

ELLA- ulipristal acetate tablet

HRA PHARMA AMERICA, INC.

HIGHLIGHTS OF PRESCRIBING INFORMATION

These highlights do not include all the information needed to use ella safely and effectively. See full prescribing information for ella.

ELLA (ulipristal acetate) tablet, for oral use
Initial U.S. Approval: 2010

RECENT MAJOR CHANGES

- | | |
|--|---------|
| • Dosage and Administration (2.1, 2.2, 2.3) | 06/2021 |
| • Warnings and Precautions (5.5) | 06/2021 |

INDICATIONS AND USAGE

Ella is a progesterone agonist/antagonist emergency contraceptive indicated for prevention of pregnancy following unprotected intercourse or a known or suspected contraceptive failure. **Ell**a is not intended for routine use as a contraceptive. (1)

DOSAGE AND ADMINISTRATION

- Take one tablet orally as soon as possible, within 120 hours (5 days) after unprotected intercourse or a known or suspected contraceptive failure. Take with or without food. Take at any time during the menstrual cycle. (2.1)
- After **ella** use, initiate or resume hormonal contraception no sooner than 5 days after the intake of **ella** and use a reliable barrier method until the next menstrual period. (2.2)
- If vomiting occurs within 3 hours of taking **ella**, consider repeating the dose. (2.3)

DOSAGE FORMS AND STRENGTHS

1. 30 mg tablet (3)

CONTRAINDICATIONS

- Known or suspected pregnancy (4)

WARNINGS AND PRECAUTIONS

- Existing Pregnancy: **ella** is not indicated for termination of an existing pregnancy. (5.1)
- Ectopic pregnancy: Evaluate women who become pregnant or complain of lower abdominal pain after taking **ella** for ectopic pregnancy. (5.2)
- Fertility Following Use: Rapid return of fertility is likely. Subsequent acts of intercourse should be protected by a reliable barrier method of contraception until the next menstrual period. (5.5)
- Effect on Menstrual Cycle: **ella** may alter the next expected menses. If menses is delayed beyond 1 week, rule out pregnancy. (5.6)
- **Ell**a does not protect against STI/HIV. (5.7)

ADVERSE REACTIONS

The most common adverse reactions ($\geq 5\%$) in the clinical trials were headache (18%), abdominal pain (12%), nausea (12%), dysmenorrhea (9%), fatigue (6%) and dizziness (5%). (6) To report SUSPECTED ADVERSE REACTIONS, contact HRA Pharma America Inc., at 844-994-0329 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

DRUG INTERACTIONS

- Drugs or herbal products that induce CYP3A4 decrease the effectiveness of **ella**. (7.1)
- Initiation of progestin-containing contraceptives may impair the ability of **ella** to delay ovulation. (7.1)

See 17 for PATIENT COUNSELING INFORMATION and FDA-approved patient labeling.

Revised: 1/2025

FULL PRESCRIBING INFORMATION: CONTENTS*

1 INDICATIONS AND USAGE

2 DOSAGE AND ADMINISTRATION

- 2.1 Recommended Dosage and Administration
- 2.2 Recommendations Regarding Use with Hormonal Contraception
- 2.3 Recommendation in Case of Gastrointestinal Disturbances

3 DOSAGE FORMS AND STRENGTHS

4 CONTRAINDICATIONS

5 WARNINGS AND PRECAUTIONS

- 5.1 Existing Pregnancy
- 5.2 Ectopic Pregnancy
- 5.3 Repeated Use
- 5.4 CYP3A4 Inducers
- 5.5 Fertility Following Use
- 5.6 Effect on Menstrual Cycle
- 5.7 Sexually Transmitted Infections/HIV

6 ADVERSE REACTIONS

- 6.1 Clinical Trials Experience
- 6.2 Postmarketing Experience

7 DRUG INTERACTIONS

- 7.1 Changes in Emergency Contraceptive Effectiveness Associated with Co-Administration of Other Products
- 7.2 Increase in Plasma Concentrations of ella Associated with Co-Administered Drugs
- 7.3 Effects of ella on Co-Administered Drugs

8 USE IN SPECIFIC POPULATIONS

- 8.1 Pregnancy
- 8.2 Lactation
- 8.3 Females and Males of Reproductive Potential
- 8.4 Pediatric Use
- 8.5 Geriatric Use
- 8.6 Race
- 8.7 Hepatic Impairment
- 8.8 Renal Impairment

10 OVERDOSAGE

11 DESCRIPTION

12 CLINICAL PHARMACOLOGY

- 12.1 Mechanism of Action
- 12.2 Pharmacodynamics
- 12.3 Pharmacokinetics

13 NONCLINICAL TOXICOLOGY

- 13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility

14 CLINICAL STUDIES

- 14.1 Open-Label Study
- 14.2 Single-Blind Comparative Study
- 14.3 Pooled Analysis

16 HOW SUPPLIED/STORAGE AND HANDLING

17 PATIENT COUNSELING INFORMATION

* Sections or subsections omitted from the full prescribing information are not listed.

FULL PRESCRIBING INFORMATION

1 INDICATIONS AND USAGE

Ellais is a progesterone agonist/antagonist emergency contraceptive indicated for prevention of pregnancy following unprotected intercourse or a known or suspected contraceptive failure [see *Dosage and Administration (2.1)*]. **Ellais** is not intended for routine use as a contraceptive.

2 DOSAGE AND ADMINISTRATION

2.1 Recommended Dosage and Administration

- Take one tablet of **ella** orally as soon as possible within 120 hours (5 days) after unprotected intercourse or a known or suspected contraceptive failure.
- Take **ella** with or without food.
- Take **ella** at any time during the menstrual cycle.

2.2 Recommendations Regarding Use with Hormonal Contraception

After **ella** use, initiate or resume hormonal contraception no sooner than 5 days after the intake of **ella** and use a reliable barrier method until the next menstrual period. For known or suspected failure of hormonal contraception refer to the hormonal contraceptive's prescribing information for instructions on what to do [see *Warnings and Precautions (5.5)*, *Drug Interactions (7.1)* and *Clinical Pharmacology (12.2)*].

2.3 Recommendation in Case of Gastrointestinal Disturbances

If vomiting occurs within 3 hours of taking **ella**, consider repeating the dose.

