**TAKE ACTION- levonorgestrel tablet**
**Foundation Consumer Healthcare LLC**

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**Take Action®**

***Drug Facts***

**Active ingredient**

Levonorgestrel 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.

**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**
- **do not use if carton is open or tear strip is removed or blister seal is broken or missing**
- store at 20-25°C (68-77°F)

**Inactive ingredients**

colloidal silicon dioxide, corn starch, lactose monohydrate, magnesium stearate, potato starch, talc

**Questions?**

Call **1-888-919-0780**

Manufactured by:
Gedeon Richter, Ltd., Budapest, Hungary

**PRINCIPAL DISPLAY PANEL - 1.5 mg Tablet Blister Pack Box**

NDC 69536-200-88

Emergency Contraceptive

This item is
electronically
protected

take
action®
LEVONORGESTREL 1.5 mg

Reduces chance of pregnancy
after unprotected sex.

NOT FOR REGULAR BIRTH CONTROL

One Tablet.
One Dose.

1 Tablet
Levonorgestrel

1.5 mg

take action®
LEVONORGESTREL 1.5 mg

Emergency Contraceptive
One Tablet. One Dose.

This item is electronically protected

NDC 69536-200-88

Emergency Contraceptive

TEAR HERE TO OPEN

# take action®

## LEVONORGESTREL 1.5 mg

Reduces chance of pregnancy after unprotected sex.

**NOT FOR REGULAR BIRTH CONTROL**

**One Tablet.
One Dose.**

1 Tablet
Levonorgestrel
1.5 mg

133339

133339 Rev. 11/2022

Emergency Contraceptive

take action®

LEVONORGESTREL 1.5 mg

- 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

## Drug Facts

**Active ingredient** — **Purpose**

Levonorgestrel 1.5 mg ..... 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)

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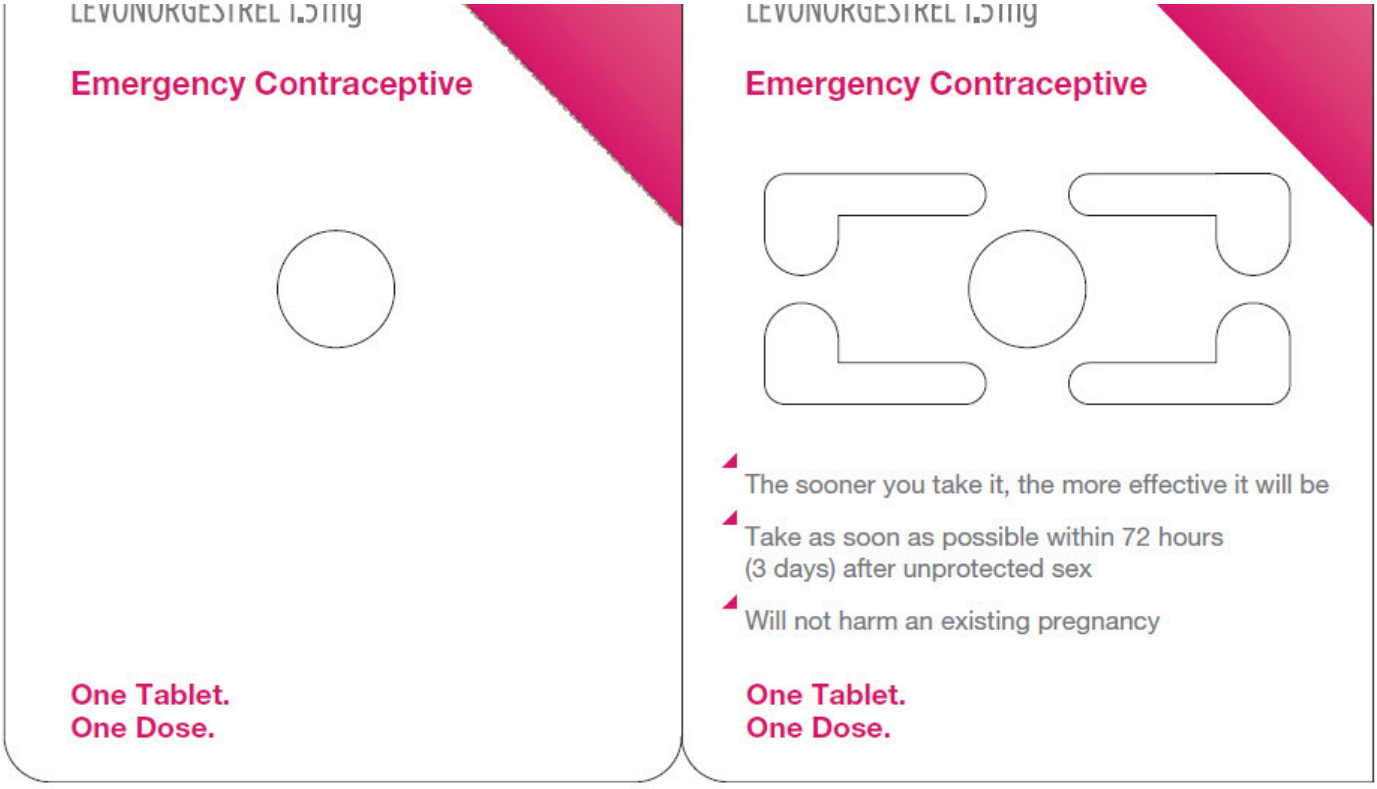
take action®

LEVONORGESTREL 1.5 mg

take action®

LEVONORGESTREL 1.5 mg




# TAKE ACTION

levonorgestrel tablet

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:69536-200 |
| **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, POTATO** (UNII: 8I089SAH3T) | |
| **MAGNESIUM STEARATE** (UNII: 70097M6I30) | |
| **TALC** (UNII: 7SEV7J4R1U) | |
| **STARCH, CORN** (UNII: O8232NY3SJ) | |
| **LACTOSE MONOHYDRATE** (UNII: EWQ57Q8I5X) | |

| Product Characteristics | | | |
|---|---|---|---|
| **Color** | WHITE | **Score** | no score |
| **Shape** | ROUND | **Size** | 8mm |

| | | | |
|---|---|---|---|
| **Flavor** | | **Imprint Code** | G00 |
| **Contains** | | | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:69536-200-88 | 1 in 1 BOX, UNIT-DOSE | 05/10/2018 | |
| **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 |
|---|---|---|---|
| NDA authorized generic | NDA021998 | 05/10/2018 | |

## Labeler - Foundation Consumer Healthcare LLC (079675882)

Revised: 1/2023 Foundation Consumer Healthcare LLC