

TAKE ACTION- levonorgestrel tablet
Foundation Consumer Healthcare LLC

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**
- 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 20-25°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

**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

NDC 69536-200-88

Emergency Contraceptive

TEAR HERE TO OPEN

take
action[®]
LEVONORGESTREL 1.5mg

take
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1 Tablet
Levonorgestrel
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799-30-101179

Rev. 03/2018

LEVONORGESTREL 1.5 mg
take
 action®

Emergency Contraceptive
 One Tablet, One Dose

**Emergency
 Contraceptive**

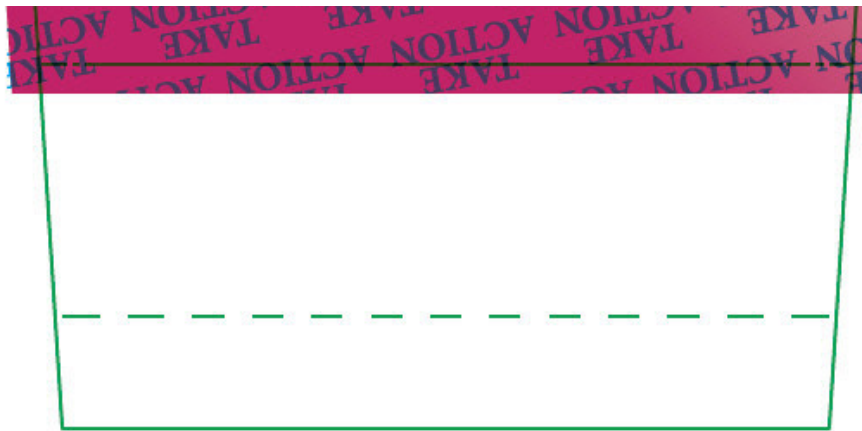
take
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 LEVONORGESTREL 1.5 mg

- ▲ 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)	
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	
<ul style="list-style-type: none"> ■ if you are already pregnant (because it will not work) ■ for regular birth control 	
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Inactive ingredients colloidal silicon dioxide, corn starch, lactose monohydrate, magnesium stearate, potato starch, talc	
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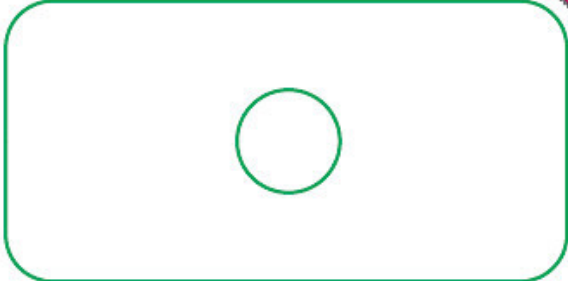


Manufactured by
 Redon Pharma Ltd., Budapest, Hungary



**take
action[®]**
 LEVONORGESTREL 1.5mg

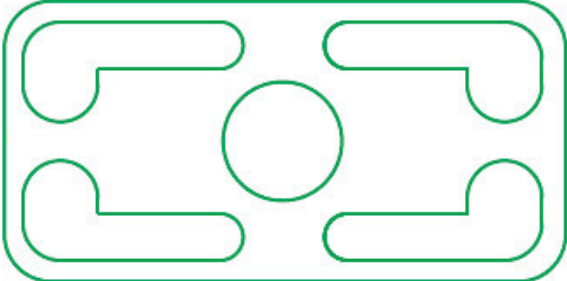
Emergency Contraceptive



**One Tablet.
One Dose.**

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 LEVONORGESTREL 1.5mg

Emergency Contraceptive



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One Dose.**

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: 10/2019

Foundation Consumer Healthcare LLC