Patient Assistance Program
250 Phillips Blvd, Ste 250, Ewing, NJ 08618
1-800-425-3122 Telephone 1-800-685-2577 Fax
Hours of Operation: Monday through Friday, 8:30 AM to 5:30 PM EST

Seasonique® Patient Assistance Program Eligibility Requirements

A 91-day supply of Seasonique® will be provided free of charge to patients who meet program eligibility requirements:

- Patient must be a US resident
- Patient must be 18 years of age or older
- Patient's gross annual household income must be at or below 200% HHS Poverty Guidelines*
- Patient must provide proof of gross annual household income
    - Financial documentation must be included with the Qualification Form
    - Proof of income includes copies of both:
        - a) federal tax return (Form 1040 or 1040EZ) for prior tax year, and
        - b) all other recent documents that show income paid to patient (and/or spouse if married), such as: wage and tax statements (W-2 forms), Social Security, Pension, or Railroad Retirement statements (SSA-1099 or similar), Statements of interest, dividends, or other income (1099-INT, 1099, 1099-DIV, or other forms)
- Patient has no insurance (public or private) or third-party payer prescription drug coverage (in whole or in part), including Medicaid or Medicare Part D
    - If patient has coverage for any prescription drug (not only Seasonique), the patient is ineligible for this program

Additional requirements:

- Program Qualification Form must be completed in its entirety by the healthcare professional caring for the patient
- Both patient and healthcare professional must sign the Qualification Form in the appropriate section
- Patient must sign and submit the Authorization to Disclose Form
- Healthcare professional must have a current valid state license

* Income criterion is based on Health and Human Services Poverty Guidelines. These guidelines can be revised each new year, usually around February. Website is: http://aspe.hhs.gov/poverty/index.shtml

Please see full prescribing information.

Teva Women's Health, Inc. reserves the right to limit enrollment of patients to the **Seasonique Patient Assistance Program** at any time.

The program administrators reserve the right any time and without notice to modify the application form, modify or discontinue any or all of the program and the related eligibility criteria; or at any time terminate assistance provided by the program.

seasonique
levonorgestrel/ethinyl estradiol 0.15 mg/0.03 mg TABLETS
ethinyl estradiol 0.01 mg TABLETS

Seasonique® Patient Assistance Program

250 Phillips Blvd, Ste 250, Ewing, NJ 08618
Phone: 1.800.425.3122 Fax: 1.800.685.2577
Qualification Form

PhRMA

PATIENT INFORMATION (Please Print) Patient must be a U.S. resident

First Name: _____ MI: _____ Last Name: _____

Address: _____ City: _____

State: _____ Zip Code: _____ Phone: _____ Social Security #: _____

Date of Birth: (mm/dd/yyyy) _____ (Patient must be 18 years of age or older) Patient's Diagnosis (ICD.9 Code): _____

PATIENT'S INCOME:

Current gross annual household income: $ _____ Number of household members dependent on income (including patient) _____ Number of children: _____

Patient financial documentation must be included with this application. Proof of income includes copies of both: a) your federal tax return (Form 1040 or 1040EZ) for prior tax year, **and** b) all other recent documents that show income paid to you (or your spouse if married), such as: wage and tax statements (W-2 forms), Social Security, Pension, or Railroad Retirement statements (SSA-1099 or similar), Statements of interest, dividends, or other income (1099-INT, 1099, 1099-DIV, or other forms)

PATIENT'S INSURANCE AND PRESCRIPTION COVERAGE (PARTIAL OR FULL) Check all that apply:

☐	Medicare	☐	Medicare Advantage (MA)	☐	Includes Rx	☐ Private Foundation	☐ Includes Rx
☐	Medicaid	☐	State/Local Government	☐	Includes Rx	☐ Medicare Medigap	☐ Includes Rx
☐	Medicaid QMB	☐	Federal Program	☐	Includes Rx	☐ Private Prescription Drug Plan (PDP)	
☐	Uninsured	☐	Private Insurance / HMO	☐	Includes Rx	☐ Other:	Specify:

If Rx Coverage is Yes, name of insurance carrier: _____ If Rx Coverage is Yes, is Seasonique covered? ☐ Yes

PATIENT/APPLICANT DECLARATION

I understand that completing this form does not ensure that I will qualify for this program. I verify that the information provided in this qualification form is complete and accurate. I agree to notify the Seasonique Patient Assistance Program if I obtain prescription drug coverage or if I no longer meet the income criteria. I understand that the program administrators reserve the right any time and without notice to modify the application form, modify or discontinue any or all of the program and the related eligibility criteria; or terminate assistance provided by the program at any time. No claim may be made to any third party payer for payment for product or administration of product provided under the Program.

