

**CAMILA - norethindrone tablet**  
**Physicians Total Care, Inc.**

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**Revised AUGUST 2002**  
**1007150102**

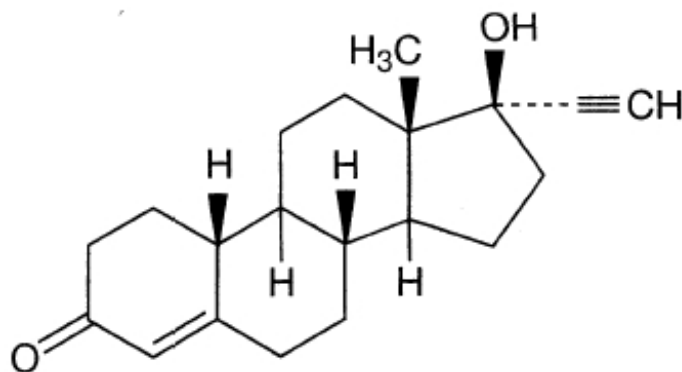
**Rx only**

**Patients should be counseled that this product does not protect against HIV infection (AIDS) and other sexually transmitted diseases.**

**DESCRIPTION**

Each light pink norethindrone tablet provides a continuous oral contraceptive regimen of 0.35 mg norethindrone daily, and has the following inactive ingredients: corn starch, FD&C red no. 40 aluminum lake, lactose monohydrate, magnesium stearate, povidone, and sodium starch glycolate.

The chemical name for norethindrone is 17-Hydroxy-19-Nor-17 $\alpha$ -pregn-4-en-20-yn-3-one. The structural formula follows:



Norethindrone Therapeutic class=oral contraceptive

**CLINICAL PHARMACOLOGY**

1. Mode of Action:

Progestin-only oral contraceptives prevent conception by suppressing ovulation in approximately half of users, thickening the cervical mucus to inhibit sperm penetration, lowering the midcycle LH and FSH peaks, slowing the movement of the ovum through the fallopian tubes, and altering the endometrium.

2. Pharmacokinetics:

Serum progestin levels peak about two hours after oral administration, followed by rapid distribution and elimination. By 24 hours after drug ingestion, serum levels are near baseline, making efficacy dependent upon rigid adherence to the dosing schedule. There are large variations in serum levels among individual users. Progestin-only administration results in lower steady-state serum progestin levels and a shorter elimination half-life than concomitant administration with estrogens.

**INDICATIONS AND USAGE**

1. Indications:

Progestin-only oral contraceptives are indicated for the prevention of pregnancy.

2. Efficacy:

If used perfectly, the first-year failure rate for progestin-only oral contraceptives is 0.5%. However,

the typical failure rate is estimated to be closer to 5%, due to late or omitted pills. The following table lists the pregnancy rates for users of all major methods of contraception.

**Table 1 Comparison of reversible contraceptive methods: Percent of women experiencing a contraceptive failure (pregnancy) during the first year of use.**

Method	Percent of women experiencing a pregnancy within the first year of use	
	Average Use	Perfect Use
No contraception	85	85
Spermicides	21	6
Periodic abstinence	20	1-9†
Withdrawal	19	4
Cervical caps		
Given birth	36	26
Never given birth	18	9
Diaphragms	18	6
Condoms		
Female	21	5
Male	12	3
Pills		
Progestin-only	3	0.5
Combined		0.1
IUDs		
Progesterone	2	1.5
Copper T 380A	0.8	0.6
Injectables	0.3	0.3
Implant	0.09	0.09

Adapted with permission. \*\*Hatcher RA, Trussel J, Stewart F, Stewart GK, Kowal D, Guest F, Cates W, Pollcar M. Contraceptive Technology 1994-1996. New York, NY: Irvington Publishers. 1994. †Depending on method (calendar, ovulation, symptom-thermal, post-ovulation).

## CONTRAINDICATIONS

Progestin-only oral contraceptives should not be used by women who currently have the following conditions:

- Known or suspected pregnancy
- Known or suspected carcinoma of the breast
- Undiagnosed abnormal genital bleeding
- Hypersensitivity to any component of this product
- Benign or malignant liver tumors
- Acute liver disease

## WARNINGS

Cigarette smoking greatly increases the possibility of suffering heart attacks and strokes. Women who use oral contraceptives are strongly advised not to smoke.