3 DOSAGE FORMS AND STRENGTHS

The **ella** tablet is supplied as a white to off-white, round, curved tablet containing 30 mg of ulipristal acetate and is marked " **ella** " on both sides.

4 CONTRAINDICATIONS

Ellais is contraindicated for use in the case of known or suspected pregnancy [see *Use in Specific Populations (8.1)*].

5 WARNINGS AND PRECAUTIONS

5.1 Existing Pregnancy

Ellais not indicated for termination of an existing pregnancy.

5.2 Ectopic Pregnancy

A history of ectopic pregnancy is not a contraindication to use of this emergency contraceptive method. Healthcare providers, however, should consider the possibility of ectopic pregnancy in women who become pregnant or complain of lower abdominal pain after taking **ella**. A follow-up physical or pelvic examination is recommended if there is any doubt concerning the general health or pregnancy status of any woman after taking **ella**.

5.3 Repeated Use

Ellais for occasional use as an emergency contraceptive. It should not replace a regular method of contraception. Repeated use of **ella** within the same menstrual cycle is not recommended, as safety and efficacy of repeat use within the same cycle has not been evaluated.

5.4 CYP3A4 Inducers

A CYP3A4 inducer, rifampin, decreases the plasma concentration of **ella** significantly. **Ellas** should not be administered with CYP3A4 inducers [see *Drug interactions (7.1)* and *Clinical Pharmacology (12.3)*].

5.5 Fertility Following Use

A rapid return of fertility is likely following treatment with **ella** for emergency contraception.

After use of **ella**, a reliable barrier method of contraception should be used with subsequent acts of intercourse until the next menstrual period.

After using **ella**, if a woman wishes to **initiate** hormonal contraception as a regular method, she can do so, no sooner than 5 days after the intake of **ella** and she should use a reliable barrier method until the next menstrual period [see *Dosage and Administration (2.2)*, *Drug Interactions (7.1 and 7.3)* and *Clinical Pharmacology (12.2)*].

Progestin-containing contraceptives may impair the ability of **ella** to delay ovulation. Advise women to follow the instructions on the initiation or resumption of hormonal contraceptives after **ella** intake [see *Dosage and Administration (2.2)*].

5.6 Effect on Menstrual Cycle

After **ella** intake, menses sometimes occur earlier or later than expected by a few days.

In clinical trials, cycle length was increased by a mean of 2.5 days but returned to normal in the subsequent cycle. Seven percent of subjects reported menses occurring more than 7 days earlier than expected, and 19% reported a delay of more than 7 days. If there is a delay in the onset of expected menses beyond 1 week, rule out pregnancy.

Nine percent of women studied reported intermenstrual bleeding after use of **ella**.

5.7 Sexually Transmitted Infections/HIV

Ellas does not protect against HIV infection (the virus that causes AIDS) or other sexually transmitted infections (STIs).

6 ADVERSE REACTIONS

6.1 Clinical Trials Experience

Because clinical trials are conducted under widely varying conditions, adverse reaction rates observed in the clinical trials of a drug cannot be directly compared to rates in the clinical trials of another drug and may not reflect the rates observed in clinical practice.

ella was studied in an open-label multicenter trial (Open-Label Study) and in a comparative, randomized, single-blind, multicenter trial (Single-Blind Comparative Study). In these studies, a total of 2,637 (1,533 + 1,104) women in the 30 mg ulipristal acetate groups were included in the safety analysis. The mean age of women who received ulipristal acetate was 24.5 years and the mean body mass index (BMI) was 25.3. The racial demographics of those enrolled were 67% Caucasian, 20% Black or African American, 2% Asian, and 12% other.

The most common adverse reactions ($\geq 10\%$) in the clinical trials for women receiving **ella** were headache (18% overall), nausea (12% overall) and abdominal and upper abdominal pain (12% overall). Table 1 lists those adverse reactions that were reported in $\geq 5\%$ of subjects in the clinical studies (14).

Table 1: Adverse Reactions in $\geq 5\%$ of Women (%) Receiving a Single Dose of ella (30 mg Ulipristal Acetate)

Most Common Adverse Reactions	Open-Label Study	Single-Blind Comparative Study
	N = 1,533	N = 1,104
Headache	18	19
Nausea	12	13
Abdominal and upper abdominal pain	15	8
Dysmenorrhea	7	13
Fatigue	6	6
Dizziness	5	5

6.2 Postmarketing Experience

Adolescents: the safety profile observed in adolescents aged 17 and younger in studies and post-marketing is similar to the safety profile in adults [see *Pediatric Use (8.4)*].

The following adverse reactions have been identified during post-approval use of **ella**:

Skin and Subcutaneous Tissue Disorders: Acne

Hypersensitivity reactions, including rash, urticaria, pruritis, and angioedema

Because these reactions are reported voluntarily from a population of uncertain size, it is not always possible to reliably estimate their frequency or establish a causal relationship

to drug exposure.

7 DRUG INTERACTIONS

Several *in vivo* drug interaction studies have shown that **ella** is predominantly metabolized by CYP3A4.

7.1 Changes in Emergency Contraceptive Effectiveness Associated with Co-Administration of Other Products

CYP3A inducers

Drugs or herbal products that induce CYP3A4 decrease the plasma concentrations of **ella**, and may decrease its effectiveness [see *Warnings and Precautions (5.4)* and *Clinical Pharmacology (12.3)*]. Avoid co-administration of **ella** and drugs or herbal products such as:

- barbiturates
- bosentan
- carbamazepine
- felbamate
- griseofulvin
- oxcarbazepine
- phenytoin
- rifampin
- St. John's Wort
- topiramate

Hormonal contraceptives

Progestin-containing contraceptives may impair the ability of **ella** to delay ovulation. After using **ella**, if a woman wishes to **initiate** or **resume** hormonal contraception, she can do so, no sooner than 5 days after the intake of **ella** and she should use a reliable barrier method until the next menstrual period. If a woman used **ella** due to a known or suspected failure of her hormonal contraception refer to the hormonal contraceptive's prescribing information for instructions on what to do [see *Dosage and Administration (2.2)*, *Warnings and Precautions (5.5)* and *Clinical Pharmacology (12.2)*].

7.2 Increase in Plasma Concentrations of ella Associated with Co-Administered Drugs

CYP3A4 inhibitors such as itraconazole or ketoconazole increase plasma concentrations of **ella** [see *Pharmacokinetics (12.3)*].