Patient's Original Signature: _____ Date: (mm/dd/yyyy) _____

HEALTHCARE PROFESSIONAL'S INFORMATION (Please Print)

First Name: _____ MI: _____ Last Name: _____

Facility: _____ Office Contact Name: _____

Street: _____ Bldg/Suite/Floor/Room: _____

City: _____ State: _____ Zip Code: _____ Phone: _____

Fax: _____ Specialty: _____ State License #: _____

E-Mail Address: _____ When you provide your e-mail address, you agree that Teva Women's Health, Inc. and its agents may contact you regarding the Seasonique PAP.

This section of the form will serve as the Seasonique® prescription:

Rx: SEASONIQUE® (levonorgestrel/ethinyl estradiol tablets 0.15 mg/0.03 mg and ethinyl estradiol tablets 0.01 mg) QD sig – one tablet

[x] **Seasonique® Extended-Cycle Tablet Dispenser**	**Quantity: 1 (91 individual pills)**	**Indicate number of prescribed refills, if any:**	Refill: ☐ 1 Time	Refill: ☐ 2 Times	Refill: ☐ 3 Times

HEALTHCARE PROFESSIONAL ATTESTATION

I represent that the information contained in this application is complete and accurate to the best of my knowledge. To the best of my knowledge, this patient has no prescription insurance coverage for the requested medication, including Medicaid or other public programs, and the patient has insufficient financial resources to pay for the prescribed therapy. No claim may be made to any third party payer for payment of Seasonique provided by this Patient Assistance Program. Seasonique received for this patient may not be sold or traded, may not be returned for credit, and is not a sample. I understand that the Seasonique Patient Assistance Program has the right to modify or discontinue this program and its eligibility requirements, or to terminate assistance, at any time and without prior notice. Please indicate that you agree to these terms by signing below. Your signature also confirms that there is a valid medical need for this patient's prescription for Seasonique.

Healthcare Professional's Original Signature: _____ Date: (mm/dd/yyyy) _____

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PATIENT ASSISTANCE PROGRAM
250 Phillips Blvd, Ste 250, Ewing, NJ 08618
Phone: 1.800.425.3122 Fax: 1.800.685.2577

Patient Authorization to Disclose Protected Health Information

To the Patient: During the course of your participation in the Seasonique® Patient Assistance Program, you or your caregiver and your health care professional will provide personal identifying information to Teva Women's Health, Inc., its affiliated companies and subcontractors on a need to know basis for purposes of administering the Seasonique® Patient Assistance Program (the "Program"). This information may constitute Protected Health Information (PHI) under the privacy rules of the Health Insurance Portability and Accountability Act (HIPAA), and you need to authorize your health care professional and caregiver, if any, to release your PHI to the Teva Team and authorize the Teva Team to use the PHI for the Program.

Authorization Statement

I, (Patient's Name) , authorize my prescribing healthcare professional,
(HCP's Name)
(HCP's Address)

and caregiver as deemed necessary to disclose any personal identifying information to Teva Women's Health Inc., its affiliated companies and subcontractors (the "Teva Team") on a need to know basis to use for purposes of administering the Program for the duration of my participation in the Program. Although the Teva Team values my privacy and intends to take reasonable and appropriate measures to protect the information provided from inappropriate disclosure and to use the information only for administering the Program or as required by law, I understand that once information is disclosed to the Teva Team, it may no longer be protected under federal privacy laws and could be redisclosed to others.

I understand that I may refuse to sign this authorization, and my right to treatment, insurance enrollment, eligibility for insurance benefits or my receipt of Seasonique® are not conditioned on my signing this authorization. However, if I do not sign this authorization, I will not be able to participate in the Program.

