

MEDROXYPROGESTERONE ACETATE- medroxyprogesterone acetate injection, suspension
Preferred Pharmaceuticals, Inc

HIGHLIGHTS OF PRESCRIBING INFORMATION

These highlights do not include all the information needed to use Medroxyprogesterone Acetate Injectable Suspension, USP safely and effectively. See full prescribing information for Medroxyprogesterone Acetate Injectable Suspension, USP.

Medroxyprogesterone Acetate Injectable Suspension, USP, for intramuscular use

Initial U.S. Approval: 1959

WARNING: LOSS OF BONE MINERAL DENSITY

See full prescribing information for complete boxed warning.

- Women who use Medroxyprogesterone Acetate Injectable Suspension, USP may lose significant bone mineral density. Bone loss is greater with increasing duration of use and may not be completely reversible. (5.1)
- It is unknown if use of Medroxyprogesterone Acetate Injectable Suspension, USP during adolescence or early adulthood, a critical period of bone accretion, will reduce peak bone mass and increase the risk for osteoporotic fracture in later life. (5.1)
- Medroxyprogesterone Acetate Injectable Suspension, USP should not be used as a long-term birth control method (i.e., longer than 2 years) unless other birth control methods are considered inadequate. (5.1)

----- **RECENT MAJOR CHANGES** -----

| | |
|--|---------|
| Boxed Warning: Loss of Bone Mineral Density | 10/2010 |
| Warnings and Precautions; Loss of Bone Mineral Density (5.1) | 10/2010 |

----- **INDICATIONS AND USAGE** -----

- Medroxyprogesterone Acetate Injectable Suspension, USP is a progestin injectable contraceptive indicated only for the prevention of pregnancy. (1)

----- **DOSAGE AND ADMINISTRATION** -----

- The recommended dose is 150 mg of Medroxyprogesterone Acetate Injectable Suspension, USP every 3 months (13 weeks) administered by deep, intramuscular (IM) injection in the gluteal or deltoid muscle. (2.1)

----- **DOSAGE FORMS AND STRENGTHS** -----

- Vials containing sterile aqueous suspension: 150 mg per mL (3)
- Prefilled syringes: prefilled syringes are available packaged with 22-gauge × 1 1/2 inch BD SafetyGlide Needles (3)

----- **CONTRAINDICATIONS** -----

- Known or suspected pregnancy or as a diagnostic test for pregnancy. (4)
- Active thrombophlebitis, or current or past history of thromboembolic disorders, or cerebral vascular disease. (4)
- Known or suspected malignancy of breast. (4)
- Known hypersensitivity to Medroxyprogesterone Acetate Injectable Suspension, USP or any of its ingredients. (4)
- Significant liver disease. (4)
- Undiagnosed vaginal bleeding. (4)

----- **WARNINGS AND PRECAUTIONS** -----

- Thromboembolic Disorders: Discontinue Medroxyprogesterone Acetate Injectable Suspension, USP in patients who develop thrombosis (5.2)
- Cancer Risks: Monitor women with breast nodules or a strong family history of breast cancer carefully. (5.3)
- Ectopic Pregnancy: Consider ectopic pregnancy if a woman using Medroxyprogesterone Acetate Injectable Suspension, USP becomes pregnant or complains of severe abdominal pain. (5.4)
- Anaphylaxis and Anaphylactoid Reactions: Provide emergency medical treatment. (5.5)
- Liver Function: Discontinue Medroxyprogesterone Acetate Injectable Suspension, USP if jaundice or disturbances of

liver function develop (5.6)

- Carbohydrate Metabolism: Monitor diabetic patients carefully. (5.11)

----- **ADVERSE REACTIONS** -----

Most common adverse reactions (incidence >5%) are: menstrual irregularities (bleeding or spotting) 57% at 12 months, 32% at 24 months, abdominal pain/discomfort 11%, weight gain > 10 lbs at 24 months 38%, dizziness 6%, headache 17%, nervousness 11%, decreased libido 6%. (6.1)

To report SUSPECTED ADVERSE REACTIONS, contact Pfizer Inc. at 1-800-438-1985 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

----- **DRUG INTERACTIONS** -----

Drugs or herbal products that induce certain enzymes, including CYP3A4, may decrease the effectiveness of contraceptive drug products. Counsel patients to use a back-up method or alternative method of contraception when enzyme inducers are used with Medroxyprogesterone Acetate Injectable Suspension, USP. (7.1)