7.3 Effects of ella on Co-Administered Drugs

Hormonal contraceptives: **ella** may impact the effect of the progestin component of hormonal contraceptives. Therefore, if a woman wishes to use hormonal contraception after using **ella**, she should use a reliable barrier method for subsequent acts of intercourse until her next menstrual period [see *Dosage and Administration (2.2)*, *Warnings and Precautions (5.5)* and *Clinical Pharmacology (12.2)*].

8 USE IN SPECIFIC POPULATIONS

8.1 Pregnancy

Risk Summary

Ellais contraindicated for use during an existing or suspected pregnancy. No signal of concern regarding pregnancy complications was found in postmarketing studies [see *Data*]. Isolated cases of major malformations in **ella**-exposed pregnancies were identified; however, the data are not sufficient to determine a risk for birth defects with inadvertent use of **ella** during pregnancy. Miscarriage was reported in 14% of the known pregnancy outcomes; a rate that is similar to the U.S. background rate for miscarriage. In the U.S. general population, the estimated background risk of major birth defects and miscarriage in clinically recognized pregnancies is 2-4% and 15-20%, respectively.

In animal reproduction studies, no malformations were observed during repeated administration of ulipristal acetate to pregnant rats, rabbits and monkeys at daily drug exposures $\frac{1}{3}$, $\frac{1}{2}$, and 3 times respectively, the human exposure at a dose of 30 mg [see *Data*] .

Data

Human Data

Ellap pregnancy exposure data was collected in the U.S. and Europe from 1999 to 2015 and analyzed post-marketing using data from interventional clinical trials, observational studies and pharmacovigilance reports. Known pregnancy outcomes were available for 462/784 pregnancies in which women received **ella** at doses of 30 mg or greater during the conception cycle or during pregnancy. Data of pregnancies with known outcome were analyzed prospectively for 272 cases and retrospectively for 190 cases. Pregnancy outcomes included 302 elective abortions (2 for fetal anomalies including 1 with trisomy 21), 63 spontaneous abortions, and 13 ectopic pregnancies. No maternal or fetal deaths were reported. 84 pregnancies continued until birth, with congenital anomalies reported in 5 infants, including 4 major malformations (2/4 with genetic syndromes). Although these data do not allow estimation of the prevalence rate of congenital anomalies associated with inadvertent use of **ella** in pregnancy or determination of a causal relationship between reported anomalies and **ella**, they show that **ella**-exposed pregnancies were not associated with a pattern of increased risk of adverse outcomes.

Animal Data

Ulipristal acetate was administered repeatedly to pregnant rats and rabbits during the period of organogenesis. Embryo/fetal loss was noted in all pregnant rats and in half of the pregnant rabbits following 12 and 13 days of dosing, at daily drug exposures $\frac{1}{3}$ and $\frac{1}{2}$ the human exposure, respectively, based on body surface area (mg/m^2). There were no malformations of the surviving fetuses in these studies. Adverse effects were not observed in the offspring of pregnant rats administered ulipristal acetate during the period of organogenesis through lactation at drug exposures $\frac{1}{24}$ the human exposure based on AUC. Administration of ulipristal acetate to pregnant monkeys for 4 days during the first trimester caused pregnancy termination in 2/5 animals at daily drug exposures 3 times the human exposure based on body surface area.

8.2 Lactation

Risk Summary

Ulipristal acetate and its active metabolite, monodemethyl-ulipristal acetate, are present in human milk in small amounts (see *Data*). Based on the levels of drug and active metabolite measured in breastmilk, a fully breastfed child would receive a weight-adjusted dosage of approximately 0.8% of ulipristal acetate and monodemethyl-ulipristal acetate on Day 1 of drug administration and an approximate total of 1% of the maternal dose over a 5-day period after drug administration. There is no information on the effects on the breastfed child or the effects on milk production. The developmental and health benefits of breastfeeding should be considered along with the mother's clinical need for **ella** and any potential adverse effects on the breastfed child from **ella** or from the underlying maternal condition.

Data

The breast milk of 12 lactating women following administration of **ella** was collected in 24-hour increments to measure the concentrations of ulipristal acetate and the active metabolite monodemethyl-ulipristal acetate in breast milk. The mean daily concentrations of ulipristal acetate in breast milk were 22.7 ng/mL [0-24 hours], 2.96 ng/mL [24-48 hours], 1.56 ng/mL [48-72 hours], 1.04 ng/mL [72-96 hours], and 0.69 ng/mL [96-120 hours]. The mean daily concentrations of monodemethyl-ulipristal acetate in breast milk were 4.49 ng/mL [0-24 hours], 0.62 ng/mL [24-48 hours], 0.28 ng/mL [48-72 hours], 0.17 ng/mL [72-96 hours], and 0.10 ng/mL [96-120 hours]. Using these data, a fully breastfed infant would receive approximately 4.1 mcg/kg of ulipristal acetate and monodemethyl-ulipristal acetate on Day 1 following drug administration and approximately 5.2 mcg/kg over a five day period following drug administration.

8.3 Females and Males of Reproductive Potential

Contraception

Progestin-containing contraceptives may impair the ability of **ella** to delay ovulation. Advise females to use a reliable barrier method for subsequent acts of intercourse until her next menstrual period.

After using **ella**, if a woman wishes to **initiate** or **resume** hormonal contraception, she can do so, no sooner than 5 days after the intake of **ella** and she should use a reliable barrier method of contraception until the next menstrual period. If a woman used **ella** due to a known or suspected failure of her hormonal contraception refer to the hormonal contraceptive's prescribing information for instructions on what to do [see *Dosage and Administration (2.2)*, *Warnings and Precautions (5.5)*, *Drug Interactions (7)*, and *Clinical Pharmacology (12.2, 12.3)*].

8.4 Pediatric Use

There is no relevant use of ulipristal acetate for children of prepubertal age in the indication emergency contraception.

Adolescents:

Safety and efficacy of **ella** have been established in women of reproductive age. The clinical trials of **ella** enrolled 41 females under age 18, and a post-marketing observational study evaluating effectiveness and safety of **ella** in adolescents enrolled

279 females under age 18, including 76 under age 16 years. In these studies, the safety and efficacy profile observed in adolescents aged 17 and younger was similar to that in adults. Use of **ella** before menarche is not indicated.