I understand that I may revoke this authorization, in writing, at any time, except to the extent action has been taken in reliance on it, by addressing such revocation to Seasonique® Patient Assistance Program 250 Phillips Blvd, Ste 250, Ewing, NJ 08618 (your healthcare professional will be advised) and that only a written revocation addressed to the Program will constitute an effective withdrawal of my authorization. I understand that I may request a copy of this form from the Seasonique® Assistance Program.

Required Signature _____ _____
Signature of patient or legal representative **Date**

If signed by patient's legal representative, complete the following:

Print name of legal representative: _____

Describe representative's authority to act for patient: _____

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Important:

To the Patient:

Once you have completed and signed this authorization form, please give it to your healthcare professional. Do not send it to the Seasonique® Patient Assistance Program.

To the Healthcare Professional:

Retain the original copy of the Patient Authorization to Disclose Protected Health Information for your records. Please return a copy of this signed form along with the completed Qualification application form to the Seasonique® Patient Assistance Program, 250 Phillips Blvd, Ste 250, Ewing, NJ 08618, or fax to 1.800.685.2577.

PhRMA

SEASONIQUE®

(levonorgestrel / ethinyl estradiol tablets) 0.15 mg / 0.03 mg and (ethinyl estradiol tablets) 0.01 mg

Brief Summary. See full package brochure for complete information.

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

CONTRAINDICATIONS: Oral contraceptives should not be used in women who currently have the following conditions: • Thrombophlebitis or thromboembolic disorders • A past history of deep vein thrombophlebitis or thromboembolic disorders • Cerebrovascular or coronary artery disease (current or history) • Valvular heart disease with thrombogenic complications • Uncontrolled hypertension • Diabetes with vascular involvement • Headaches with focal neurological symptoms • Major surgery with prolonged immobilization • Known or suspected carcinoma of the breast or personal history of breast cancer • Carcinoma of the endometrium or other known or suspected estrogen dependent neoplasia • Undiagnosed abnormal genital bleeding • Cholestatic jaundice of pregnancy or jaundice with prior pill use • Hepatic adenomas or carcinomas, or active liver disease • Known or suspected pregnancy • Hypersensitivity to any component of this product

WARNINGS

Cigarette smoking increases the risk of serious cardiovascular side effects from oral contraceptive use. This risk increases with age and with heavy smoking (15 or more cigarettes per day) and is quite marked in women over 35 years of age. Women who use oral contraceptives should be strongly advised not to smoke.

The use of oral contraceptives is associated with increased risk of several serious conditions including venous and arterial thrombotic and thromboembolic events (such as myocardial infarction, thromboembolism, and stroke), hepatic neoplasia, gallbladder disease, and hypertension. The risk of serious morbidity or mortality is very small in healthy women without underlying risk factors. The risk of morbidity and mortality increases significantly in the presence of other underlying risk factors such as certain inherited thrombophilias, hypertension, hyperlipidemias, obesity and diabetes.

Practitioners prescribing oral contraceptives should be familiar with the following information relating to these risks. The information contained in this brief summary is principally based on studies carried out in patients who used oral contraceptives with higher formulations of estrogens and progestogens than those in common use today. The effect of long-term use of the oral contraceptives with lower doses of both estrogens and progestogens remains to be determined. Throughout this labeling, epidemiological studies reported are of two types: retrospective or case control studies and prospective or cohort studies. Case control studies provide a measure of the relative risk of a disease, namely, a ratio of the incidence of a disease among oral contraceptive users to that among nonusers. The relative risk does not provide information on the actual clinical occurrence of a disease. Cohort studies provide a measure of attributable risk, which is the difference in the incidence of disease between oral contraceptive users and nonusers. The attributable risk does provide information about the actual occurrence of a disease in the population. For further information, the reader is referred to a text on epidemiological methods.

