# COLD ZONE PAIN RELIEVING GEL- DG HEALTH- menthol gel Velocity Pharma LLC

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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#### Cold Zone Pain Relieving Gel- DG Health

#### **Active Ingredients**

Menthol USP 4%

#### **Purpose**

Cooling Pain Relief

#### Uses

Temporary relief from minor aches and pains of sore muscles and joints associated with: - arthritis - backache - strains - sprains

#### Warnings:

For external use only

#### Flammable:

Keep away from excessive heat or open flame.

#### Ask a doctor before use if you have:

sensitive skin

#### When using this product:

- Avoid contact with the eyes or mucous membranes
- Do not apply to wounds or damaged skin
- Do not use with other ointments, creams, sprays or liniments
- Do not apply to irritated skin or if excessive irritated develops
- Do not bandage
- Wash hands after use with cool water
- Do not use with heating pad or device

#### Stop use and ask a doctor if:

Condition worsens, or if symptoms persist for more than 7 days, or clear up and recur

#### If pregnant or breast-feeding:

Ask a health professional before use

#### Keep out of reach of children:

If accidentally ingested, get medical help or contact a Poison Control Center immediately

#### Directions:

- **Adults and children 2 years of age and older:** Rub a thin film over affected areas not more than 4 times daily; massage not necessary
- Children under 2 years of age: Consult physician

#### Other Information:

Store in a cool dry place with lid closed tightly

#### **Inactive Ingredients**

Purified water, Glycerin, Isopropyl alcohol, Iso propyl myristate, Carbopol usp, Aloevera gel, Arnica montana flower extract, Articum lappa root (burdock) extract, Boswellia carterii resin extract, Calendula officinalis extract, Camellia sinensis leaf extract, Iiex paraguariensis leaf extract, Melissa officinalis (lemon balm) leaf extract, Tocopheryl acetate, Triethanolamine, Colloidal silicon dioxide, Sorbic acid, Camphor, Brillient blue, Tartrazin supra

#### Cold Zone Pain Relieving Gel- DG Health



### COLD ZONE PAIN RELIEVING GEL- DG HEALTH

menthol gel

| Product Information     |                |                    |               |
|-------------------------|----------------|--------------------|---------------|
| Product Type            | HUMAN OTC DRUG | Item Code (Source) | NDC:76168-401 |
| Route of Administration | TOPICAL        |                    |               |

| Basis of Strength | Strength      |
|-------------------|---------------|
| MENTHOL           | 40 mg in 1 mL |
|                   |               |

| Inactive Ingredients                            |          |
|---|----------|
| Ingredient Name                                 | Strength |
| CAMPHOR (NATURAL) (UNII: N20 HL7Q941)           |          |
| BRILLIANT BLUE G (UNII: M1ZRX790SI)             |          |
| PETROLATUM (UNII: 4T6H12BN9U)                   |          |
| CARBO XYPO LYMETHYLENE (UNII: 0 A5MM307FC)      |          |
| ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)            |          |
| ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)        |          |
| ARCTIUM LAPPA WHO LE (UNII: 730 70 DU1LA)       |          |
| CALENDULA OFFICINALIS FLOWER (UNII: POM7O4Y7YD) |          |
| FRANKINCENSE (UNII: R9 XLF1R1WM)                |          |
| GREEN TEA LEAF (UNII: W2ZU1RY8B0)               |          |
| ILEX PARAGUARIENSIS LEAF (UNII: 1Q953B4O4F)     |          |
| MELISSA OFFICINALIS LEAF (UNII: 50 D2ZE9219)    |          |
| ISOPROPYL ALCOHOL (UNII: ND2M416302)            |          |
| ISOPROPYL MYRISTATE (UNII: 0 RE8 K4LNJS)        |          |
| ALPHATO COPHEROL ACETATE (UNII: 9E8X80D2L0)     |          |
| SILICON DIO XIDE (UNII: ETJ7Z6 XBU4)            |          |
| TROLAMINE (UNII: 9O3K93S3TK)                    |          |
| SORBIC ACID (UNII: X045WJ989B)                  |          |
| WATER (UNII: 059QF0KO0R)                        |          |
| FD&C BLUE NO. 1 (UNII: H3R47K3TBD)              |          |

| l | Packaging        |   |                             |                    |
|---|------------------|---|-----------------------------|--------------------|
| l | # Item Code      | Package Description                                       | <b>Marketing Start Date</b> | Marketing End Date |
| l | 1 NDC:76168-401- | 1 59.14 mL in 1 BOTTLE; Type 0: Not a Combination Product | 0 1/22/20 18                |                    |

| Marketing Information   |  |                      |                    |
|-------------------------|--|----------------------|--------------------|
| Marketing Category      | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
| OTC monograph not final | part348                                  | 0 1/22/20 18         |                    |
|                         |  |                      |                    |

## Labeler - Velocity Pharma LLC (962198409)

### Registrant - Velocity Pharma LLC (962198409)

| Establishment        |         |           |                        |  |
|----------------------|---------|-----------|------------------------|--|
| Name                 | Address | ID/FEI    | Business Operations    |  |
| Yash Pharmaceuticals |         | 871409551 | manufacture(76168-401) |  |

Revised: 1/2018 Velocity Pharma LLC