----- **USE IN SPECIFIC POPULATIONS** -----

- *Nursing Mothers*: Detectable amounts of drug have been identified in the milk of mothers receiving Medroxyprogesterone Acetate Injectable Suspension, USP. (8.3)
- *Pediatric Patients*: Medroxyprogesterone Acetate Injectable Suspension, USP is not indicated before menarche. (8.4)

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

Revised: 9/2011

FULL PRESCRIBING INFORMATION: CONTENTS*

WARNING: LOSS OF BONE MINERAL DENSITY

1 INDICATIONS AND USAGE

2 DOSAGE AND ADMINISTRATION

2.1 Prevention of Pregnancy

2.2 Switching from other Methods of Contraception

3 DOSAGE FORMS AND STRENGTHS

4 CONTRAINDICATIONS

5 WARNINGS AND PRECAUTIONS

5.1 Loss of Bone Mineral Density

5.2 Thromboembolic Disorders

5.3 Cancer Risks

5.4 Ectopic Pregnancy

5.5 Anaphylaxis and Anaphylactoid Reaction

5.6 Liver Function

5.7 Convulsions

5.8 Depression

5.9 Bleeding Irregularities

5.10 Weight Gain

5.11 Carbohydrate Metabolism

5.12 Lactation

5.13 Fluid Retention

5.14 Return of Fertility

5.15 Sexually Transmitted Diseases

5.16 Pregnancy

5.17 Monitoring

5.18 Interference with Laboratory Tests

6 ADVERSE REACTIONS

6.1 Clinical Trials Experience

6.2 Post-marketing Experience

7 DRUG INTERACTIONS

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

7.2 Laboratory Test Interactions

8 USE IN SPECIFIC POPULATIONS

8.1 Pregnancy

8.3 Nursing Mothers

8.4 Pediatric Use

8.5 Geriatric Use

8.6 Renal Impairment

8.7 Hepatic Impairment

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 Contraception

14.2 Bone Mineral Density (BMD) Changes in Adult Women

14.3 Bone Mineral Density Changes in Adolescent Females (12–18 years of age)

14.4 Relationship of fracture incidence to use of Medroxyprogesterone Acetate Injectable Suspension, USP or non-use by women of reproductive age

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

WARNING: LOSS OF BONE MINERAL DENSITY

Women who use Medroxyprogesterone Acetate Injectable Suspension, USP may lose significant bone mineral density. Bone loss is greater with increasing duration of use and may not be completely reversible.

It is unknown if use of Medroxyprogesterone Acetate Injectable Suspension, USP during adolescence or early adulthood, a critical period of bone accretion, will reduce peak bone mass and increase the risk for osteoporotic fracture in later life.

Medroxyprogesterone Acetate Injectable Suspension, USP should not be used as a long-term birth control method (i.e., longer than 2 years) unless other birth control methods are considered inadequate. (See *Warnings and Precautions (5.1)*).

1 INDICATIONS AND USAGE

Medroxyprogesterone Acetate Injectable Suspension, USP is indicated only for the prevention of pregnancy. The loss of bone mineral density (BMD) in women of all ages and the impact on peak bone mass in adolescents should be considered, along with the decrease in BMD that occurs during pregnancy and/or lactation, in the risk/benefit assessment for women who use Medroxyprogesterone Acetate Injectable Suspension, USP long-term [see *Warnings and Precautions (5.1)*].

2 DOSAGE AND ADMINISTRATION

2.1 Prevention of Pregnancy

Both the 1 mL vial and the 1 mL prefilled syringe of Medroxyprogesterone Acetate Injectable Suspension, USP should be vigorously shaken just before use to ensure that the dose being administered represents a uniform suspension.

The recommended dose is 150 mg of Medroxyprogesterone Acetate Injectable Suspension, USP every 3 months (13 weeks) administered by deep IM injection in the gluteal or deltoid muscle.

Medroxyprogesterone Acetate Injectable Suspension, USP should not be used as a long-term birth control method (i.e. longer than 2 years) unless other birth control methods are considered inadequate. Dosage does not need to be adjusted for body weight [See *Clinical Studies (14.1)*].

To ensure the patient is not pregnant at the time of the first injection, the first injection should be given ONLY during the first 5 days of a normal menstrual period; ONLY within the first 5-days postpartum if not breast-feeding; and if exclusively breast-feeding, ONLY at the sixth postpartum week. If the time interval between injections is greater than 13 weeks, the physician should determine that the patient is not pregnant before administering the drug. The efficacy of Medroxyprogesterone Acetate Injectable Suspension, USP depends on adherence to the dosage schedule of administration.