1. Thromboembolic Disorders and Other Vascular Problems: Use of Seasonique® provides women with more hormonal exposure on a yearly basis than conventional monthly oral contraceptives containing similar strength synthetic estrogens and progestins (an additional 13 weeks of exposure to birth control pill hormones per year). • a. Myocardial Infarction: An increased risk of myocardial infarction has been attributed to oral contraceptive use. This risk is primarily in smokers or women with other underlying risk factors for coronary artery disease such as hypertension, hypercholesterolemia, morbid obesity, and diabetes. The relative risk of heart attack for current oral contraceptive users has been estimated to be two to six. The risk is very low under the age of 30. Smoking in combination with oral contraceptive use has been shown to contribute substantially to the incidence of myocardial infarction in women in their mid-thirties or older with smoking accounting for the majority of excess cases. Mortality rates associated with circulatory disease have been shown to increase substantially in smokers over the age of 35 and nonsmokers over the age of 40 among women who use oral contraceptives. Oral contraceptives may compound the effects of well-known risk factors, such as hypertension, diabetes, hyperlipidemias, age and obesity. In particular, some progestogens are known to decrease HDL cholesterol and cause glucose intolerance, while estrogens may create a state of hyperinsulinism. Oral contraceptives have been shown to increase blood pressure among users (see section 9 in **WARNINGS**). The severity and number of risk factors increase heart disease risk. Oral contraceptives must be used with caution in women with cardiovascular disease risk factors. • b. Thromboembolism: An increased risk of thromboembolic and thrombotic disease associated with the use of oral contraceptives is well established. Case control studies have found the relative risk of users compared to nonusers to be 3 for the first episode of superficial venous thrombosis, 4 to 11 for deep vein thrombosis or pulmonary embolism, and 1.5 to 6 for women with predisposing conditions for venous thromboembolic disease. Cohort studies have shown the relative risk to be somewhat lower, about 3 for new cases and about 4.5 for new cases requiring hospitalization. The approximate incidence of deep vein thrombosis and pulmonary embolism in users of low dose (<50 µg ethinyl estradiol) combination oral contraceptives is up to 4 per 10,000 woman-years compared to 0.5-3 per 10,000 woman-years for non-users. However, the incidence is less than that associated with pregnancy (6 per 10,000 woman-years). The risk of thromboembolic disease due to oral contraceptives is not related to length of use and disappears after pill use is stopped. A two- to four-fold increase in relative risk of postoperative thromboembolic complications has been reported with the use of oral contraceptives. The relative risk of venous thrombosis in women who have predisposing conditions is twice that of women without such medical conditions. If feasible, oral contraceptives should be discontinued at least four weeks prior to and for two weeks after elective surgery of a type associated with an increase in risk of thromboembolism and during and following prolonged immobilization. Since the immediate postpartum period is also associated with an increased risk of thromboembolism, oral contraceptives should be started no earlier than four weeks after delivery in women who elect not to breast-feed. • c. Cerebrovascular Diseases: Oral contraceptives have been shown to increase both the relative and attributable risks of cerebrovascular events (thrombotic and hemorrhagic strokes), although, in general, the risk is greatest among older (>35 years), hypertensive women who also smoke. Hypertension was found to be a risk factor for both users and nonusers, for both types of strokes, while smoking interacted to increase the risk for hemorrhagic strokes. In a large study, the relative risk of thrombotic strokes has been shown to range from 3 for normotensive users to 14 for users with severe hypertension. The relative risk of hemorrhagic stroke is reported to be 1.2 for nonsmokers who used oral contraceptives, 2.6 for smokers who did not use oral contraceptives, 7.6 for smokers who used oral contraceptives, 1.8 for normotensive users and 25.7 for users with severe hypertension. The attributable risk is also greater in older women. Oral contraceptives also increase the risk for stroke in women with other underlying risk factors such as certain inherited or acquired thrombophilias, hyperlipidemias, and obesity. Women with migraine (particularly migraine with aura) who take combination oral contraceptives may be at an increased risk of stroke. • d. Dose-Related Risk of Vascular Disease from Oral Contraceptives: A positive association has been observed between the amount of estrogen and progestogen in oral contraceptives and the risk of vascular disease. A decline in serum high-density lipoproteins (HDL) has been reported with many progestational agents. A decline in serum high-density lipoproteins has been associated with an increased incidence of ischemic heart disease. Because estrogens increase HDL cholesterol, the net effect of an oral contraceptive depends on a balance achieved between doses of estrogen and progestogen and the nature and absolute amount of progestogen used in the contraceptive. The amount of both hormones should be considered in the choice of an oral contraceptive. Minimizing exposure to estrogen and progestogen is in keeping with good principles of therapeutics. For any particular estrogen/progestogen combination, the dosage regimen prescribed should be one which contains the least amount of estrogen and progestogen that is compatible with a low failure rate and the needs of the individual patient. New acceptors of oral contraceptive agents should be started on preparations containing the lowest estrogen content, which is judged appropriate for the individual patient. • e. Persistence of Risk of Vascular Disease: There are two studies, which have shown persistence of risk of vascular disease for ever-users of oral contraceptives. In a study in the United States, the risk of developing myocardial infarction after discontinuing oral contraceptives persists for at least 9 years for women 40 to 49 years old who had used oral contraceptives for five or more years, but this increased risk was not demonstrated in other age groups. In another study in Great Britain, the risk of developing cerebrovascular disease persisted for at least 6 years after discontinuation of oral contraceptives, although excess risk was very small. However, both studies were performed with oral contraceptive formulations containing 50 micrograms or higher of estrogens.