8.5 Geriatric Use

This product is not intended for use in postmenopausal women.

8.6 Race

While no formal studies have evaluated the effect of race, a cross-study comparison of two pharmacokinetic studies indicated that exposure in South Asians may exceed that in Caucasians and African Americans. However, no difference in efficacy and safety was observed for women of different races in clinical studies.

8.7 Hepatic Impairment

No studies have been conducted to evaluate the effect of hepatic disease on the disposition of **ella**.

8.8 Renal Impairment

No studies have been conducted to evaluate the effect of renal disease on the disposition of **ella**.

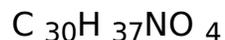
10 OVERDOSAGE

Experience with ulipristal acetate overdose is limited. In a clinical study, single doses equivalent to up to 4 times **ella** were administered to a limited number of subjects without any adverse reactions.

11 DESCRIPTION

The **ella** (ulipristal acetate) tablet for oral use contains 30 mg of a single active steroid ingredient, ulipristal acetate [17 α -acetoxy-11 β -(4-N,N-dimethylaminophenyl)-19-norpregna-4,9-diene-3,20-dione], a synthetic progesterone agonist/antagonist. The inactive ingredients are lactose monohydrate, povidone K-30, croscarmellose sodium and magnesium stearate.

Ulipristal acetate is a white to yellow crystalline powder which has a molecular weight of 475.6. The structural formula is:



12 CLINICAL PHARMACOLOGY

12.1 Mechanism of Action

When taken immediately before ovulation is to occur, **ella** postpones follicular rupture. The likely primary mechanism of action of ulipristal acetate for emergency contraception is therefore inhibition or delay of ovulation; however, alterations to the endometrium that may affect implantation may also contribute to efficacy.

12.2 Pharmacodynamics

Ulipristal acetate is a selective progesterone receptor modulator with antagonistic and partial agonistic effects (a progesterone agonist/antagonist) at the progesterone receptor. It binds the human progesterone receptor and prevents progesterone from occupying its receptor.

The pharmacodynamics of ulipristal acetate depend on the timing of administration in the menstrual cycle. Administration in the mid-follicular phase causes inhibition of folliculogenesis and reduction of estradiol concentration.

Pharmacodynamic data showed that administration of **ellato** 34 women in the late follicular phase postponed follicular rupture for at least 5 days in all (100%) of 8 subjects who took **ella** before the luteinizing hormone (LH) surge and 11 (79%) of 14 subjects who took **ella** immediately before ovulation (when LH has already started to rise). However, treatment was not effective in postponing follicular rupture when administered on the day of LH peak.

Dosing in the early luteal phase does not significantly delay endometrial maturation but decreases endometrial thickness by 0.6 ± 2.2 mm (mean \pm SD).

Hormonal Contraceptives after ella intake:

In women initiating hormonal contraception (quickstart scenario):

When a combined oral contraceptive pill (COC) containing ethinyl estradiol 30 μg + levonorgestrel 150 μg was started the day after **ella** intake during the follicular phase, **ella** did not interfere with the COC's ability to suppress ovarian activity, as assessed by measurement of follicle size via transvaginal ultrasound, combined with serum

progesterone and estradiol levels: ovarian activity was suppressed in 61.5% (24/39) of subjects receiving **ella**plus COC and 62.2% (23/37) of subjects receiving a placebo plus the COC. The incidence of ovulation was similar between the group that received **ella**plus the COC [33.3% (13/39)] and the group that received a placebo plus the COC [32.4% (12/37)]. [see *Dosage and Administration (2.2)*, *Warnings and Precautions (5.5)* and *Drug Interactions (7.3)*].

The initiation of a desogestrel 75 µg progestin-only pill (POP) the day after **ella**intake during the follicular phase was associated with a higher incidence of ovulation in the six days following **ella**intake compared to an **ella**-only treatment group, and a relatively slower onset (3 to 4 days) of thickened cervical mucus compared to a group given desogestrel without prior **ella**intake (2 days), suggesting an effect of prior use of **ella** on the ability of desogestrel to inhibit mucus permeability. [See *Dosage and Administration (2.2)*, *Warnings and Precautions (5.5)* and *Drug Interactions (7.1; 7.3)*].

When a COC containing ethinyl estradiol 30 µg + levonorgestrel 150 µg was started 2 days after **ella**intake, **ella**'s ability to delay ovulation, as assessed by transvaginal ultrasound, was reduced: follicular rupture occurred in 9/33 subjects (27%) in less than 5 days, compared to 1/33 subjects after **ella**alone (3%). [See *Dosage and Administration (2.2)*, *Warnings and Precautions (5.5)* and *Drug Interactions (7.1)*].

In women resuming combined hormonal contraception (missed pills scenario):

The effects on ovarian activity of delaying versus immediately resuming COCs after **ella**intake were investigated in 49 women who had been using COCs containing ethinyl estradiol 30 µg + levonorgestrel 150 µg once daily for 21 days followed by 7 days of placebo pills for at least one cycle. All subjects missed three consecutive pills (Days 5-7) during the first week of pills in the subsequent cycle and took **ella** on the following day (Day 8). These subjects were randomized to resume their COCs either on the same day as **ella**intake versus five days later. No ovulations with potential risk of pregnancy occurred in either group in the five days following **ella**intake. However, in the group that waited five days to resume taking COCs, 4/23 (17.4%) women did ovulate later in the cycle (Days 18-26) whereas no ovulations (0/26) occurred in the group that resumed COC intake on the same day as **ella** intake. Whether similar results can be expected with other COC products containing different progestins or missed active pills during different weeks in a treatment cycle is unknown.

12.3 Pharmacokinetics

Absorption

Following a single dose administration of **ella** in 20 women under fasting conditions, maximum plasma concentrations of ulipristal acetate and the active metabolite, monodemethyl-ulipristal acetate, were 176 and 69 ng/ml and were reached at 0.9 and 1 hour, respectively.

