#### RAPIDOL ARNICA- arnica montana gel Pharmadel LLC

Disclaimer: This homeopathic product has not been evaluated by the Food and Drug Administration for safety or efficacy. FDA is not aware of scientific evidence to support homeopathy as effective.

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**Rapidol Arnica Gel (Hwell)** 

**Drug Facts** 

## Active ingredient & Purposes

| Active Ingredient **    | Purposes*                        |
|-------------------------|----------------------------------|
| Arnica montana 1 X HPUS | Muscle aches and pains, bruises, |
|                         | swelling from injuries           |

The letters *"HPUS"* indicate that the components in this product are officially included in the Homeopathic Pharmacopoeia of the United States.

#### Uses

For the temporary relief of minor aches and pains of muscles and joints associated with

- simple backaches
- arthritis
- strains
- bruises
- sprains

## Warnings

# FOR EXTERNAL USE ONLY. Avoid contact with the eyes.

#### Do not use

- on wounds or damaged skin
- on a bandage tightly

## Stop use and consult a doctor

- condition worsens
- symptoms persist for more than 7 days or clear up and occur again within a few days

## if pregnant or breastfeeding,

consult a doctor before using this product.

# KEEP OUT OF THE REACH OF CHILDREN.

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

# Directions

- adults and children 2 years of age and older: apply to the affected area not more than 3 to 4 times daily.
- children under 2 years of age: do not use, consult a doctor

# Other information

- store in a cool dry place
- do not use if foil seal under cap is torn, broken, or missing

# Inactive ingredients

aloe vera leaf, capryloyl glycine, hexylene glycol, hydroxyethyl cellulose, phenoxyethanol, potassium sorbate, methyl salicylate, water.

## Questions or comments?

1-866-359-3478 (M-F) 9AM to 5 PM EST or www.pharmadel.com

# Distributed by:

PHARMADEL LLC. New Castle, DE, 19720

## Homeopathic Statements

\*These "USES" claims have not been evaluated by the Food and Drug Administration (FDA). They are based upon traditional homeopathic practices. \*\*C, K, CK, and X are homeopathic dilutions.

# Principal Display Panel



| Product Information      |                |                    |               |
|--------------------------|----------------|--------------------|---------------|
| Product Type             | HUMAN OTC DRUG | Item Code (Source) | NDC:55758-403 |
| Route of Administration  | TOPICAL        |                    |               |
|                          |                |                    |               |
| Active Ingredient/Active | Moiety         |                    |               |
| Ingr                     | edient Name    | Basis<br>Streng    | Strongth      |

| Ina     | active Ingre                      | dients                          |  |     |                         |     |                     |
|---------|-----------------------------------|---------------------------------|--|-----|-------------------------|-----|---------------------|
|         |                                   | In                              | gredient Name                                  |     |                         |     | Strength            |
| CA      | PRYLOYL GLYCI                     | NE (UNII: 8TY5YO42NJ            | ))   |     |                         |     |                     |
| 4L(     | OE VERA LEAF (                    | UNII: ZY81Z83H0X)               |  |     |                         |     |                     |
|         |                                   | <b>TE</b> (UNII: LAV5U5022Y     | <i>(</i> )                                     |     |                         |     |                     |
|         |                                   | <b>L</b> (UNII: HIE492ZZ3T)     |  |     |                         |     |                     |
|         |                                   | ATE (UNII: 1VPU26JZZ            | (4)  |     |                         |     |                     |
|         |                                   | <b>L</b> (UNII: KEH0A3F75J)     |  |     |                         |     |                     |
| IY      | DROXYETHYL C                      | ELLULOSE, UNSPEC                | IFIED (UNII: T4V6TWG28D                        | ))  |                         |     |                     |
|         |                                   |                                 |  |     |                         |     |                     |
|         | ickaging                          |                                 | <b>.</b>                                       |     | Marketing Start         | Маг | rketina End         |
| Pa<br># | ickaging<br>Item Code             | Package                         | e Description                                  | 1   | Marketing Start<br>Date | Mai | rketing End<br>Date |
| #       | Item Code                         |                                 | e <b>Description</b><br>e 0: Not a Combination |     | -                       | Maı |                     |
| #       | Item Code<br>NDC:55758-403-       | 57 g in 1 TUBE; Type            | -  |     | Date                    | Maı |                     |
| #<br>1  | Item Code<br>NDC:55758-403-<br>02 | 57 g in 1 TUBE; Type            | -  |     | Date                    | Maı | •                   |
| #       | Item Code<br>NDC:55758-403-<br>02 | 57 g in 1 TUBE; Type<br>Product | -  | 06/ | Date                    |     | •                   |

# Labeler - Pharmadel LLC (030129680)

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Pharmadel LLC