2. Estimates of Mortality from Contraceptive Use: Each method of contraception has its specific benefits and risks. One study concluded that with the exception of oral contraceptive users 35 and older who smoke and 40 and older who do not smoke, mortality associated with all methods of birth control is less than that associated with childbirth. The observation of a possible increase in risk of mortality with age for oral contraceptive users is based on data gathered in the 1970's--but not reported until 1983. However, current clinical practice involves the use of lower estrogen dose formulations combined with careful restriction of oral contraceptive use to women who do not have the various risk factors listed in this labeling. Because of these changes in practice and, also, because of some limited new data which suggest that the risk of cardiovascular disease with the use of oral contraceptives may now be less than previously observed, the Fertility and Maternal Health Drugs Advisory Committee was asked to review the topic in 1989. The Committee concluded that although cardiovascular disease risks may be increased with oral contraceptive use after age 40 in healthy nonsmoking women (even with the newer low-dose formulations), there are greater potential health risks associated with pregnancy in older women and with the alternative surgical and medical procedures which may be necessary if such women do not have access to effective and acceptable means of contraception. Therefore, the Committee recommended that the benefits of oral contraceptive use by healthy nonsmoking women over 40 may outweigh the possible risks. Of course, older women, as all women who take oral contraceptives, should take the lowest possible dose formulation that is effective.

3. Carcinoma of the Reproductive Organs and Breasts: Although the risk of having breast cancer diagnosed may be slightly increased among current and recent users of combined oral contraceptives (RR=1.24), this excess risk decreases over time after combination oral contraceptive discontinuation and by 10 years after cessation the increased risk disappears. The risk does not increase with duration of use and no consistent relationships have been found with dose or type of steroid. The patterns of risk are also similar regardless of a woman's reproductive history or her family breast cancer history. The subgroup for whom risk has been found to be significantly elevated is women who first used oral contraceptives before age 20, but because breast cancer is so rare at these young ages, the number of cases attributable to this early oral contraceptive use is extremely small. Breast cancers diagnosed in current or previous oral contraceptive users tend to be less clinically advanced than in never-users. Women who currently have or have had breast cancer should not use oral contraceptives because breast cancer is a hormone sensitive tumor. Some studies suggest that oral contraceptive use has been associated with an increase in the risk of cervical intraepithelial neoplasia or invasive cervical cancer 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. In spite of many studies of the relationship between oral contraceptive use and breast cancer and cervical cancers, a cause-and-effect relationship has not been established.

4. Hepatic Neoplasia: Benign hepatic adenomas are associated with oral contraceptive use, although their occurrence is rare in the United States. Indirect calculations have estimated the attributable risk to be in the range of 3.3 cases/100,000 for users, a risk that increases after four or more years of use. Rupture of hepatic adenomas may cause death through intra-abdominal hemorrhage. Studies from Britain have shown an increased risk of developing hepatocellular carcinoma in long-term (>8 years) oral contraceptive users. However, these cancers are extremely rare in the U.S., and the attributable risk (the excess incidence) of liver cancers in oral contraceptive users approaches less than one per million users.

5. Ocular Lesions: There have been clinical case reports of retinal thrombosis associated with the use of oral contraceptives that may lead to partial or complete loss of vision. Oral contraceptives should be discontinued if there is unexplained partial or complete loss of vision; onset of proptosis or diplopia; papilledema; or retinal vascular lesions. Appropriate diagnostic and therapeutic measures should be undertaken immediately.

