MY CHOICE TM- levonorgestrel tablet Proficient Rx LP

My Choice™

Drug Facts

Active ingredient

Levonorgestrel, USP 1.5 mg

Purpose

Emergency contraceptive

Use

for women to reduce chance of pregnancy af ter unprotected sex (if a contraceptive failed or if you did not use bir th 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 bir th 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 fer tilization of a released egg (joining of sperm and egg) or at tachment of a fer tilized 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

Relabeled by:

Proficient Rx LP

Thousand Oaks, CA 91320

PRINCIPAL DISPLAY PANEL - 1.5 mg Tablet Blister Pack Carton

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

NDC 71205-120-01

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.





NDC 71205-120-01

Relabeled By: Proficient Rx LP Thousand Oaks, CA 91320

My Choice (Levonorgestrel) 1.5mg #01 Tablets SN# MASTER Lot #:00000 Exp:00/00/00 NDC 71205-120-01

My Choice (Levonorgestrel) 1.5mg #01 Tablets SN# MASTER Lot #:00000 Exp:00/00/00 NDC 71205-120-01

My Choice (Levonorgestrel) 1.5mg #01 Tablets SN# MASTER Lot #:00000 Exp:00/00/00 NDC 71205-120-01



GTIN: 00371205120016 SN# MASTER Exp. 00/00/00 Lot #:00000

My Choice (Levonorgestrel) 1.5mg

#01

Tablets

Each tablet contains: Levonorgestrel, USP 1.5 mg Emergency contraceptive

White (white to off-white), round, unscored with imprint code 718

Product ID: SM012001

Dist. By: Ohm Laboratories Inc. New Brunswick, NJ 08901 MADE IN INDIA Store at 20° to 25°C (68° to 77°F)

Keep medication out of the reach of children

MY CHOICE TM

levonorgestrel tablet

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:71205-120(NDC:62756-720)

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: 7SEV7J4R1U)	

Product Characteristics			
Color	WHITE (white to off-white)	Score	no score
Shape	ROUND (circular)	Size	8mm
Flavor		Imprint Code	718
Contains			

F	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:71205-120- 01	1 in 1 CARTON	09/04/2018		
1		1 in 1 BLISTER PACK; Type 0: Not a Combination Product			

Marketing Information			
Marketing Application Number or Monograph Category Citation		Marketing Start Date	Marketing End Date
ANDA	ANDA202635	04/01/2018	

Labeler - Proficient Rx LP (079196022)

Establishment				
Name	Address	ID/FEI	Business Operations	
Proficient Rx LP		079196022	RELABEL(71205-120)	

Revised: 9/2022 Proficient Rx LP