

**ODYNIA-UULTRA PATCH ULTRA PATCH- lidocaine, capsicum patch**  
**Ursh Pharmaceutical 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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**Drug Facts**

**Active Ingredient**

Lidocaine 4 %

Capsicum 0.03 %

**Purpose**

Topical Analgesic

External Analgesic

**Uses** temporarily relieves minor aches and pains associated with: -arthritis -simple backache --  
bursitis --tendonitis

--muscle strains --sprains --bruises --cramps

**Warnings**

**For external use only**

**When using this product** -use only as directed -do not bandage tightly or use with a heating pad

-avoid contact with eyes and mucus membranes -do not apply to wounds or damaged, broken or irritated skin

**Stop use and ask a doctor if** -condition worsens -redness is present -irritation develops

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

**If pregnant or breast feeding**, ask a health professional before use

**Keep out of reach of children.** If swallowed, get medical help or contact a Poison Control Center right away

**Directions**

**adults and children over 12 years:**

-remove backing from patch by firmly grasping both ends and gently pulling until backing separates in middle

-carefully remove smaller portion of backing from patch and apply exposed portion of patch to affected area

-once exposed portion of patch is positioned, carefully remove remaining backing to completely apply patch to affected area

-wear one Icy Hot Patch up to 8 hours

-repeat as necessary, but no more than 3 times daily

**children under 12 years or younger:** ask a doctor

**Inactive ingredients** aluminum hydroxide, carmellose sodium, glycerin, isopropyl myristate, methyl

acrylate / 2 -ethylhexyl acrylate copolymer, nonoxynol-30, polyacrylic acid, polysorbate 80, sodium polyacrylate, sorbitan

sesquioleate, starch / acrylic acid graft copolymer sodium salt, talc, tartaric acid, titanium dioxide, water

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| <b>Active ingredient</b>   |
| Lidocaine 4%<br>Capsicum 0.03%   |
| <b>Uses</b> temporarily relieves minor aches and pains associated with:<br>• arthritis<br>• simple backache<br>• muscle strains<br>• sprains<br>• bruises<br>• cramps  |
| <b>Warnings</b><br>For external use only   |
| When using this product:<br>• use only as directed<br>• do not bandage tightly or use with a heating pad<br>• avoid contact with eyes and mucous membranes<br>• do not apply to wounds or damaged, broken or irritated skin  |
| Stop use and ask a doctor if:<br>• condition worsens<br>• redness is present<br>• irritation develops<br>• symptoms persist for more than 7 days or clear up and occur again within a few days   |
| If pregnant or breast-feeding, ask a health professional before use.<br>Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.   |
| <b>Directions</b>  |
| adults and children over 12 years:<br>• remove backing from patch by firmly grasping both ends and gently pulling until backing separates in middle<br>• carefully remove smaller portion of backing from patch and apply exposed portion of patch to affected area<br>• once exposed portion of patch is positioned, carefully remove remaining backing to completely apply patch to affected area<br>• wear one Ice Hot Patch up to 8 hours<br>• repeat as necessary, but no more than 3 times daily |
| children 12 years or younger: ask a doctor   |
| <b>Inactive ingredients</b> aluminum hydroxide, carmellose sodium, glycerin, isopropyl myristate, methyl methacrylate/2-ethylhexyl acrylate copolymer, nonoxynol-30, polyacrylic acid, polysorbate 80, sodium polyacrylate, starch/acrylic acid graft copolymer sodium salt, talc, tartaric acid, titanium dioxide, water  |

## ODYNIA-UULTRA PATCH ULTRA PATCH

lidocaine, capsicum patch

### Product Information

|                         |                |                    |               |
|-------------------------|----------------|--------------------|---------------|
| Product Type            | HUMAN OTC DRUG | Item Code (Source) | NDC:69647-002 |
| Route of Administration | TOPICAL        |                    |               |

**Active Ingredient/Active Moiety**

| Ingredient Name  | Basis of Strength | Strength       |
|--|-------------------|----------------|
| LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987) | LIDOCAINE         | 4 g in 100 g   |
| CAPSICUM (UNII: 00UK7646FG) (CAPSICUM - UNII:00UK7646FG)   | CAPSICUM          | .03 g in 100 g |

**Inactive Ingredients**

| Ingredient Name  | Strength |
|--|----------|
| ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)                              |          |
| CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED FORM (UNII: K679OBS311) |          |
| GLYCERIN (UNII: PDC6A3C0OX)  |          |
| ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)                             |          |
| METHYL ACRYLATE (UNII: WC487PR91H)                                 |          |
| 2-ETHYLHEXYL ACRYLATE (UNII: HR49R9S6XG)                           |          |
| NONOXYNOL-30 (UNII: JJX07DG188)                                    |          |
| POLYACRYLIC ACID (250000 MW) (UNII: 9G2MAD7J6W)                    |          |
| POLYSORBATE 80 (UNII: 6OZP39ZG8H)                                  |          |
| SODIUM POLYACRYLATE (2500000 MW) (UNII: 05H5JN2J)                  |          |
| SORBITAN SESQUIOLEATE (UNII: 0W8RRI5W5A)                           |          |
| ACRYLIC ACID (UNII: J94PBK7X8S)                                    |          |
| SODIUM (UNII: 9NEZ333N27)  |          |
| SODIUM CHLORIDE (UNII: 451W47IQ8X)                                 |          |
| TALC (UNII: 7SEV7J4R1U)  |          |
| TARTARIC ACID (UNII: W4888119H)                                    |          |
| TITANIUM DIOXIDE (UNII: 15FIX9V2JP)                                |          |
| WATER (UNII: 059QF0K00R)   |          |

**Packaging**

| # | Item Code        | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|---------------------|----------------------|--------------------|
| 1 | NDC:69647-002-01 | 15 in 1 BOX         |                      |                    |
| 1 |                  | 10 g in 1 PATCH     |                      |                    |

**Marketing Information**

| Marketing Category      | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|-------------------------|--|----------------------|--------------------|
| OTC monograph not final | part348                                  | 03/12/2015           |                    |

**Labeler** - Ursh Pharmaceutical Inc. (079715344)

Revised: 3/2015

Ursh Pharmaceutical Inc.