6. Oral Contraceptive Use Before or During Early Pregnancy: Because women using Seasonique® will likely have withdrawal bleeding only 4 times per year, pregnancy should be ruled out at the time of any missed menstrual period. Oral contraceptive use should be discontinued if pregnancy is confirmed. Extensive epidemiological studies have revealed no increased risk of birth defects in women who have used oral contraceptives prior to pregnancy. Studies also do not suggest a teratogenic effect, particularly in so far as cardiac anomalies and limb-reduction defects are concerned, when taken inadvertently during early pregnancy (see **CONTRAINDICATIONS**). The administration of oral contraceptives to induce withdrawal bleeding should not be used as a test for pregnancy. Oral contraceptives should not be used during pregnancy to treat threatened or habitual abortion.

7. Gallbladder Disease: Earlier studies have reported an increased lifetime relative risk of gallbladder surgery in users of oral contraceptives and estrogens. More recent studies, however, have shown that the relative risk of developing gallbladder disease among oral contraceptive users may be minimal. The recent findings of minimal risk may be related to the use of oral contraceptive formulations containing lower hormonal doses of estrogens and progestogens.

8. Carbohydrate and Lipid Metabolic Effects: Oral contraceptives have been shown to cause glucose intolerance in a significant percentage of users. Oral contraceptives containing greater than 75 micrograms of estrogens cause hyperinsulinism, while lower doses of estrogen cause less glucose intolerance. Progestogens increase insulin secretion and create insulin resistance, this effect varying with different progestational agents. However, in the nondiabetic woman, oral contraceptives appear to have no effect on fasting blood glucose. Because of these demonstrated effects, prediabetic and diabetic women should be carefully observed

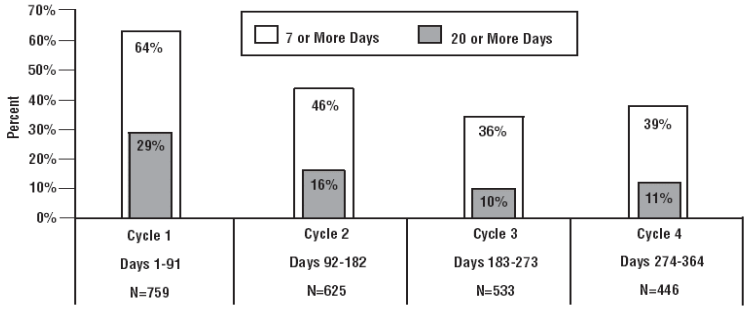
while taking oral contraceptives. A small proportion of women will have persistent hypertriglyceridemia while on the pill. As discussed earlier (see **WARNINGS** 1a. and 1d.), changes in serum triglycerides and lipoprotein levels have been reported in oral contraceptive users.

9. Elevated Blood Pressure: Women with significant hypertension should not be started on hormonal contraceptive. An increase in blood pressure has been reported in women taking oral contraceptives and this increase is more likely in older oral contraceptive users and with continued use. Data from the Royal College of General Practitioners and subsequent randomized trials have shown that the incidence of hypertension increases with increasing concentrations of progestogens. Women with a history of hypertension or hypertension-related diseases, or renal disease should be encouraged to use another method of contraception. If women with hypertension elect to use oral contraceptives, they should be monitored closely, and if significant elevation of blood pressure occurs, oral contraceptives should be discontinued (see **CONTRAINDICATIONS**). For most women, elevated blood pressure will return to normal after stopping oral contraceptives, and there is no difference in the occurrence of hypertension among ever- and never-users.

10. Headache: The onset or exacerbation of migraine or development of headache with a new pattern that is recurrent, persistent, or severe requires discontinuation of oral contraceptives and evaluation of the cause. (See **WARNINGS**, 1c.)

11. Bleeding Irregularities: When prescribing Seasonique® the convenience of fewer planned menses (4 per year instead of 13 per year) should be weighed against the inconvenience of increased intermenstrual bleeding and/or spotting. The primary clinical trial (PSE-301) that evaluated the efficacy of Seasonique® also assessed intermenstrual bleeding. The participants in the study (N=1,006) were composed primarily of women who had used oral contraceptives previously (89.3%) as opposed to new users (10.7%). A total of 82 (8.2%) of the women discontinued Seasonique®, at least in part, due to bleeding or spotting. The following figure shows the percentage of Seasonique® subjects participating in trial PSE-301 with $\geq$ 7 days or $\geq$ 20 days of intermenstrual bleeding or spotting during each treatment cycle. During the first 91 day treatment cycle, 64% of subjects experienced 7 or more days of intermenstrual bleeding or spotting with 29% of this cohort experiencing 20 or more days of intermenstrual bleeding or spotting. During the fourth 91-day treatment cycle, these percentages were 39% and 11%, respectively.

