

**IMPROVUE LUBRICANT- hypromellose 2208 (15000 mpa.s) solution/ drops**  
**Oculus Surgical, Inc.**

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**ImproVue®**  
**Ophthalmic Lubricant Drops**

***Drug Facts***

**Active Ingredients**

Hypromellose (1.7%)

**Purpose**

Ophthalmic Lubricant

**Indications**

- For the temporary relief of burning and irritation due to dryness of the eye.
- For use as a lubricant to prevent further irritation or to relieve dryness of the eye.

**Warnings**

- For external use only.
- To avoid contamination do not touch tip of container or applicator to any surface.
- Do not reuse. Once opened, discard.

**Stop use and ask a doctor if**

- If you experience eye pain, changes in vision, continued redness or irritation of the eye, or if the condition worsens or persists for more than 72 hours, discontinue use and consult a doctor.
- If solution changes color or becomes cloudy, do not use.

**Keep out of reach of children**

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

**Directions**

- Remove cap from syringe and screw on applicator tip.
- Instill 1 or 2 drops in the affected eye(s) as needed.

**Other Information**

- Do not use if pouch is damaged or has been previously opened.
- Protect from freezing
- Store at or below 25°C (77°F)

## **Inactive Ingredients**

Calcium chloride dihydrate, hydrochloric acid<sup>1</sup>, magnesium chloride hexahydrate, potassium chloride, purified water, sodium acetate trihydrate, sodium chloride, sodium citrate dihydrate, and sodium hydroxide<sup>1</sup>

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<sup>1</sup> May contain one or more of these ingredients for pH adjustment

## **Questions or Comments?**

(855) 734-2466 or (772) 236-2622 or log onto [www.oculussurgical.com](http://www.oculussurgical.com)

## **PRINCIPAL DISPLAY PANEL - 6 Pouch Carton**

### **ImproVue®**

#### **Ophthalmic Lubricant Drops**

- Sterile
- Preservative-free
- Single use, disposable container

Contains:

6 Pouches, each Pouch containing  
1 Single Use Syringe, 0.07 fl oz (2 ml)  
1 Single Use Applicator Tip

**OCULUS®**

**SURGICAL**

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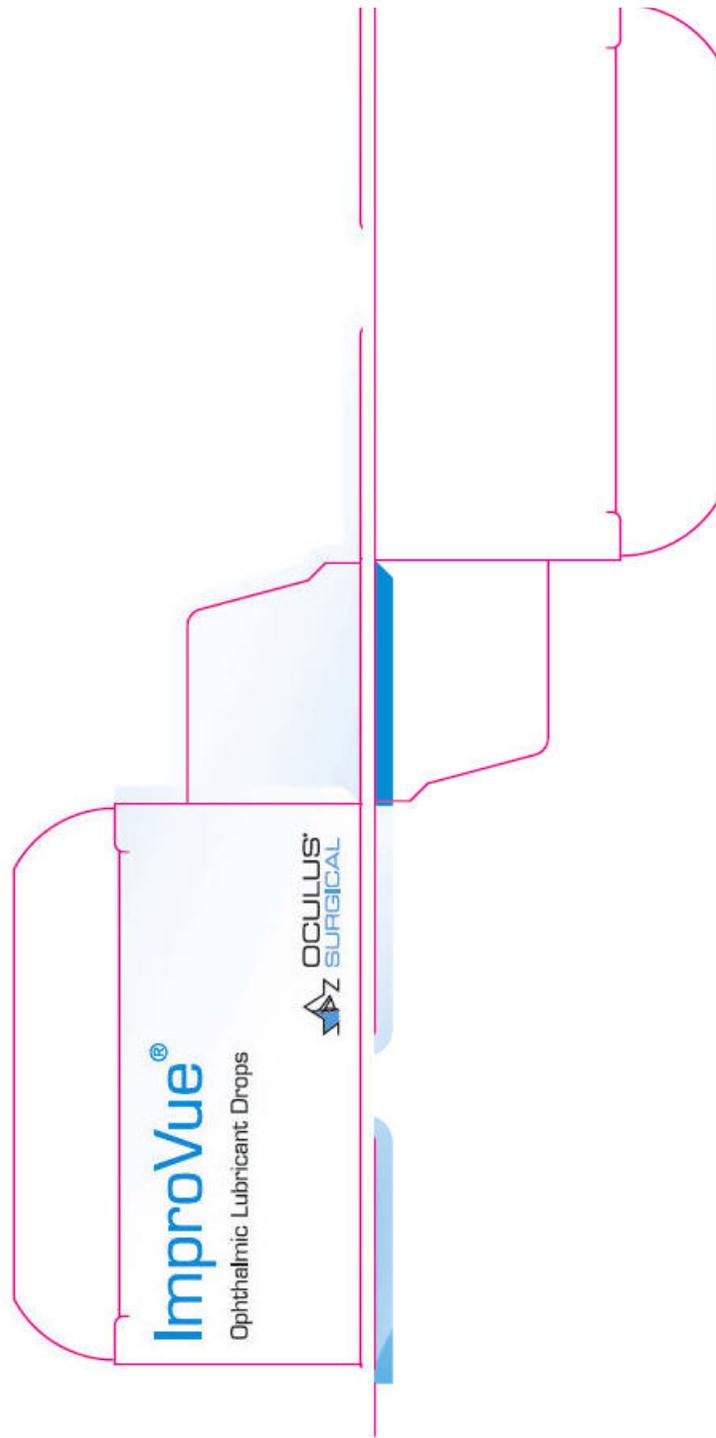
|   |   |
|---|---|
| <b>Drug Facts</b>   | <b>Purpose</b><br>Ophthalmic Lubricant  |
| <b>Active Ingredients</b><br>Hydroxyethylcellulose (1.7%) |   |
| <b>Indications</b>  | For the temporary relief of burning and irritation due to dryness of the eye.<br>For use as a lubricant to prevent further irritation or to relieve dryness of the eye.   |
| <b>Warnings</b>   | For external use only.<br>To avoid contamination do not touch tip of container or applicator to any surface.<br>Do not reuse. Once opened, discard.   |
| <b>Stop use and ask a doctor if</b>                       | If you experience eye pain, changes in vision, continued redness or irritation of the eye, or if the condition worsens or persists for more than 72 hours, discontinue use and consult a doctor.<br>If solution changes color or becomes cloudy, do not use.                        |
| <b>Keep out of reach of children</b>                      | If swallowed, get medical help or contact a Poison Control Center right away.   |
| <b>Directions</b>   | Remove cap from syringe and screw on applicator tip.<br>Instill 1 or 2 drops in the affected eye(s) as needed.  |
| <b>Other Information</b>                                  | Do not use if pouch is damaged or has been previously opened.<br>Protect from freezing.<br>Store at or below 25°C (77°F)  |
| <b>Inactive Ingredients</b>                               | Calcium chloride dihydrate, hydrochloric acid*, magnesium chloride hexahydrate, potassium chloride, purified water, sodium acetate trihydrate, sodium chloride, sodium citrate dihydrate, and sodium hydroxide*<br>* May contain one or more of these ingredients for pH adjustment |
| <b>Questions or Comments?</b>                             | (855) 734-2468 or (772) 236-2622 or log onto www.oculussurgical.com   |

