAY PANEL - 57 g Canister Carton

NEW STRONGEST

ACTIVON® ARTHRITIS

ACTIVON®

Topical Analgesic

ULTRA

STRENGTH

ARTHRITIS

**Powerful Pain Relief
for Arthritis &
Joint & Muscle Pain**

No-Mess

NDC 51068-507-01

NET WT 2 OZ (57 g)

ACTIVON[®]

Topical Analgesic

ULTRA STRENGTH ARTHRITIS

ACTIVON[®]

Topical Analgesic

ULTRA STRENGTH ARTHRITIS

- ✦ NO UNPLEASANT ODOR
- ✦ NON-GREASY
- ✦ NO NEED TO RUB IN
- ✦ CLEAR, NON-STAINING
- ✦ NO-MESS APPLICATOR

DEEP PENETRATING PAIN RELIEF



NEW STRONGEST
ACTIVON[®] ARTHRITIS

ACTIVON[®]

Topical Analgesic

ULTRA STRENGTH ARTHRITIS





**Apply directly
where it hurts**

**FOR ARTHRITIS &
Joint & Muscle Pain**

No-Mess

NDC 51068-507-01

NET WT 2 OZ (57 g)



ACTIVON[®]
Topical Analgesic

ULTRA STRENGTH ARTHRITIS

**Powerful Arthritis &
Joint & Muscle
Pain Relief**



KNEE



SHOULDER

ACTIVON[®]
Topical Analgesic

ULTRA STRENGTH ARTHRITIS

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Keep carton. It contains important information.
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Tamper-Evident:
Do Not Use if safety wrap around tube is broken or missing.

ACTIVON ULTRA STRENGTH ARTHRITIS

menthol, unspecified form stick

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:51068-507
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	0.05138 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:51068-507-01	1 in 1 CARTON	02/15/2017	
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	02/15/2017		

Labeler - Family First Pharmaceuticals, Inc. (832435809)

Establishment

Name	Address	ID/FEI	Business Operations
RNA Pharma, LLC		079103999	MANUFACTURE(51068-507)

Revised: 2/2017

Family First Pharmaceuticals, Inc.