2.2 Switching from other Methods of Contraception

When switching from other contraceptive methods, Medroxyprogesterone Acetate Injectable Suspension, USP should be given in a manner that ensures continuous contraceptive coverage based upon the mechanism of action of both methods, (e.g., patients switching from oral contraceptives should have their first injection of Medroxyprogesterone Acetate Injectable Suspension, USP on the day after the last active tablet or at the latest, on the day following the final inactive tablet).

3 DOSAGE FORMS AND STRENGTHS

Sterile Aqueous suspension: 150mg/ml

Prefilled syringes are available packaged with 22-gauge × 1 1/2 inch BD SafetyGlide™ Needles.

4 CONTRAINDICATIONS

The use of Medroxyprogesterone Acetate Injectable Suspension, USP is contraindicated in the following conditions:

- Known or suspected pregnancy or as a diagnostic test for pregnancy.
- Active thrombophlebitis, or current or past history of thromboembolic disorders, or cerebral vascular disease [see *Warnings and Precautions (5.2)*].
- Known or suspected malignancy of breast [see *Warnings and Precautions (5.3)*].
- Known hypersensitivity to Medroxyprogesterone Acetate Injectable Suspension, USP (medroxyprogesterone acetate) or any of its other ingredients [see *Warnings and Precautions (5.5)*].
- Significant liver disease [see *Warnings and Precautions (5.6)*].
- Undiagnosed vaginal bleeding [see *Warnings and Precautions (5.9)*].

5 WARNINGS AND PRECAUTIONS

5.1 Loss of Bone Mineral Density

Use of Medroxyprogesterone Acetate Injectable Suspension, USP reduces serum estrogen levels and is associated with significant loss of bone mineral density (BMD). This loss of BMD is of particular concern during adolescence and early adulthood, a critical period of bone accretion. It is unknown if use of Medroxyprogesterone Acetate Injectable Suspension, USP by younger women will reduce peak bone mass and increase the risk for osteoporotic fracture in later life.

[After discontinuing Medroxyprogesterone Acetate Injectable Suspension, USP in adolescents, mean

BMD loss at total hip and femoral neck did not fully recover by 60 months (240 weeks) post-treatment [see *Clinical Studies (14.3)*]. Similarly, in adults, there was only partial recovery of mean BMD at total hip, femoral neck and lumbar spine towards baseline by 24 months post-treatment. [See *Clinical Studies (14.2)*.]

Medroxyprogesterone Acetate Injectable Suspension, USP should not be used as a long-term birth control method (i.e., longer than 2 years) unless other birth control methods are considered inadequate. BMD should be evaluated when a woman needs to continue to use Medroxyprogesterone Acetate Injectable Suspension, USP long-term. In adolescents, interpretation of BMD results should take into account patient age and skeletal maturity.

Other birth control methods should be considered in the risk/benefit analysis for the use of Medroxyprogesterone Acetate Injectable Suspension, USP in women with osteoporosis risk factors. Medroxyprogesterone Acetate Injectable Suspension, USP can pose an additional risk in patients with risk factors for osteoporosis (e.g., metabolic bone disease, chronic alcohol and/or tobacco use, anorexia nervosa, strong family history of osteoporosis or chronic use of drugs that can reduce bone mass such as anticonvulsants or corticosteroids). Although there are no studies addressing whether calcium and Vitamin D may lessen BMD loss in women using Medroxyprogesterone Acetate Injectable Suspension, USP, all patients should have adequate calcium and Vitamin D intake.

5.2 Thromboembolic Disorders

There have been reports of serious thrombotic events in women using Medroxyprogesterone Acetate Injectable Suspension, USP. However, Medroxyprogesterone Acetate Injectable Suspension, USP has not been causally associated with the induction of thrombotic or thromboembolic disorders. Any patient who develops thrombosis while undergoing therapy with Medroxyprogesterone Acetate Injectable Suspension, USP should discontinue treatment unless she has no other acceptable options for birth control.

Do not re-administer Medroxyprogesterone Acetate Injectable Suspension, USP pending examination if there is a sudden partial or complete loss of vision or if there is a sudden onset of proptosis, diplopia, or migraine. Do not re-administer if examination reveals papilledema or retinal vascular lesions.

5.3 Cancer Risks

Breast Cancer

Women who currently have or have had breast cancer should not use hormone contraceptives, including Medroxyprogesterone Acetate Injectable Suspension, USP, because breast cancer may be hormonally sensitive. Women with a strong family history of breast cancer or who have breast nodules should be monitored with particular care.