Norethindrone Tablets do not contain estrogen and, therefore, this insert does not discuss the serious

health risks that have been associated with the estrogen component of combined oral contraceptives. The healthcare provider is referred to the prescribing information of combined oral contraceptives for a discussion of those risks, including, but not limited to, an increased risk of serious cardiovascular disease in women who smoke, carcinoma of the breast and reproductive organs, hepatic neoplasia, and changes in carbohydrate and lipid metabolism. The relationship between progestin-only oral contraceptives and these risks have not been established and there are no studies definitely linking progestin-only pill (POP) use to an increased risk of heart attack or stroke.

The physician should remain alert to the earliest manifestation of symptoms of any serious disease and discontinue oral contraceptive therapy when appropriate.

#### 1. Ectopic Pregnancy

The incidence of ectopic pregnancies for progestin-only oral contraceptive users is 5 per 1000 woman-years. Up to 10% of pregnancies reported in clinical studies of progestin-only oral contraceptive users are extrauterine. Although symptoms of ectopic pregnancy should be watched for, a history of ectopic pregnancy need not be considered a contraindication to use of this contraceptive method. Health providers should be alert to the possibility of an ectopic pregnancy in women who become pregnant or complain of lower abdominal pain while on progestin-only oral contraceptives.

#### 2. Delayed Follicular Atresia/Ovarian Cysts:

If follicular development occurs, atresia of the follicle is sometimes delayed, and the follicle may continue to grow beyond the size it would attain in a normal cycle. Generally these enlarged follicles disappear spontaneously. Often they are asymptomatic; in some cases they are associated with mild abdominal pain. Rarely they may twist or rupture, requiring surgical intervention.

#### 3. Irregular Genital Bleeding:

Irregular menstrual patterns are common among women using progestin-only oral contraceptives. If genital bleeding is suggestive of infection, malignancy or other abnormal conditions, such nonpharmacologic causes should be ruled out. If prolonged amenorrhea occurs, the possibility of pregnancy should be evaluated.

#### 4. Carcinoma of the Breast and Reproductive Organs:

Some epidemiologic studies of oral contraceptive users have reported an increased relative risk of developing breast cancer, particularly at a younger age and apparently related to duration of use. These studies have predominantly involved combined oral contraceptives and there is insufficient data to determine whether the use of POPs similarly increases the risk. Women with breast cancer should not use oral contraceptives because the role of female hormones in breast cancer has not been fully determined.

Some studies suggest that oral contraceptive use has been associated with an increase in the risk of cervical intraepithelial neoplasia in some populations of women. However, there continues to be controversy about the extent to which such findings may be due to differences in sexual behavior and other factors. There is insufficient data to determine whether the use of POPs increases the risk of developing cervical intraepithelial neoplasia.

#### 5. Hepatic Neoplasia:

Benign hepatic adenomas are associated with combined oral contraceptive use, although the incidence of benign tumors is rare in the United States. Rupture of benign, hepatic adenomas may cause death through intraabdominal hemorrhage

Studies from Britain and the U.S. have shown an increased risk of developing hepatocellular carcinoma in combined oral contraceptive users. However, these cancers are rare. There is insufficient data to determine whether POPs increase the risk of developing hepatic neoplasia.

## **PRECAUTIONS**

### 1. General:

Patients should be counseled that this product does not protect against HIV infection (AIDS) and other sexually transmitted diseases.

#### 2. Physical Examination and Follow-up:

It is considered good medical practice for sexually active women using oral contraceptives to have annual history and physical examinations. The physical examination may be deferred until after initiation of oral contraceptives if requested by the woman and judged appropriate by the clinician.

#### 3. Carbohydrate and Lipid Metabolism:

Some users may experience slight deterioration in glucose tolerance, with increases in plasma insulin but women with diabetes mellitus who use progestin-only oral contraceptives do not generally experience changes in their insulin requirements. Nonetheless, prediabetic and diabetic women in particular should be carefully monitored while taking POPs.

Lipid metabolism is occasionally affected in that HDL, HDL<sub>2</sub>, and apolipoprotein A-I and A-II may be decreased; hepatic lipase may be increased. There is no effect on total cholesterol, HDL<sub>3</sub>, LDL, or VLDL.

#### 4. Drug Interactions:

The effectiveness of progestin-only pills is reduced by hepatic enzyme-inducing drugs such as the anticonvulsants phenytoin, carbamazepine, and barbiturates, and the anti-tuberculosis drug rifampin. No significant interaction has been found with broad-spectrum antibiotics.

