

LEVONORGESTREL- levonorgestrel tablet
Ohm Laboratories Inc.

Levonorgestrel

Drug Facts

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
- breast pain
- nausea
- headache
- vomiting
- lower stomach (abdominal) pain
- dizziness

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

NDC 51660-723-11

My After Plan®

Levonorgestrel

1.5 mg Tablet

EMERGENCY

CONTRACEPTIVE

- Reduces the chance of pregnancy after unprotected sex
- Not for regular birth control

One Step

Contains 1 Tablet

Levonorgestrel 1.5 mg



52151 21

NDC 51660-723-11



My After Plan®

Levonorgestrel
1.5 mg Tablet

EMERGENCY CONTRACEPTIVE

- ◆ Reduces the chance of pregnancy after unprotected sex
- ◆ Not for regular birth control



One Step

Contains 1 Tablet
Levonorgestrel 1.5 mg

Lot No.

Expiration Date:

NON VARNISH

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MY AFTER PLAN is a registered trademark of Sun Pharmaceutical Industries Limited.

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New Brunswick, NJ 08901

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LEVONORGESTREL

levonorgestrel tablet

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:51660-723
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
Levonorgestrel (UNII: 5W7SIA7YZW) (Levonorgestrel - UNII:5W7SIA7YZW)	Levonorgestrel	1.5 mg

Inactive Ingredients	
Ingredient Name	Strength
silicon dioxide (UNII: ETJ7Z6XBU4)	
starch, corn (UNII: O8232NY3SJ)	
hypromellose, unspecified (UNII: 3NXW29V3WO)	
lactose monohydrate (UNII: EWQ57Q8I5X)	
magnesium stearate (UNII: 70097M6I30)	
talc (UNII: 7SEV7J4RIU)	

Product Characteristics			
Color	WHITE	Score	no score
Shape	ROUND	Size	8mm
Flavor		Imprint Code	
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51660-723-11	1 in 1 CARTON	11/01/2017	
1		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA202635	11/01/2017	

Labeler - Ohm Laboratories Inc. (184769029)

Establishment

Name	Address	ID/FEI	Business Operations
Sun Pharmaceutical Industries Limited		725959238	MANUFACTURE(51660-723)