A pooled analysis from two case-control studies, the World Health Organization Study and the New Zealand Study, reported the relative risk (RR) of breast cancer for women who had ever used Medroxyprogesterone Acetate Injectable Suspension, USP as 1.1 (95% confidence interval [CI] 0.97 to 1.4). Overall, there was no increase in risk with increasing duration of use of Medroxyprogesterone Acetate Injectable Suspension, USP. The RR of breast cancer for women of all ages who had initiated use of Medroxyprogesterone Acetate Injectable Suspension, USP within the previous 5 years was estimated to be 2.0 (95% CI 1.5 to 2.8).

The World Health Organization Study, a component of the pooled analysis described above, showed an increased RR of 2.19 (95% CI 1.23 to 3.89) of breast cancer associated with use of Medroxyprogesterone Acetate Injectable Suspension, USP in women whose first exposure to drug was within the previous 4 years and who were under 35 years of age. However, the overall RR for ever-users of Medroxyprogesterone Acetate Injectable Suspension, USP was 1.2 (95% CI 0.96 to 1.52).

The National Cancer Institute reports an average annual incidence rate for breast cancer for US women, all races, age 15 to 34 years of 8.7 per 100,000. A RR of 2.19, thus, increases the possible risk from 8.7 to 19.0 cases per 100,000 women. The attributable risk, thus, is 31.8 per 100,000 women per year.

Cervical Cancer

A statistically nonsignificant increase in RR estimates of invasive squamous-cell cervical cancer has been associated with the use of Medroxyprogesterone Acetate Injectable Suspension, USP in women who were first exposed before the age of 35 years (RR 1.22 to 1.28 and 95% CI 0.93 to 1.70). The overall, nonsignificant relative rate of invasive squamous-cell cervical cancer in women who ever used Medroxyprogesterone Acetate Injectable Suspension, USP was estimated to be 1.11 (95% CI 0.96 to 1.29). No trends in risk with duration of use or times since initial or most recent exposure were observed.

Other Cancers

Long-term case-controlled surveillance of users of Medroxyprogesterone Acetate Injectable Suspension, USP found no overall increased risk of ovarian or liver cancer.

5.4 Ectopic Pregnancy

Be alert to the possibility of an ectopic pregnancy among women using Medroxyprogesterone Acetate Injectable Suspension, USP who become pregnant or complain of severe abdominal pain.

5.5 Anaphylaxis and Anaphylactoid Reaction

Anaphylaxis and anaphylactoid reaction have been reported with the use of Medroxyprogesterone Acetate Injectable Suspension, USP. Institute emergency medical treatment if an anaphylactic reaction occurs.

5.6 Liver Function

Discontinue Medroxyprogesterone Acetate Injectable Suspension, USP use if jaundice or acute or chronic disturbances of liver function develop. Do not resume use until markers of liver function return to normal and Medroxyprogesterone Acetate Injectable Suspension, USP causation has been excluded.

5.7 Convulsions

There have been a few reported cases of convulsions in patients who were treated with Medroxyprogesterone Acetate Injectable Suspension, USP. Association with drug use or pre-existing conditions is not clear.

5.8 Depression

Monitor patients who have a history of depression and do not readminister Medroxyprogesterone Acetate Injectable Suspension, USP if depression recurs.

5.9 Bleeding Irregularities

Most women using Medroxyprogesterone Acetate Injectable Suspension, USP experience disruption of menstrual bleeding patterns. Altered menstrual bleeding patterns include amenorrhea, irregular or unpredictable bleeding or spotting, prolonged spotting or bleeding, and heavy bleeding. Rule out the possibility of organic pathology if abnormal bleeding persists or is severe, and institute appropriate treatment.

As women continue using Medroxyprogesterone Acetate Injectable Suspension, USP, fewer experience irregular bleeding and more experience amenorrhea. In clinical studies of Medroxyprogesterone Acetate Injectable Suspension, USP, by month 12 amenorrhea was reported by 55% of women, and by month 24, amenorrhea was reported by 68% of women using Medroxyprogesterone Acetate Injectable Suspension, USP.

5.10 Weight Gain

Women tend to gain weight while on therapy with Medroxyprogesterone Acetate Injectable Suspension, USP. From an initial average body weight of 136 lb, women who completed 1 year of therapy with

Medroxyprogesterone Acetate Injectable Suspension, USP gained an average of 5.4 lb. Women who completed 2 years of therapy gained an average of 8.1 lb. Women who completed 4 years gained an average of 13.8 lb. Women who completed 6 years gained an average of 16.5 lb. Two percent of women withdrew from a large-scale clinical trial because of excessive weight gain.