#### 5. Interactions with Laboratory Tests:

The following endocrine tests may be affected by progestin-only oral contraceptive use

- Sex hormone-binding globulin (SHBG) concentrations may be decreased.
- Thyroxine concentrations may be decreased, due to a decrease in thyroid binding globulin (TBG).

#### 6. Carcinogenesis:

See WARNINGS section.

#### 7. Pregnancy:

Many studies have found no effects on fetal development associated with long-term use of contraceptive doses of oral progestins. The few studies of infant growth and development that have been conducted have not demonstrated significant adverse effects. It is nonetheless prudent to rule out suspected pregnancy before initiating any hormonal contraceptive use.

#### 8. Nursing Mothers:

No adverse effects have been found on breastfeeding performance or on the health, growth or development of the infant. Small amounts of progestin pass into the breast milk, resulting in steroid levels in infant plasma of 1-6% of the levels of maternal plasma.

#### 9. Fertility Following Discontinuation:

The limited available data indicate a rapid return of normal ovulation and fertility following discontinuation of progestin-only oral contraceptives.

#### 10. Headache:

The onset or exacerbation of migraine or the development of severe headache with focal neurological symptoms which is recurrent or persistent requires discontinuation of progestin-only contraceptives and evaluation of the cause.

#### 11. Pediatric Use:

Safety and efficacy of norethindrone have been established in women of reproductive age. Safety and efficacy are expected to be the same for postpubertal adolescents under the age of 16 and for users 16 years and older. Use of this product before menarche is not indicated.

## **INFORMATION FOR THE PATIENT**

1. See PATIENT LABELING for detailed information.
2. **Counseling Issues:**

The following points should be discussed with prospective users before prescribing progestin-only oral contraceptives:

- The necessity of taking pills at the same time every day, including throughout all bleeding episodes.
- The need to use a backup method such as condoms and spermicides for the next 48 hours whenever a progestin-only oral contraceptive is taken 3 or more hours late.
- The potential side effects of progestin-only oral contraceptives, particularly menstrual irregularities.
- The need to inform the clinician of prolonged episodes of bleeding, amenorrhea or severe abdominal pain.
- The importance of using a barrier method in addition to progestin-only oral contraceptives if a woman is at risk of contracting or transmitting STDs/HIV.

### **ADVERSE REACTIONS**

- Menstrual irregularity is the most frequently reported side effect.
- Frequent and irregular bleeding are common, while long duration of bleeding episodes and amenorrhea are less likely.
- Headache, breast tenderness, nausea, and dizziness are increased among progestin-only oral contraceptive users in some studies.
- Androgenic side effects such as acne, hirsutism, and weight gain occur rarely.

### **OVERDOSAGE**

There have been no reports of serious ill effects from overdosage, including ingestion by children.

### **DOSAGE AND ADMINISTRATION**

To achieve maximum contraceptive effectiveness, Norethindrone Tablets must be taken exactly as directed. One tablet is taken every day, at the same time. Administration is continuous, with no interruption between pill packs. See PATIENT LABELING for detailed instructions.

### **HOW SUPPLIED**

**Camila™**(norethindrone tablets, USP 0.35 mg) are packaged in cartons of six blister cards each containing 28 tablets. Each light pink, round, flat-faced, unscored tablet is debossed with stylized **b** on one side and **715** on the other side. (NDC 54868-4814-0)

Store at controlled room temperature 15°-30°C (59°-86°F) [See USP].

### **REFERENCES**

McCann M, and Potter L. Progestin-Only Oral Contraceptives: A Comprehensive Review. Contraception, 50:60 (Suppl. 1), December 1994.

### **DETAILED INFORMATION FOR THE PATIENT**

**This product (like all oral contraceptives) is used to prevent pregnancy. It does not protect against HIV infection (AIDS) and other sexually transmitted diseases.**

### **INTRODUCTION**

This leaflet is about birth control pills that contain one hormone, a progestin. Please read this leaflet before you begin to take your pills. It is meant to be used along with talking with your doctor or clinic.

Progestin-only pills are often called "POPs" or "the minipill." POPs have less progestin than the combined birth control pill (or "the pill") which contains both an estrogen and a progestin.

### HOW EFFECTIVE ARE POPS?

About 1 in 200 POPs users will get pregnant in the first year if they all take POPs perfectly (that is, on time, every day). About 1 in 20 "typical" POPs users (including women who are late taking pills or miss pills) gets pregnant in the first year of use. The following table will help you compare the efficacy of different methods.