Manufactured for:  
 OCULUS<sup>®</sup>  
 SURGICAL  
 Port St. Lucie, FL 34986 USA



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## IMPROVUE LUBRICANT

hypromellose 2208 (15000 mpa.s) solution/ drops

### Product Information

|                                |                |                           |                |
|--------------------------------|----------------|---------------------------|----------------|
| <b>Product Type</b>            | HUMAN OTC DRUG | <b>Item Code (Source)</b> | NDC:62144-5510 |
| <b>Route of Administration</b> | OPHTHALMIC     |                           |                |

### Active Ingredient/Active Moiety

| Ingredient Name   | Basis of Strength               | Strength      |
|---|---------------------------------|---------------|
| <b>HYPROMELLOSE 2208 (15000 MPA.S)</b> (UNII: Z78RG6M2N2) (HYPROMELLOSE 2208 (15000 MPA.S) - UNII:Z78RG6M2N2) | HYPROMELLOSE 2208 (15000 MPA.S) | 17 mg in 1 mL |

### Inactive Ingredients

| Ingredient Name                                       | Strength |
|---|----------|
| <b>CALCIUM CHLORIDE</b> (UNII: M4I0D6VV5M)            |          |
| <b>HYDROCHLORIC ACID</b> (UNII: QTT17582CB)           |          |
| <b>MAGNESIUM CHLORIDE</b> (UNII: 02F3473H90)          |          |
| <b>POTASSIUM CHLORIDE</b> (UNII: 660YQ98I10)          |          |
| <b>WATER</b> (UNII: 059QF0KO0R)                       |          |
| <b>SODIUM ACETATE</b> (UNII: 4550K0SC9B)              |          |
| <b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)             |          |
| <b>TRISODIUM CITRATE DIHYDRATE</b> (UNII: B22547B95K) |          |
| <b>SODIUM HYDROXIDE</b> (UNII: 55X04QC32I)            |          |

### Packaging

| # | Item Code        | Package Description  | Marketing Start Date | Marketing End Date |
|---|------------------|--|----------------------|--------------------|
| 1 | NDC:62144-5510-5 | 6 in 1 CARTON  | 06/25/2014           |                    |
| 1 |                  | 1 in 1 POUCH   |                      |                    |
| 1 |                  | 2 mL in 1 SYRINGE, PLASTIC; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.) |                      |                    |

### Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|--------------------|--|----------------------|--------------------|
| OTC MONOGRAPH DRUG | M018                                     | 06/25/2014           |                    |

**Labeler** - Oculus Surgical, Inc. (015409582)

### Establishment

| Name                | Address | ID/FEI    | Business Operations     |
|---------------------|---------|-----------|-------------------------|
| OASIS Medical, Inc. |         | 194121018 | MANUFACTURE(62144-5510) |

### Establishment

| Name                | Address | ID/FEI    | Business Operations |
|---------------------|---------|-----------|---------------------|
| OASIS Medical, Inc. |         | 024362989 | PACK(62144-5510)    |