5.11 Carbohydrate Metabolism

A decrease in glucose tolerance has been observed in some patients on Medroxyprogesterone Acetate Injectable Suspension, USP treatment. Monitor diabetic patients carefully while receiving Medroxyprogesterone Acetate Injectable Suspension, USP.

5.12 Lactation

Detectable amounts of drug have been identified in the milk of mothers receiving Medroxyprogesterone Acetate Injectable Suspension, USP. In nursing mothers treated with Medroxyprogesterone Acetate Injectable Suspension, USP, milk composition, quality, and amount are not adversely affected. Neonates and infants exposed to medroxyprogesterone from breast milk have been studied for developmental and behavioral effects through puberty. No adverse effects have been noted.

5.13 Fluid Retention

Because progestational drugs including Medroxyprogesterone Acetate Injectable Suspension, USP may cause some degree of fluid retention, monitor patients with conditions that might be influenced by this condition, such as epilepsy, migraine, asthma, and cardiac or renal dysfunction.

5.14 Return of Fertility

Return to ovulation and fertility is likely to be delayed after stopping Medroxyprogesterone Acetate Injectable Suspension, USP. In a large US study of women who discontinued use of Medroxyprogesterone Acetate Injectable Suspension, USP to become pregnant, data are available for 61% of them. Of the 188 women who discontinued the study to become pregnant, 114 became pregnant. Based on Life-Table analysis of these data, it is expected that 68% of women who do become pregnant may conceive within 12 months, 83% may conceive within 15 months, and 93% may conceive within 18 months from the last injection. The median time to conception for those who do conceive is 10 months following the last injection with a range of 4 to 31 months, and is unrelated to the duration of use. No data are available for 39% of the patients who discontinued Medroxyprogesterone Acetate Injectable Suspension, USP to become pregnant and who were lost to follow-up or changed their mind.

5.15 Sexually Transmitted Diseases

Patients should be counseled that Medroxyprogesterone Acetate Injectable Suspension, USP does not protect against HIV infection (AIDS) and other sexually transmitted diseases.

5.16 Pregnancy

Although Medroxyprogesterone Acetate Injectable Suspension, USP should not be used during pregnancy, there appears to be little or no increased risk of birth defects in women who have inadvertently been exposed to medroxyprogesterone acetate injections in early pregnancy. Neonates exposed to medroxyprogesterone acetate in-utero and followed to adolescence showed no evidence of any adverse effects on their health including their physical, intellectual, sexual or social development.

5.17 Monitoring

A woman who is taking hormonal contraceptive should have a yearly visit with her healthcare provider for a blood pressure check and for other indicated healthcare.

5.18 Interference with Laboratory Tests

The use of Medroxyprogesterone Acetate Injectable Suspension, USP may change the results of some laboratory tests, such as coagulation factors, lipids, glucose tolerance, and binding proteins. *[See Drug*

Interactions (7.2)].

6 ADVERSE REACTIONS

The following important adverse reactions observed with the use of Medroxyprogesterone Acetate Injectable Suspension, USP are discussed in greater detail in the *Warnings and Precautions* section (5):

- Loss of Bone Mineral Density [see *Warnings and Precautions* (5.1)]
- Thromboembolic disease [see *Warnings and Precautions* (5.2)]
- Breast Cancer [see *Warnings and Precautions* (5.3)]
- Anaphylaxis and Anaphylactoid Reactions [see *Warnings and Precautions* (5.5)]
- Bleeding Irregularities [see *Warnings and Precautions* (5.9)]
- Weight Gain [see *Warnings and Precautions* (5.10)]

6.1 Clinical Trials Experience

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

In the two clinical trials with Medroxyprogesterone Acetate Injectable Suspension, USP, over 3,900 women, who were treated for up to 7 years, reported the following adverse reactions, which may or may not be related to the use of Medroxyprogesterone Acetate Injectable Suspension, USP. The population studied ranges in age from 15 to 51 years, of which 46% were White, 50% Non-White, and 4.9% Unknown race. The patients received 150 mg Medroxyprogesterone Acetate Injectable Suspension, USP every 3-months (90 days). The median study duration was 13 months with a range of 1–84 months. Fifty eight percent of patients remained in the study after 13 months and 34% after 24 months.