**Table 1 Comparison of reversible contraceptive methods: Percent of women who become pregnant during the first year of use.**

Method	Percent of woman experiencing a pregnancy within the first year of use	
	Average Use	Perfect Use
No contraception	85	85
Spermicides	21	6
Periodic abstinence	20	1-9†
Withdrawal	19	4
Cervical caps		
Given birth	36	26
Never given birth	18	9
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### HOW DO POPS WORK?

- They make the cervical mucus at the entrance to the womb (the uterus) too thick for the sperm to get through to the egg.
- They prevent ovulation (release of the egg from the ovary) in about half the time.
- They also affect other hormones, the fallopian tubes and the lining of the uterus.

### YOU SHOULD NOT TAKE POPS

- If there is any chance you may be pregnant.

- If you have breast cancer.
- If you have bleeding between your periods which has not been diagnosed.
- If you are taking certain drugs for epilepsy (seizures) or for TB. (See USING POPS WITH OTHER MEDICINES below.)
- If you are hypersensitive or allergic to any component of this product.
- If you have liver tumors, either benign or cancerous.
- If you have acute liver disease.

## **RISKS OF TAKING POPS**

**WARNING:** If you have sudden or severe pain in your lower abdomen or stomach area, you may have an ectopic pregnancy or an ovarian cyst. If this happens, you should contact your doctor or clinic immediately.

### **1. Ectopic Pregnancy:**

An ectopic pregnancy is a pregnancy outside the womb. Because POPS protect against pregnancy, the chance of having a pregnancy outside the womb is very low. If you do get pregnant while taking POPS, you have a slightly higher chance that the pregnancy will be ectopic than do users of some other birth control methods.

### **2. Ovarian Cysts:**

These cysts are small sacs of fluid in the ovary. They are more common among POP users than among users of most other birth control methods. They usually disappear without treatment and rarely cause problems.

### **3. Cancer of the Reproductive Organs and Breasts:**

Some studies in women who use combined oral contraceptives that contain both estrogen and a progestin have reported an increase in the risk of developing breast cancer, particularly at a younger age and apparently related to duration of use. There is insufficient data to determine whether the use of POPS similarly increases this risk.

Some studies have found an increase in the incidence of cancer of the cervix in women who use oral contraceptives. However, this finding may be related to factors other than the use of oral contraceptives and there is insufficient data to determine whether the use of POPS increases the risk of developing cancer of the cervix.

### **4. Liver Tumors:**

In rare cases, combined oral contraceptives can cause benign but dangerous liver tumors. These benign liver tumors can rupture and cause fatal internal bleeding. In addition, a possible but not definite association has been found with combined oral contraceptives and liver cancers in studies in which a few women who developed these very rare cancers were found to have used combined oral contraceptives for long periods of time. There is insufficient data to determine whether POPS increase the risk of liver tumors.

## **SEXUALLY-TRANSMITTED DISEASES (STDs)**

**WARNING:** POPS do not protect against getting or giving someone HIV (AIDS) or any other STD, such as chlamydia, gonorrhea, genital warts or herpes.

## **SIDE EFFECTS**

### **1. Irregular Bleeding:**

The most common side effect of POPS is a change in menstrual bleeding. Your periods may be either early or late, and you may have some spotting between periods. Taking pills late or missing pills can also result in some spotting or bleeding.

### **2. Other Side Effects:**

Less common side effects include headaches, tender breasts, nausea and dizziness. Weight gain, acne and extra hair on your face and body have been reported, but are rare.

If you are concerned about any of these side effects, check with your doctor or clinic.

## **USING POPS WITH OTHER MEDICINES**

Before taking a POP, inform your healthcare provider of any other medication, including over-the-counter medicine, that you may be taking.

If you are taking medicines for seizures (epilepsy) or tuberculosis (TB), tell your doctor or clinic. These medicines can make POPs less effective:

Medicines for seizures:

- Phenytoin (Dilantin®)
- Carbamazepine (Tegretol®)
- Phenobarbital

Medicine for TB:

- Rifampin (Rifampicin)

Before you begin taking any new medicines be sure your doctor or clinic knows you are taking birth control pills that contain a progestin.