Figure: Percentage of Women Taking Seasonique® Reporting Intermenstrual Bleeding and/or Spotting.



As in any case of bleeding irregularities, nonhormonal causes should always be considered and adequate diagnostic measures taken to rule out malignancy or pregnancy. In the event of amenorrhea, pregnancy should be ruled out. Some women may encounter post-pill amenorrhea or oligomenorrhea (possibly with anovulation), especially when such a condition was preexistent.

PRECAUTIONS

1. Sexually Transmitted Diseases: 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: A periodic history and physical examination are appropriate for all women, including women using oral contraceptives. The physical examination, however, may be deferred until after initiation of oral contraceptives if requested by the woman and judged appropriate by the clinician. The physical examination should include special reference to blood pressure, breasts, abdomen and pelvic organs, including cervical cytology, and relevant laboratory tests. In case of undiagnosed, persistent or recurrent abnormal vaginal bleeding, appropriate diagnostic measures should be conducted to rule out malignancy. Women with a strong family history of breast cancer or who have breast nodules should be monitored with particular care.

3. Lipid Disorders: Women who are being treated for hyperlipidemias should be followed closely if they elect to use oral contraceptives. Some progestogens may elevate LDL levels and may render the control of hyperlipidemias more difficult. (See **WARNINGS** 1d.) In patients with familial defects of lipoprotein metabolism receiving estrogen-containing preparations, there have been case reports of significant elevations of plasma triglycerides leading to pancreatitis.

4. Liver Function: If jaundice develops in any woman receiving such drugs, the medication should be discontinued. Steroid hormones may be poorly metabolized in patients with impaired liver function.

5. Fluid Retention: Oral contraceptives may cause some degree of fluid retention. They should be prescribed with caution, and only with careful monitoring, in patients with conditions, which might be aggravated by fluid retention.

6. Emotional Disorders: Women with a history of depression should be carefully observed and the drug discontinued if depression recurs to a serious degree. Patients becoming significantly depressed while taking oral contraceptives should stop the medication and use an alternate method of contraception in an attempt to determine whether the symptom is drug related.

7. Contact Lenses: Contact-lens wearers who develop visual changes or changes in lens tolerance should be assessed by an ophthalmologist.

8. Drug Interactions: Changes in contraceptive effectiveness associated with co-administration of other products: • a. Anti-infective agents and anticonvulsants: Contraceptive effectiveness may be reduced when hormonal contraceptives are co-administered with antibiotics, anticonvulsants, and other drugs that increase the metabolism of contraceptive steroids. This could result in unintended pregnancy or breakthrough bleeding. Examples include rifampin, barbiturates, phenylbutazone, phenytoin, carbamazepine, felbamate, oxcarbazepine, topiramate, and griseofulvin. Several cases of contraceptive failure and breakthrough bleeding have been reported in the literature with concomitant administration of antibiotics such as ampicillin and tetracyclines. However, clinical pharmacology studies investigating drug interaction between combined oral contraceptives and these antibiotics have reported inconsistent results. • b. Anti-HIV protease inhibitors: Several of the anti-HIV protease inhibitors have been studied with co-administration of oral combination hormonal contraceptives; significant changes (increase and decrease) in the plasma levels of the estrogen and progestin have been noted in some cases. The safety and efficacy of combination oral contraceptive products may be affected with co-administration of anti-HIV protease inhibitors. Healthcare providers should refer to the label of the individual anti-HIV protease inhibitors for further drug-drug interaction information. • c. Herbal products: Herbal products containing St. John's Wort (hypericum perforatum) may induce hepatic enzymes (cytochrome P450) and p-glycoprotein transporter and may reduce the effectiveness of contraceptive steroids. This may also result in breakthrough bleeding. **Increase in plasma levels of estradiol associated with co-administered drugs:** Co-administration of atorvastatin and certain combination oral contraceptives containing ethinyl estradiol increase AUC values for ethinyl estradiol by approximately 20%. Ascorbic acid and acetaminophen may increase plasma ethinyl estradiol levels, possibly by inhibition of conjugation. CYP 3A4 inhibitors such as itraconazole or ketoconazole may increase plasma hormone levels. **Changes in plasma levels of co-administered drugs:** Combination hormonal contraceptives containing some synthetic estrogens (e.g., ethinyl estradiol) may inhibit the metabolism of other compounds. Increased plasma concentrations of cyclosporin, prednisolone, and theophylline have been reported with concomitant administration of combination oral contraceptives. Decreased plasma concentrations of acetaminophen and increased clearance of temazepam, salicylic acid, morphine and clofibric acid, due to induction of conjugation have been noted when these drugs were administered with combination oral contraceptives.