Table 1 Adverse Reactions that Were Reported by More than 5% of Subjects

| Body System* | Adverse Reactions (Incidence (%)) |
|-----------------------|---|
| Body as a Whole | Headache (16.5%) Abdominal pain/discomfort (11.2%) |
| Metabolic/Nutritional | Increased weight > 10lbs at 24 months (37.7%) |
| Nervous | Nervousness (10.8%) Dizziness (5.6%) Libido decreased (5.5%) |
| Urogenital | Menstrual irregularities: (bleeding (57.3% at 12 months, 32.1% at 24 months) amenorrhea (55% at 12 months, 68% at 24 months) |

* Body System represented from COSTART medical dictionary.

Table 2 Adverse Reactions that Were Reported by between 1 and 5% of Subjects

| Body System* | Adverse Reactions (Incidence (%)) |
|-----------------------|---|
| Body as a Whole | Asthenia/fatigue (4.2%) Backache (2.2%) Dysmenorrhea (1.7%) Hot flashes (1.0%) |
| Digestive | Nausea (3.3%) Bloating (2.3%) |
| Metabolic/Nutritional | Edema (2.2%) |
| Musculoskeletal | Leg cramps (3.7%) |

| | |
|----------------------------|--|
| Musculoskeletal | Arthralgia (1.0%) |
| Nervous | Depression (1.5%) Insomnia (1.0%) |
| Skin and Appendages | Acne (1.2%) No hair growth/alopecia (1.1%) Rash (1.1%) |
| Urogenital | Leukorrhea (2.9%) Breast pain (2.8%) Vaginitis (1.2%) |

* Body System represented from COSTART medical dictionary.

Adverse reactions leading to study discontinuation in $\geq 2\%$ of subjects: bleeding (8.2%), amenorrhea (2.1%), weight gain (2.0%)

6.2 Post-marketing Experience

The following adverse reactions have been identified during post approval use of Medroxyprogesterone Acetate Injectable Suspension, USP. 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.

There have been cases of osteoporosis including osteoporotic fractures reported post-marketing in patients taking Medroxyprogesterone Acetate Injectable Suspension, USP.

Table 3 Adverse Reactions Reported during Post-Marketing Experience

| Body System* | Adverse Reactions |
|----------------------------------|---|
| Body as a Whole | Chest pain, Allergic reactions, Fever, Pain at injection site, Chills, Axillary swelling |
| Cardiovascular | Syncope, Tachycardia, Thrombophlebitis, Deep vein thrombosis, Pulmonary embolus, Varicose veins |
| Digestive | Changes in appetite, Gastrointestinal disturbances, Jaundice, Excessive thirst, Rectal bleeding |
| Hematologic and Lymphatic | Anemia, Blood dyscrasia |
| Musculoskeletal | Osteoporosis |
| Nervous | Paralysis, Facial palsy, Paresthesia, Drowsiness |
| Respiratory | Dyspnea and asthma, Hoarseness |
| Skin and Appendages | Hirsutism, Excessive sweating and body odor, Dry skin, Scleroderma |
| Urogenital | Cervical cancer, Breast cancer, Lack of return to fertility, Unexpected pregnancy, Prevention of lactation, Changes in breast size, Breast lumps or nipple bleeding, Galactorrhea, Melasma, Chloasma, Increased libido, Uterine hyperplasia, Genitourinary infections, Vaginal cysts, Dyspareunia |

* Body System represented from COSTART medical dictionary.

7 DRUG INTERACTIONS

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

If a woman on hormonal contraceptives takes a drug or herbal product that induces enzymes, including CYP3A4, that metabolize contraceptive hormones, counsel her to use additional contraception or a

different method of contraception. Drugs or herbal products that induce such enzymes may decrease the plasma concentrations of contraceptive hormones, and may decrease the effectiveness of hormonal contraceptives. Some drugs or herbal products that may decrease the effectiveness of hormonal contraceptives include:

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

HIV protease inhibitors and non-nucleoside reverse transcriptase inhibitors: Significant changes (increase or decrease) in the plasma levels of progestin have been noted in some cases of co-administration of HIV protease inhibitors. Significant changes (increase or decrease) in the plasma levels of the progestin have been noted in some cases of co-administration with non-nucleoside reverse transcriptase inhibitors.

Antibiotics: There have been reports of pregnancy while taking hormonal contraceptives and antibiotics, but clinical pharmacokinetic studies have not shown consistent effects of antibiotics on plasma concentrations of synthetic steroids.

Consult the labeling of all concurrently-used drugs to obtain further information about interactions with hormonal contraceptives or the potential for enzyme alterations.