Figure 1: Mean (± SD) Plasma Concentration-time Profile of Ulipristal Acetate and Monodemethyl-ulipristal Acetate Following Single Dose Administration of 30 mg Ulipristal Acetate

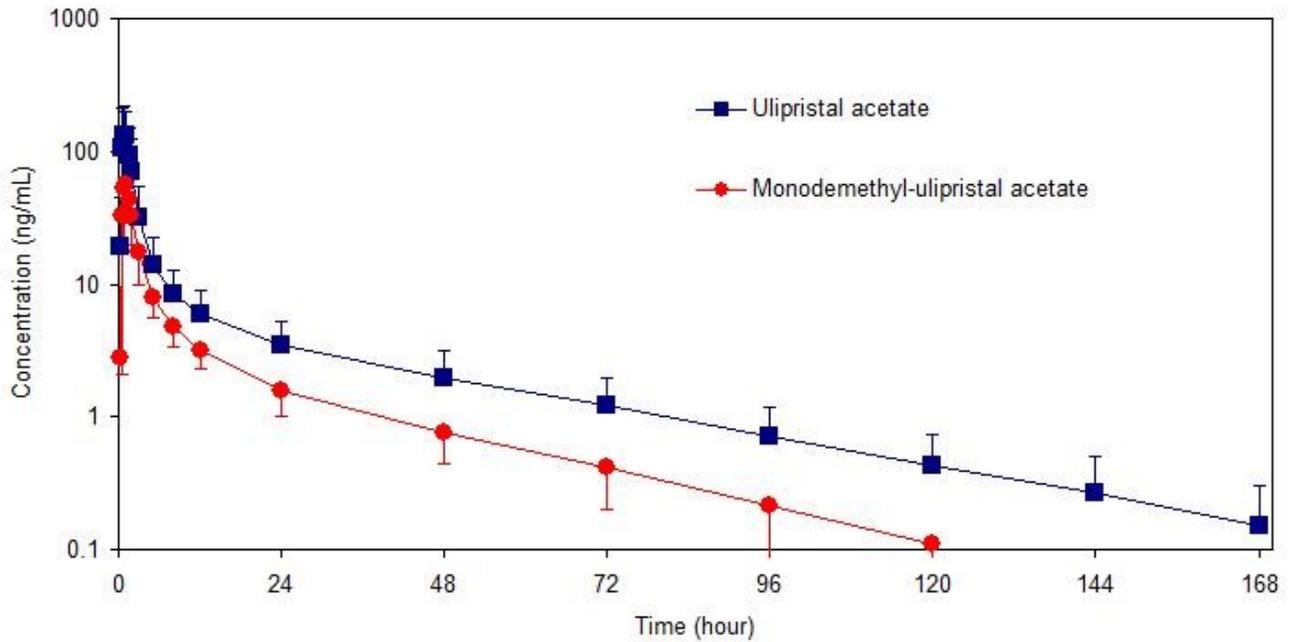


Table 2: Pharmacokinetic Parameter Values Following Administration of ella (ulipristal acetate) Tablet 30 mg to 20 Healthy Female Volunteers under Fasting Conditions

	Mean (\pm SD)				
	C_{max} (ng/ml)	AUC _{0-t} (ng•hr/ml)	AUC _{0-∞} (ng•hr/ml)	t_{max} (hr)*	$t_{1/2}$ (hr)
Ulipristal acetate	176 (89)	548 (259)	556 (260)	0.9 (0.5-2.0)	32 (6.3)
Monodemethyl-ulipristal acetate	69 (26)	240 (59)	246 (59)	0.9 (0.8-2.0)	27 (6.9)

C_{max} = maximum concentration

AUC_{0-t} = area under the drug concentration curve from time 0 to time of last determinable concentration

AUC_{0-∞} = area under the drug concentration curve from time 0 to infinity

t_{max} = time to maximum concentration

$t_{1/2}$ = elimination half-life

* Median (range)

Effect of food: Administration of **ella** together with a high-fat breakfast resulted in approximately 40 - 45% lower mean C_{max} , a delayed t_{max} (from a median of 0.75 hours to 3 hours) and 20 - 25% higher mean AUC_{0-∞} of ulipristal acetate and monodemethyl-ulipristal acetate compared with administration in the fasting state. These differences are not expected to impair the efficacy or safety of **ella** to a clinically significant extent; therefore, **ella** can be taken with or without food.

Distribution

Ulipristal acetate is highly bound (> 94%) to plasma proteins, including high density

lipoprotein, alpha-l-acid glycoprotein, and albumin.

Metabolism

Ulipristal acetate is metabolized to mono-demethylated and di-demethylated metabolites. *In vitro* data indicate that this is predominantly mediated by CYP3A4. The mono-demethylated metabolite is pharmacologically active.

Excretion

The terminal half-life of ulipristal acetate in plasma following a single 30 mg dose is estimated to be 32.4 ± 6.3 hours.

Drug interactions

CYP3A4 inducers: When a single 30 mg dose of ulipristal acetate was administered following administration of the strong CYP3A4 inducer, rifampin 600 mg once daily for 9 days, C_{max} and AUC of ulipristal acetate decreased by 90% and 93% respectively. The C_{max} and AUC of monodemethyl-ulipristal acetate decreased by 84% and 90% respectively [see Drug Interactions (7.1)] .

CYP3A4 inhibitors: When a single 10 mg dose of ulipristal acetate was administered following administration of the strong CYP3A4 inhibitor, ketoconazole 400 mg once daily for 7 days, C_{max} and AUC of ulipristal acetate increased by 2- and 5.9- fold, respectively. While the AUC of monodemethyl-ulipristal acetate increased by 2.4-fold, C_{max} of monodemethyl-ulipristal acetate decreased by 47%. There was no *in vivo* drug-drug interaction study between ulipristal acetate 30 mg and CYP3A4 inhibitors [see Drug Interactions (7.1)] .

In vitro studies demonstrated that **ella** does not induce or inhibit the activity of cytochrome P450 enzymes.

P-glycoprotein (P-gp) transporter: *In vitro* data indicate that ulipristal may be an inhibitor of P-gp at clinically relevant concentrations. When a single 60 mg dose of fexofenadine, a substrate of P-gp glycoprotein, was administered 1.5 hours after the administration of a single 10 mg dose of ulipristal acetate, there was no increase in C_{max} or AUC of fexofenadine.

Breast Cancer Resistance Protein (BCRP) transporter: *In vitro* data indicate that ulipristal acetate may be an inhibitor of BCRP at the intestinal level.

