

**ZYLOTROL MAXIMUM PAIN PATCH- lidocaine 4%, menthol 1% patch**  
**Whitestone Products LLC**

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**Zylotrol for Pain Control Lidocaine 4% and Menthol 1%**

Lidocaine 4%

Menthol 1%

Topical Analgesic

**For external use only** not intended for ingestion.

- in large quantities, particularly over raw surfaces, or blistered areas.
- Avoid contact with eyes.
- condition worsens, or if symptoms persist for more than 7 days or clear up and occur again within a few days.

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

ask a health professional before use.

**DIRECTIONS**

- Adults and children 2 years of age and older: Apply to affected area not more than 3 to 4 times a daily.
- Children under 2 years of age: consult a doctor.
- Store at 20-25oC (68-77oF) and protect from moisture.

polyacrylamide, vinol, sodium polyacrylate; acrylate polymerization; purified water.

**Questions?**

(310) 320-0100

**USES**

For the temporary relief of pain.

5.2244 w x 6.7461h x 1.75"deep



## ZYLOTROL MAXIMUM PAIN PATCH

lidocaine 4%, menthol 1% patch

### Product Information

|                                |                |                           |               |
|--------------------------------|----------------|---------------------------|---------------|
| <b>Product Type</b>            | HUMAN OTC DRUG | <b>Item Code (Source)</b> | NDC:81902-101 |
| <b>Route of Administration</b> | TOPICAL        |                           |               |

### Active Ingredient/Active Moiety

| Ingredient Name   | Basis of Strength         | Strength        |
|---|---------------------------|-----------------|
| <b>LIDOCAINE</b> (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)                                 | LIDOCAINE                 | 4 g<br>in 100 g |
| <b>MENTHOL, UNSPECIFIED FORM</b> (UNII: L7T10EIP3A) (MENTHOL, UNSPECIFIED FORM - UNII:L7T10EIP3A) | MENTHOL, UNSPECIFIED FORM | 1 g<br>in 100 g |

### Inactive Ingredients

| Ingredient Name                 | Strength |
|---------------------------------|----------|
| <b>WATER</b> (UNII: 059QF0KO0R) |          |

|  |  |
|--|--|
| <b>ACRYLATES CROSSPOLYMER-6</b> (UNII: 4GXD0Q3OS3)         |  |
| <b>POLYACRYLAMIDE (10000 MW)</b> (UNII: E2KR9C9V2I)        |  |
| <b>SODIUM POLYACRYLATE (2500000 MW)</b> (UNII: 05I15JNI2J) |  |
| <b>ETHENYL</b> (UNII: PQ2K3G3591)                          |  |

### Packaging

| # | Item Code        | Package Description                               | Marketing Start Date | Marketing End Date |
|---|------------------|---|----------------------|--------------------|
| 1 | NDC:81902-101-15 | 15 in 1 BOX                                       | 07/30/2021           |                    |
| 1 |                  | 4 g in 1 PATCH; Type 0: Not a Combination Product |                      |                    |

### Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|--------------------|--|----------------------|--------------------|
| OTC Monograph Drug | 505G(a)(3)                               | 07/30/2021           |                    |

**Labeler** - Whitestone Products LLC (118064415)

Revised: 10/2023

Whitestone Products LLC