9. Interactions with Laboratory Tests - See Package Insert for complete information.

10. Carcinogenesis: See **WARNINGS**. **11. Pregnancy:** Pregnancy Category X. See **CONTRAINDICATIONS and WARNINGS**. **12. Nursing Mothers:** Small amounts of oral contraceptive steroids and/or metabolites have been identified in the milk of nursing mothers, and a few adverse effects on the child have been reported, including jaundice and breast enlargement. In addition, oral contraceptives given in the postpartum period may interfere with lactation by decreasing the quantity and quality of breast milk. If possible, the nursing mother should be advised not to use oral contraceptives but to use other forms of contraception until she has completely weaned her child. **13. Pediatric Use:** Safety and efficacy of Seasonique® tablets have been established in women of reproductive age. Safety and efficacy are expected to be the same in postpubertal adolescents under the age of 16 and users 16 and older. Use of Seasonique® before menarche is not indicated. **14. Geriatric Use:** Seasonique® tablets have not been studied in women who have reached menopause.

INFORMATION FOR THE PATIENT: See Package Brochure or complete information.

ADVERSE REACTIONS: An increased risk of the following serious adverse reactions has been associated with the use of oral contraceptives (see **WARNINGS**): • Thrombophlebitis • Arterial thromboembolism • Pulmonary embolism • Myocardial infarction • Cerebral hemorrhage • Cerebral thrombosis • Hypertension • Gallbladder disease • Hepatic adenomas or benign liver tumors. There is evidence of an association between the following conditions and the use of oral contraceptives: • Mesenteric thrombosis • Retinal thrombosis. The following adverse reactions have been reported in patients receiving oral contraceptives and are believed to be drug related: • Nausea • Vomiting • Gastrointestinal symptoms (such as abdominal cramps and bloating) • Breakthrough bleeding • Spotting • Change in menstrual flow • Amenorrhea • Temporary infertility after discontinuation of treatment • Edema/fluid retention • Melasma/chloasma which may persist • Breast changes: tenderness, enlargement, and secretion • Change in weight or appetite (increase or decrease) • Change in cervical ectropion and secretion • Possible diminution in lactation when given immediately postpartum • Cholestatic jaundice • Migraine headache • Rash (allergic) • Mood changes, including depression • Vaginitis, including candidiasis • Change in corneal curvature (steepening) • Intolerance to contact lenses • Decrease in serum folate levels • Exacerbation of systemic lupus erythematosus • Exacerbation of porphyria • Exacerbation of chorea • Aggravation of varicose veins • Anaphylactic/anaphylactoid reactions, including urticaria, angioedema, and severe reactions with respiratory and circulatory symptoms. The following adverse reactions have been reported in users of oral contraceptives and the association has been neither confirmed nor refuted: • Premenstrual syndrome • Cataracts • Optic neuritis which may lead to partial or complete loss of vision • Cystitis-like syndrome • Headache • Nervousness • Dizziness • Hirsutism • Loss of scalp hair • Erythema multiforme • Erythema nodosum • Hemorrhagic eruption • Impaired renal function • Hemolytic uremic syndrome • Budd-Chiari syndrome • Acne • Changes in libido • Colitis • Pancreatitis • Dysmenorrhea

OVERDOSAGE: Serious ill effects have not been reported following acute ingestion of large doses of oral contraceptives by young children. Overdosage may cause nausea, and withdrawal bleeding may occur in females.

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