The effects of **ella** on P-gp and BCRP transporters are unlikely to have any clinical consequences when considering **ella**'s single dose treatment regimen, although there was no *in vivo* drug interaction study between ulipristal acetate 30 mg (**ella**) and substrates of P-gp and BCRP transporters.

13 NONCLINICAL TOXICOLOGY

13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility

Carcinogenicity:

Carcinogenicity potential was evaluated in rats and mice.

Sprague Dawley rats were exposed to ulipristal acetate daily for 99-100 weeks at doses of 1, 3, or 10 mg/kg/day, representing exposures up to 31 times higher than exposures

at the maximum recommended human dose (MRHD). There were no drug-related neoplasms in male rats. In female rats, potential treatment-related neoplastic findings were limited to adrenal cortical adenomas in the intermediate dose group (3 mg/kg/day). Despite the increase, this incidence of adrenal cortical adenomas in females may not be relevant to clinical use.

Tg.rasH2 transgenic mice were exposed to ulipristal acetate for 26 weeks at doses of 5, 45, or 130 mg/kg/day, representing exposures 100 times higher than exposures at the MRHD. There was no drug-related increase in neoplasm incidence in male or female mice.

Genotoxicity: Ulipristal acetate was not genotoxic in the Ames assay, *in vitro* mammalian assays utilizing mouse lymphoma cells and human peripheral blood lymphocytes, and in an *in vivo* micronucleus assay in mice.

Impairment of Fertility: Single oral doses of ulipristal acetate prevented ovulation in 50% of rats at 2 times the human exposure based on body surface area (mg/m²). Single doses of ulipristal acetate given on post-coital days 4 or 5 prevented pregnancy in 80-100% of rats and in 50% of rabbits when given on post-coital days 5 or 6 at drug exposures 4 and 12 times the human exposure based on body surface area. Lower doses administered for 4 days to rats and rabbits were also effective at preventing ovulation and pregnancy.

14 CLINICAL STUDIES

Two multicenter clinical studies evaluated the efficacy and safety of **ella**. An open-label study provided the primary data to support the efficacy and safety of ulipristal acetate for emergency contraception when taken 48 to 120 hours after unprotected intercourse. A single-blind comparative study provided the primary data to support the efficacy and safety of ulipristal acetate for emergency contraception when taken 0 to 72 hours after unprotected intercourse and provided supportive data for ulipristal acetate for emergency contraception when taken > 72 to 120 hours after unprotected intercourse. Women in both studies were required to have a negative pregnancy test prior to receiving emergency contraception. The primary efficacy analyses were performed on subjects less than 36 years of age who had a known pregnancy status after taking study medication.

Table 3: Summary of Clinical Trial Results for Women Who Received a Single Dose of ella (30 mg Ulipristal Acetate)

	Open-Label Study 48 to 120 Hours *	Single-Blind Comparative Study 0 to 72 Hours *
	N = 1,242	N = 844
Expected Pregnancy Rate **	5.5	5.6
Observed Pregnancy Rate ** (95% confidence interval)	2.2 (1.5, 3.2)	1.9 (1.1, 3.1)

* Time after unprotected intercourse when **ella** was taken

** Number of pregnancies per 100 women at risk for pregnancy

14.1 Open-Label Study

This study was a multicenter open-label trial conducted at 40 family planning clinics in the United States. Healthy women with a mean age of 24 years who requested emergency contraception 48 to 120 hours after unprotected intercourse received a dose of 30 mg ulipristal acetate (**ella**). The median BMI for the study subjects was 25.3 and ranged from 16.1 to 61.3 kg/m².

Twenty-seven pregnancies occurred in 1,242 women aged 18 to 35 years evaluated for efficacy. The number of pregnancies expected without emergency contraception was calculated based on the timing of intercourse with regard to each woman's menstrual cycle. **Ella** statistically significantly reduced the pregnancy rate, from an expected rate of 5.5% to an observed rate of 2.2%, when taken 48 to 120 hours after unprotected intercourse.

14.2 Single-Blind Comparative Study

This study was a multicenter, single-blind, randomized comparison of the efficacy and safety of 30 mg ulipristal acetate (**ella**) to levonorgestrel (another form of emergency contraception). Subjects were enrolled at 35 sites in the U.S., the United Kingdom and Ireland, with the majority (66%) having been enrolled in the U.S. Healthy women with a mean age of 25 years who requested emergency contraception within 120 hours of unprotected intercourse were enrolled and randomly allocated to receive **ella** or levonorgestrel 1.5 mg. The median BMI for the study subjects was 25.3 and ranged from 14.9 to 70.0 kg/m².

In the **ella** group, 16 pregnancies occurred in 844 women aged 16 to 35 years when emergency contraception was taken 0 to 72 hours after unprotected intercourse. The number of pregnancies expected without emergency contraception was calculated based on the timing of intercourse with regard to each woman's menstrual cycle; **ella** statistically significantly reduced the pregnancy rate, from an expected 5.6% to an observed 1.9%, when taken within 72 hours after unprotected intercourse. There were no pregnancies observed in the women who were administered **ella** more than 72 hours after unprotected intercourse (10% of women who received **ella**).

14.3 Pooled Analysis

Data from the two studies were pooled to provide a total efficacy population of women treated with ulipristal acetate up to 120 hours after unprotected intercourse. Time Trend analysis for the five 24-hour intervals from 0 to 120 hours between unprotected intercourse and treatment was conducted. There were no significant differences in the observed pregnancy rates across the five time intervals.

Subgroup analysis of the pooled data by BMI showed that for women with BMI > 30 kg/m² (16% of all subjects), the observed pregnancy rate was 3.1% (95% CI: 1.7, 5.7), which was not significantly reduced compared to the expected pregnancy rate of 4.5% in the absence of emergency contraception taken within 120 hours after unprotected intercourse. In the comparative study, a similar effect was seen for the comparator emergency contraception drug, levonorgestrel 1.5 mg. For levonorgestrel, when used by women with BMI > 30 kg/m², the observed pregnancy rate was 7.4% (95% CI: 3.9, 13.4), compared to the expected pregnancy rate of 4.4% in the absence of emergency

contraception taken within 72 hours after unprotected intercourse.