7.2 Laboratory Test Interactions

The pathologist should be advised of progestin therapy when relevant specimens are submitted.

The following laboratory tests may be affected by progestins including Medroxyprogesterone Acetate Injectable Suspension, USP:

- (a) Plasma and urinary steroid levels are decreased (e.g., progesterone, estradiol, pregnanediol, testosterone, cortisol).
- (b) Gonadotropin levels are decreased.
- (c) Sex-hormone-binding-globulin concentrations are decreased.
- (d) Protein-bound iodine and butanol extractable protein-bound iodine may increase. T₃-uptake values may decrease.
- (e) Coagulation test values for prothrombin (Factor II), and Factors VII, VIII, IX, and X may increase.
- (f) Sulfbromophthalein and other liver function test values may be increased.
- (g) The effects of medroxyprogesterone acetate on lipid metabolism are inconsistent. Both increases and decreases in total cholesterol, triglycerides, low-density lipoprotein (LDL) cholesterol, and high-density lipoprotein (HDL) cholesterol have been observed in studies.

8 USE IN SPECIFIC POPULATIONS

8.1 Pregnancy

Medroxyprogesterone Acetate Injectable Suspension, USP should not be administered during pregnancy. [See *Contraindications and Warnings and Precautions (5.16)*.]

8.3 Nursing Mothers

Detectable amounts of drug have been identified in the milk of mothers receiving Medroxyprogesterone Acetate Injectable Suspension, USP. [See *Warnings and Precautions* (5.12).]

8.4 Pediatric Use

Medroxyprogesterone Acetate Injectable Suspension, USP is not indicated before menarche. Use of Medroxyprogesterone Acetate Injectable Suspension, USP is associated with significant loss of BMD. This loss of BMD is of particular concern during adolescence and early adulthood, a critical period of bone accretion. In adolescents, interpretation of BMD results should take into account patient age and skeletal maturity. It is unknown if use of Medroxyprogesterone Acetate Injectable Suspension, USP by younger women will reduce peak bone mass and increase the risk of osteoporotic fractures in later life. Other than concerns about loss of BMD, the safety and effectiveness are expected to be the same for postmenarchal adolescents and adult women.

8.5 Geriatric Use

This product has not been studied in post-menopausal women and is not indicated in this population.

8.6 Renal Impairment

The effect of renal impairment on Medroxyprogesterone Acetate Injectable Suspension, USP pharmacokinetics has not been studied.

8.7 Hepatic Impairment

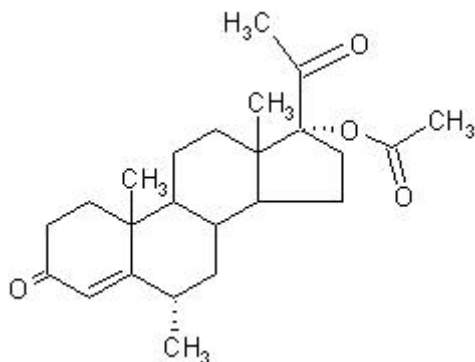
The effect of hepatic impairment on Medroxyprogesterone Acetate Injectable Suspension, USP pharmacokinetics has not been studied. Medroxyprogesterone Acetate Injectable Suspension, USP should not be used by women with significant liver disease and should be discontinued if jaundice or disturbances of liver function occur. [See *Contraindications* (4) and *Warnings and Precautions* (5.6).]

11 DESCRIPTION

Medroxyprogesterone Acetate Injectable Suspension, USP contains medroxyprogesterone acetate, a derivative of progesterone, as its active ingredient. Medroxyprogesterone acetate is active by the parenteral and oral routes of administration. It is a white to off-white; odorless crystalline powder that is stable in air and that melts between 200°C and 210°C. It is freely soluble in chloroform, soluble in acetone and dioxane, sparingly soluble in alcohol and methanol, slightly soluble in ether, and insoluble in water.

The chemical name for Medroxyprogesterone Acetate is pregn-4-ene-3,20-dione, 17-(acetyloxy)-6-methyl-, (6 α -).

The structural formula is as follows:



Medroxyprogesterone Acetate Injectable Suspension, USP for intramuscular (IM) injection is available

in vials and prefilled syringes, each containing 1 mL of medroxyprogesterone acetate sterile aqueous suspension 150 mg/mL.

