Active ingredient
Levonorgestrel, USP 1.5 mg

Purpose
Emergency contraceptive

Use
for women to reduce chance of pregnancy after unprotected sex (if a contraceptive failed or if you did not use birth control)

Warnings

Allergy alert
Do not use if you have ever had an allergic reaction to levonorgestrel

Sexually transmitted diseases (STDs) alert
This product does not protect against HIV/AIDS or other STDs.

Do not use
• if you are already pregnant (because it will not work)
• for regular birth control

Ask a doctor or pharmacist before use if you are taking efavirenz (HIV medication) or rifampin (tuberculosis treatment) or medication for seizures (epilepsy). These medications may reduce the effectiveness of levonorgestrel.

When using this product you may have
• menstrual changes
• tiredness
• dizziness
• nausea
• headache
• breast pain
• lower stomach (abdominal) pain
• vomiting

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away (1-800-222-1222).
Directions

• take as soon as possible within 72 hours (3 days) after unprotected sex. The sooner you take it, the better it will work.
• if you vomit within 2 hours after taking the medication, call a healthcare professional to find out if you should repeat the dose

Other information

• read the instructions, warnings, and enclosed product leaflet before use
• this product works mainly by preventing ovulation (egg release). It may also prevent fertilization of a released egg (joining of sperm and egg) or attachment of a fertilized egg to the uterus (implantation).
• do not use if carton is open or tear strip is removed or blister seal is broken or missing
• store at 20° to 25°C (68° to 77°F)

Inactive ingredients
colloidal silicon dioxide, corn starch, hypromellose, lactose monohydrate, magnesium stearate, talc

Questions or comments?
For more information, call toll free 1-800-818-4555 weekdays

Distributed by:
Ohm Laboratories Inc.
New Brunswick, NJ 08901

PRINCIPAL DISPLAY PANEL - 1.5 mg Tablet Blister Pack Carton

†Compare To
the active ingredient of
Plan B One-Step®

NDC 62756-720-60

See New Warning

My
Choice™

Levonorgestrel Tablet 1.5 mg
Emergency Contraceptive

• Reduces the chance of pregnancy after unprotected sex
• Not for regular birth control
• The sooner you take it, the more effective it will be
• Take as soon as possible within 72 hours (3 days) after unprotected sex
• Will not harm an existing pregnancy

1 Tablet Levonorgestrel 1.5 mg
One Tablet. One Step.
My Choice
Levonorgestrel Tablet 1.5 mg

Emergency Contraceptive

- Reduces the chance of pregnancy after unprotected sex
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1 Tablet Levonorgestrel 1.5 mg

One Tablet. One Step.

Drug Facts

Active ingredient: Levonorgestrel, USP 1.5 mg

Purpose: Emergency contraceptive

Uses: For women to reduce the chance of pregnancy after unprotected sex (if a contraceptive failed or if you did not use birth control)

Warnings:
- Allergy alert: Do not use if you have ever had an allergic reaction to levonorgestrel
- Sexually transmitted diseases (STDs) alert: This product does not protect against
### MY CHOICE TM
levonorgestrel tablet

#### Product Information

<table>
<thead>
<tr>
<th>Product Type</th>
<th>HUMAN OTC DRUG</th>
<th>Item Code (Source)</th>
<th>NDC: 62756-720</th>
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<tbody>
<tr>
<td>Route of Administration</td>
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#### Active Ingredient/Active Moiety

<table>
<thead>
<tr>
<th>Ingredient Name</th>
<th>Basis of Strength</th>
<th>Strength</th>
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<tbody>
<tr>
<td>LEVONORGESTREL</td>
<td>(UNII: 5W7SIA7YZW)</td>
<td>LEVONORGESTREL</td>
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<tr>
<td>(LEVONORGESTREL - UNII: 5W7SIA7YZW)</td>
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#### Inactive Ingredients

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<tr>
<th>Ingredient Name</th>
<th>Strength</th>
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<tbody>
<tr>
<td>SILICON DIOXIDE (UNII: E117Z6XBU4)</td>
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<tr>
<td>STARCH, CORN (UNII: 0B232NY3SJ)</td>
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<tr>
<td>HYDROXYMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)</td>
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<td>LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)</td>
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<td>MAGNESIUM STEARATE (UNII: 70097M6I30)</td>
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<td>TALC (UNII: 7SEV7J4RIU)</td>
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### Product Characteristics

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<tr>
<td>Shape</td>
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<td>Flavor</td>
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### Packaging

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<th>Marketing End Date</th>
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<tr>
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<td>NDC:62756-720-60</td>
<td>1 in 1 CARTON</td>
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<td>1</td>
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<td>1 in 1 BLISTER PACK; Type 0: Not a Combination Product</td>
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### Marketing Information

<table>
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<tr>
<th>Marketing Category</th>
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<th>Marketing End Date</th>
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### Labeler

SUN PHARMACEUTICAL INDUSTRIES, INC. (146974886)

### Establishment

<table>
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<tr>
<th>Name</th>
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<tr>
<td>Sun Pharmaceutical Industries Limited</td>
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<td>725959238</td>
<td>ANALYSIS(62756-720), MANUFACTURE(62756-720)</td>
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Revised: 10/2018