16 HOW SUPPLIED/STORAGE AND HANDLING

Ella(ulipristal acetate) tablet, 30 mg is supplied in a PVC-PE-PVDC or PVC-PVDC-aluminum blister. The tablet is a white to off-white, round, curved tablet marked with “**ella**” on both sides.

NDC 73302-456-01 (1 tablet unit of use package)

Store at 20-25°C (68-77°F); excursions permitted to 15-30°C (59-86°F) [See USP controlled room temperature.]

Keep the blister in the outer carton in order to protect from light. **Keep out of reach of children.**

17 PATIENT COUNSELING INFORMATION

Advise the patient to read the FDA-Approved patient Labeling (Patient Information).

Administration Instructions

- Instruct patients to take **ella** as soon as possible and not more than 120 hours after unprotected intercourse or a known or suspected contraceptive failure.
- Advise patients to contact their healthcare provider immediately in case of vomiting within 3 hours of taking the tablet, to discuss whether to take another tablet.
- Advise patients that after using **ella**, a reliable barrier method of contraception should be used for all subsequent acts of intercourse until the next menstrual period.
- Advise patients that after using **ella**, additional levonorgestrel emergency contraceptive pills should not be used within 5 days of **ella** intake.
- Advise patients that after using **ella**, hormonal contraception should be initiated or resumed no sooner than 5 days after the intake of **ella** and to use a reliable contraceptive barrier method until the next menstrual period.
- Advise patients with known or suspected failure of a hormonal contraceptive to refer to the hormonal contraceptive’s prescribing information for instructions on what to do.

Ectopic Pregnancy

Advise patients to seek medical attention if they experience severe lower abdominal pain 3 to 5 weeks after taking **ella**, in order to be evaluated for an ectopic pregnancy [see *Warnings and Precautions (5.2)*].

Effect on Menstrual Cycle

Advise women that after **ella** intake, menses may occur earlier or later than expected, by a few days. Advise patients to contact their healthcare provider and consider the possibility of pregnancy if their period is delayed after taking **ella** by more than 1 week beyond the date it was expected [see *Warnings and Precautions (5.6)*].

Sexually Transmitted Infections/HIV

Inform women that **ella** does not protect against HIV infection (the virus that causes AIDS) and other sexually transmitted diseases/infections [see *Warnings and Precautions (5.7)*].

Drug Interactions

Advise women to inform their healthcare provider if they take herbal supplements such as St. John's wort [see *Drug Interactions (7.1)*].

Patient Information **ella**(el-uh) **(ulipristal acetate)** **tablet, for oral use**

Read this Patient Information Leaflet before you take **ella**. There may be new information. This information does not take the place of talking to your healthcare provider about your medical condition or treatment.

What is ella?

- **Ellais** a prescription emergency contraceptive that reduces your chance of becoming pregnant if your birth control fails or you have unprotected sex.
- **Ellais** should not be used as your regular birth control. It is very important that you have a reliable form of birth control that is right for you.
- **Ellais** will not protect you against HIV infection (the virus that causes AIDS) and other sexually transmitted infections (STIs).

Who should not take ella?

Do not take **ella** if:

- you know or suspect you are already pregnant. **Ellais** not for use to end an existing pregnancy.

What should I tell my healthcare provider before taking ella? Before you take ella, tell your healthcare provider about all of your medical conditions, including if you:

- have a history of pregnancy outside the womb, usually in a fallopian tube (ectopic pregnancy).
- have taken **ella** during the same menstrual cycle for a different act of unprotected sex or birth control failure. **Do not** take **ella** more than 1 time in the same menstrual cycle.
- are breastfeeding or plan to breastfeed. A small amount of the hormones in **ella** may pass into your breast milk. Talk to healthcare provider about the best birth control for you while breastfeeding.

Tell your healthcare provider about all the medicines you take, including prescription and over-the-counter medicines, vitamins, and herbal supplements. Using some other medicines may make **ella** less effective. **Especially tell your healthcare provider if you take:** a hormonal form of birth control such as birth control pills, implant, IUD (intrauterine device) or patch, St. John's Wort, bosentan, griseofulvin, phenytoin, topiramate, oxcarbazepine, carbamazepine, barbiturates, rifampin, and felbamate. Know the medicines you take. Keep a list of them to show your healthcare provider and pharmacist when you get a new medicine.

How does ella work?

Ellais thought to work for emergency contraception primarily by stopping or delaying the release of an egg from the ovary. It is possible that **ella** may also work by preventing attachment (implantation) to the uterus.

How should I take ella?

- Take **ella** as soon as possible within 5 days (120 hours) after unprotected sex or if you had a birth control failure.
- **Ellais** can be taken with or without food.
- Contact your healthcare provider right away if you vomit within 3 hours of taking **ella**. Your healthcare provider may prescribe another dose of **ella** for you.

- **Ella** can be taken at any time during the menstrual cycle.
- **Ella** is meant for emergency contraception only and is not to be used frequently or as a regular birth control. If you need to use emergency contraception often, talk to your healthcare provider and learn about methods of birth control that are right for you and about sexually transmitted infection prevention.
- After using **ella**, you should use a reliable barrier birth control method (such as condom with spermicide) each time you have sex until your next menstrual period.
- After using **ella**, **do not** use any additional levonorgestrel emergency contraceptive pills within 5 days after taking **ella**.
- After using **ella**, wait 5 days to **start** or **resume** hormonal birth control as a regular method (such as the pill, patch or vaginal ring). If you used **ella** because you had a problem with your hormonal birth control, follow the instructions provided for that specific hormonal birth control product or contact your healthcare provider for further information. Be sure to use a reliable barrier birth control method (such as a condom with spermicide) each time you have sex until your next menstrual period.

What are the possible side effects of ella?

- **Pregnancy outside of the womb (ectopic pregnancy).** If you have severe lower stomach (abdominal) pain about 3 to 5 weeks after taking **ella**, you may have a pregnancy outside of the uterus (womb), which is called an ectopic or tubal pregnancy. An ectopic pregnancy is a serious condition that needs medical treatment right away. Call your healthcare provider or go to the nearest emergency room right away if you think you may have an ectopic pregnancy.
- **Changes in menstrual cycle.** Some women taking **ella** may have their next period earlier or later than expected. If your period is more than a week late, you should take a pregnancy test and follow up with your healthcare provider.