## **HOW TO TAKE POPS**

### **IMPORTANT POINTS TO REMEMBER**

- POPS must be taken at the same time every day, so choose a time and then take the pill at that same time every day. Every time you take a pill late, and especially if you miss a pill, you are more likely to get pregnant.
- Start the next pack the day after the last pack is finished. There is no break between packs. Always have your next pack of pills ready.
- You may have some menstrual spotting between periods. Do not stop taking your pills if this happens.
- If you vomit soon after taking a pill, use a backup method (such as condom and/or spermicide) for 48 hours.
- If you want to stop taking POPS, you can do so at any time, but, if you remain sexually active and don't wish to become pregnant, be certain to use another birth control method.
- If you are not sure about how to take POPS, ask your doctor or clinic.

### **STARTING POPS**

- It's best to take your first POP on the first day of your menstrual period.
- If you decide to take your first POP on another day, use a backup method (such as condom and/or spermicide) every time you have sex during the next 48 hours.
- If you have had a miscarriage or an abortion, you can start POPS the next day.

### **IF YOU ARE LATE OR MISS TAKING YOUR POPS**

- If you are more than 3 hours late or you miss one or more POPS:
  1. TAKE a missed pill as soon as you remember that you missed it,
  2. THEN go back to taking POPS at your regular time,
  3. BUT be sure to use a backup method (such as condom and/or spermicide) every time you have sex for the next 48 hours.
- If you are not sure what to do about the pills you have missed, keep taking POPS and use a backup method until you can talk to your doctor or clinic.

### **IF YOU ARE BREASTFEEDING**

- If you are fully breastfeeding (not giving your baby any food or formula), you may start your pills 6 weeks after delivery.



- If you are partially breastfeeding (giving your baby some food or formula), you should start taking pills by 3 weeks after delivery.

## **IF YOU ARE SWITCHING PILLS**

- If you are switching from the combined pills to POPs, take the first POP the day after you finish the last active combined pill. Do not take any of the 7 inactive pills from the combined pill pack. You should know that many women have irregular periods after switching to POPs, but this is normal and to be expected.
- If you are switching from POPs to the combined pills, take the first active combined pill on the first day of your period, even if your POPs pack is not finished.
- If you switch to another brand of POPs, start the new brand anytime.
- If you are breastfeeding, you can switch to another method of birth control at any time, except do not switch to the combined pills until you stop breastfeeding or at least until 6 months after delivery.

## **PREGNANCY WHILE ON THE PILL**

If you become pregnant, or think you might be, stop taking POPs and contact your physician. Even though research has shown that POPs do not cause harm to the unborn baby, it is always best not to take any drugs or medicines that you don't need when you are pregnant.

You should get a pregnancy test:

- If your period is late and you took one or more pills late or missed taking them and had sex without a backup method.
- Anytime you miss 2 periods in a row.

## **WILL POPS AFFECT YOUR ABILITY TO GET PREGNANT LATER?**

If you want to become pregnant, simply stop taking POPs. POPs will not delay your ability to get pregnant.

## **BREASTFEEDING**

If you are breastfeeding, POPs will not affect the quality or amount of your breastmilk or the health of your nursing baby.

## **OVERDOSE**

No serious problems have been reported when many pills were taken by accident, even by a small child, so there is usually no reason to treat an overdose.

## **OTHER QUESTIONS OR CONCERNS**

Cigarette smoking greatly increases the possibility of suffering heart attacks and strokes. Women who use oral contraceptives are strongly advised not to smoke.

Diabetic women taking POPs do not generally require changes in the amount of insulin they are taking. However, your physician may monitor you more closely under these conditions.

If you have any questions or concerns, check with your doctor or clinic. You can also ask for the more detailed "professional package labeling" written for doctors and other healthcare providers.

## **HOW TO STORE YOUR POPS**

Store your POPs at room temperature 15°-30°C (59°-86°F).

