## 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.

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#### 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 mediacal 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)** 

# ACCION BETT OF THE STRENGTH ARTHRITS

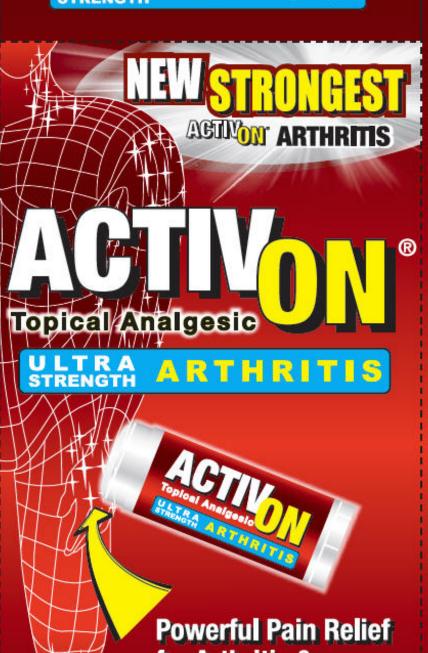
# ACTIVON Topical Analgesis

ULTRA ARTHRITIS

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

**DEEP PENETRATING PAIN RELIEF** 









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





## ACLLV **Topical Analgesic**

ULTRA STRENGTH ARTHRITIS

#### **Drug Facts**

#### **Active Ingredient**

Purpose

Menthol 5.138%.....

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,

- 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



#### 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
<b>Menthol, Unspecified Form</b> (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)		

l	Packaging			
ı	# Item Code	Package Description	<b>Marketing Start Date</b>	Marketing End Date
ı	1 NDC:51068-507-01	1 in 1 CARTON	02/15/2017	
ı	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.