The most common side effects of **ella** include:

- headache
- nausea
- stomach (abdominal) pain
- menstrual pain (dysmenorrhea)
- tiredness
- dizziness

Tell your healthcare provider if you have any side effect that bothers you or that does not go away. These are not all the possible side effects of **ella**. For more information, ask your healthcare provider or pharmacist.

Call your doctor for medical advice about side effects. You may report side effects to FDA 1-800-FDA-1088.

How should I store ella?

- Store **ella** at room temperature between 68°F to 77°F (20°C to 25°C).
- Protect **ella** from light. Keep **ella** in the blister card inside the original box until you are ready to take it.

Keep **ella** and all medicines out of the reach of children.

General information about the safe and effective use of ella:

Medicines are sometimes prescribed for purposes other than those listed in a Patient Information Leaflet. Do not use **ella** for a condition for which it was not prescribed. Do not give **ella** to other people, even if they have the same symptoms that you have. It may harm them.

This Patient Information Leaflet summarizes the most important information about **ella**. If you would like more information, talk with your healthcare provider. You can ask your pharmacist or healthcare provider for information about **ella** that is written for health

professionals.

What are the ingredients in ella?

Active ingredients: ulipristal acetate, 30 mg

Inactive ingredients: lactose monohydrate, povidone, croscarmellose sodium, and magnesium stearate

Manufactured for:

HRA Pharma America Inc.

Morristown, NJ 07960

Under License From:

Laboratoire HRA Pharma

92320 Chatillon, France

ella[®] is a registered trademark of Laboratoire HRA Pharma

Manufactured by:

Cenexi, 95520 Osny, France, or

Laboratorios León Farma S.A., 24008 León, Spain or

Delpharm Lille SAS, 59452 Lys-Lez-Lannoy, France

For more information, go to www.ella-now.com or call 844-994-0329.

This Patient Information has been approved by the U.S. Food and Drug Administration

Revised: June 2021

General information about the safe and effective use of ella:

PRINCIPAL DISPLAY PANEL - 30 mg Tablet Blister Pack Carton

NDC 73302- 456-01

ella[®]

ulipristal acetate

tablet 30 mg

HRAPharma

Rx only

Contains 1 Tablet

ella[®]

ulipristal acetate

tablet 30 mg

Each tablet contains

30 mg ulipristal acetate.

Usual dose: See package insert.

Take one tablet as soon as possible, within 5 days after unprotected intercourse.

Ella is contraindicated during pregnancy.

Store at 20-25°C (68-77°F). [See USP controlled room temperature.]

Keep the blister in the outer carton in order to protect from light.

Keep out of reach of children.

For Medical Emergencies - Contact HRA Pharma America Inc, Health and Safety Team at 844-994-0329

HRAPharma

Manufactured for
HRA Pharma America Inc.
36 Cattano Ave. Suite 400 Morristown,
NJ 07960

Under License From:
Laboratoire HRA Pharma
92330 Chatillon, France

Product of Spain
Manufactured By:
Delpharm Lille S.A.S.
PARC D'ACTIVITE ROUBAIX EST
22 RUE DE TOUFFLERS
CS 50070
59492 LYS LEZ LANNOY
FRANCE

5044762

ella[®]
ulipristal acetate
tablet 30 mg

Each tablet contains
30 mg ulipristal acetate.

NDC 73302-456-01



ella[®]
ulipristal acetate
tablet 30 mg

HRA Pharma

Rx only
Contains 1 Tablet

5044762

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ulipristal acetate
tablet 30 mg

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HRA Pharma

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36 Cattano Ave. Suite 400 Morristown,
NJ 07960

Under License From:
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PARC D'ACTIVITE ROUBAIX EST
22 RUE DE TOUFFLERS
CS 50070
59452 LYS LEZ LANNOY
FRANCE



(01)00373302456012

ulipristal acetate table 30 mg

N 73302-456-1

Push through from front

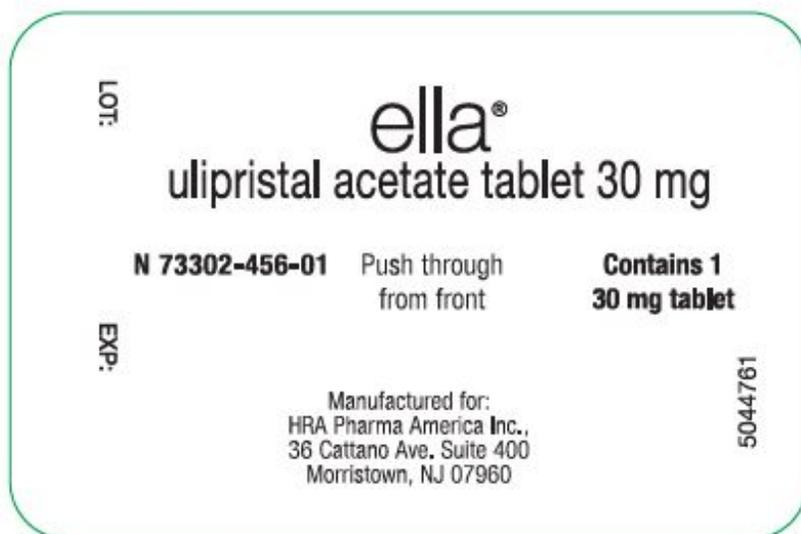
**Contains 1
30 mg tablet**

Manufactured for:
HRA Pharma America Inc.,
36 Cattano Ave. Suite 400
Morristown, NJ 07960

LOT:

EXP:

5044761



ELLA

ulipristal acetate tablet

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:73302-456
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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ULIPRISTAL ACETATE (UNII: YF7V70N02B) (ULIPRISTAL - UNII:6J5J15Q2X8)		ULIPRISTAL ACETATE	30 mg	
Inactive Ingredients				
Ingredient Name			Strength	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)				
POVIDONE K30 (UNII: U725QWY32X)				
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)				
MAGNESIUM STEARATE (UNII: 70097M6I30)				
Product Characteristics				
Color	white (white to off-white)	Score	no score	
Shape	ROUND	Size	9mm	
Flavor		Imprint Code	ella	
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73302-456-01	1 in 1 CARTON	05/11/2020	
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	NDA022474	05/11/2020		

Labeler - HRA PHARMA AMERICA, INC. (081160441)

Revised: 1/2026

HRA PHARMA AMERICA, INC.