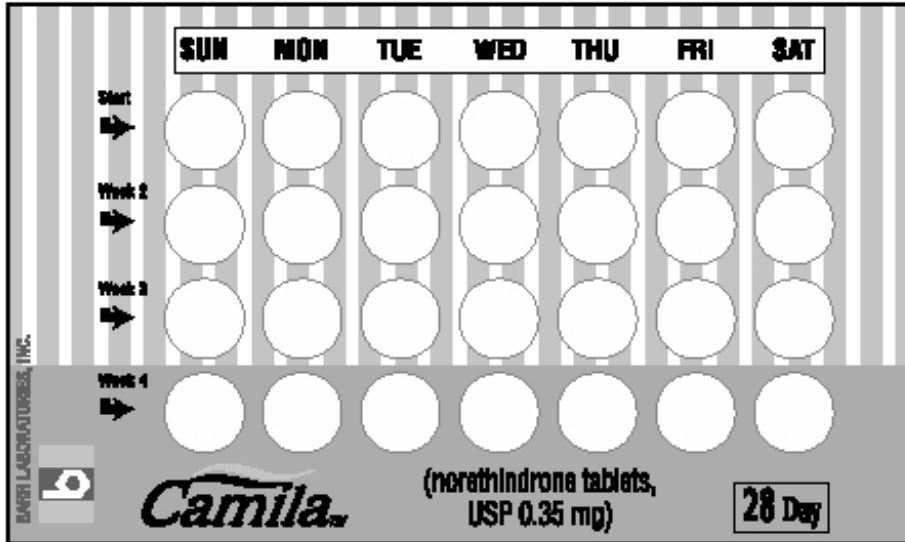
## **INSTRUCTIONS TO PATIENTS**

### **How to Use the Camila™ Tablets Blister Card:**

1. The first time you use these pills, take your first pill on the first day of your menstrual period. Pick the Days of the Week Sticker that starts the first day of your period. When you have picked the right sticker, throw away the others and place the sticker on the blister card over the pre-printed days of

the week and make sure it lines up with the pills.

- Your blister package consists of three parts, the foil pouch, wallet, and a blister card containing 28 individually sealed pills. Note that the pills are arranged in four numbered rows of 7 pills, with the pre-printed days of the week printed above them. All 28 pills are "active" birth control pills. Refer to the sample of the blister card below:
- 



- To remove a pill, push down on the pill with your thumb and forefinger so that the pill releases through the back of the blister card. Each day, take one pill. Always go from left to right along the row. Each new row will begin on the same day of the week.
- Take one pill every day for 28 days, whether bleeding or not, until you have taken all the pills. It is important that you take your pill at the same time every day.
- After you have taken all 28 pills, begin taking your pills again the next day. Be sure that the calendar day on your new package corresponds with the actual day.

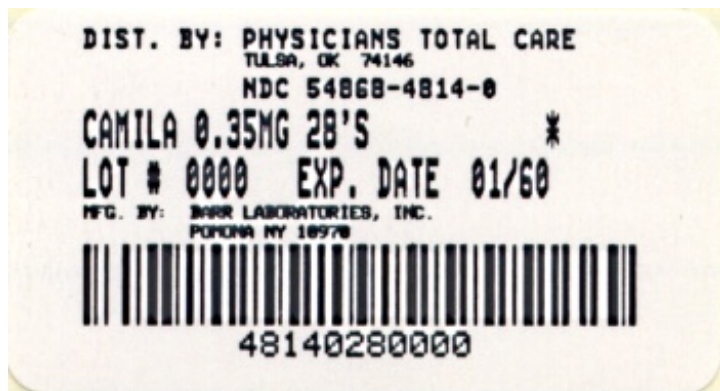
**MANUFACTURED BY  
BARR LABORATORIES, INC.  
POMONA, NY 10970**

**Rev AUGUST 2002**

**Relabeling of "Additional Barcode Label" by:**  
Physicians Total Care, Inc.  
Tulsa, OK 74146

**PRINCIPAL DISPLAY PANEL**

**Camila™**(norethindrone tablets, USP 0.35 mg)



## CAMILA

norethindrone tablet

### Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:54868-4814(NDC:0555-0715)
Route of Administration	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
NORETHINDRONE (UNII: T18F433X4S) (NORETHINDRONE - UNII:T18F433X4S)	NORETHINDRONE	0.35 mg

### Inactive Ingredients

Ingredient Name	Strength
STARCH, CORN (UNII: O8232NY3SJ)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POVIDONE (UNII: FZ989GH94E)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	

### Product Characteristics

Color	pink (pink)	Score	no score
Shape	ROUND (round)	Size	6mm
Flavor		Imprint Code	715
Contains			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54868-4814-0	28 in 1 BLISTER PACK		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA076177	06/19/2003	

**Labeler** - Physicians Total Care, Inc. (194123980)

**Establishment**

Name	Address	ID/FEI	Business Operations
Physicians Total Care, Inc.		194123980	relabel

Revised: 12/2010

Physicians Total Care, Inc.