Each mL contains:

| | |
|-----------------------------|---------------------|
| Medroxyprogesterone acetate | 150 mg |
| Polyethylene glycol 3350 | 28.9 mg |
| Polysorbate 80 | 2.41 mg |
| Sodium chloride | 8.68 mg |
| Methylparaben | 1.37 mg |
| Propylparaben | 0.150 mg |
| Water for injection | quantity sufficient |

When necessary, pH is adjusted with sodium hydroxide or hydrochloric acid, or both.

12 CLINICAL PHARMACOLOGY

12.1 Mechanism of Action

Medroxyprogesterone Acetate Injectable Suspension, USP, when administered at the recommended dose to women every 3 months, inhibits the secretion of gonadotropins which, in turn, prevents follicular maturation and ovulation and results in endometrial thinning. These actions produce its contraceptive effect.

12.2 Pharmacodynamics

No specific pharmacodynamic studies were conducted with Medroxyprogesterone Acetate Injectable Suspension, USP.

12.3 Pharmacokinetics

Absorption

Following a single 150 mg IM dose of Medroxyprogesterone Acetate Injectable Suspension, USP in eight women between the ages of 28 and 36 years old, medroxyprogesterone acetate concentrations, measured by an extracted radioimmunoassay procedure, increase for approximately 3 weeks to reach peak plasma concentrations of 1 to 7 ng/mL.

Distribution

Plasma protein binding of MPA averages 86%. MPA binding occurs primarily to serum albumin. No binding of MPA occurs with sex-hormone-binding globulin (SHBG).

Metabolism

MPA is extensively metabolized in the liver by P450 enzymes. Its metabolism primarily involves ring A and/or side-chain reduction, loss of the acetyl group, hydroxylation in the 2-, 6-, and 21-positions or a combination of these positions, resulting in more than 10 metabolites.

Excretion

The concentrations of medroxyprogesterone acetate decrease exponentially until they become undetectable (<100 pg/mL) between 120 to 200 days following injection. Using an unextracted radioimmunoassay procedure for the assay of medroxyprogesterone acetate in serum, the apparent half-life for medroxyprogesterone acetate following IM administration of Medroxyprogesterone Acetate Injectable Suspension, USP is approximately 50 days. Most medroxyprogesterone acetate metabolites are excreted in the urine as glucuronide conjugates with only minor amounts excreted as sulfates.

Specific Populations

The effect of hepatic and/or renal impairment on the pharmacokinetics of Medroxyprogesterone Acetate Injectable Suspension, USP is unknown.

13 NONCLINICAL TOXICOLOGY

13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility

[See Warnings and Precautions, (5.3, 5.14, and 5.16).]

14 CLINICAL STUDIES

14.1 Contraception

In five clinical studies using Medroxyprogesterone Acetate Injectable Suspension, USP, the 12-month failure rate for the group of women treated with Medroxyprogesterone Acetate Injectable Suspension, USP was zero (no pregnancies reported) to 0.7 by Life-Table method. The effectiveness of Medroxyprogesterone Acetate Injectable Suspension, USP is dependent on the patient returning every 3 months (13 weeks) for reinjection.

14.2 Bone Mineral Density (BMD) Changes in Adult Women

In a controlled, clinical study, adult women using Medroxyprogesterone Acetate Injectable Suspension, USP for up to 5 years showed spine and hip BMD mean decreases of 5–6%, compared to no significant change in BMD in the control group. The decline in BMD was more pronounced during the first two years of use, with smaller declines in subsequent years. Mean changes in lumbar spine BMD of -2.86%, -4.11%, -4.89%, -4.93% and -5.38% after 1, 2, 3, 4, and 5 years, respectively, were observed. Mean decreases in BMD of the total hip and femoral neck were similar.

After stopping use of Medroxyprogesterone Acetate Injectable Suspension, USP (150 mg), there was partial recovery of BMD toward baseline values during the 2-year post-therapy period. Longer duration of treatment was associated with less complete recovery during this 2-year period following the last injection. Table 4 shows the change in BMD in women after 5 years of treatment with Medroxyprogesterone Acetate Injectable Suspension, USP and in women in a control group, as well as the extent of recovery of BMD for the subset of the women for whom 2-year post treatment data were available.

Table 4. Mean Percent Change from Baseline in BMD in Adults by Skeletal Site and Cohort (5 Years of Treatment and 2 Years of Follow-Up)

| Time in Study | Spine | | Total Hip | | Femoral Neck | |
|---------------|------------------------------|-----------------|------------------------------|----------------|------------------------------|------------------|
| | Medroxyprogesterone Acetate* | Control† | Medroxyprogesterone Acetate* | Control† | Medroxyprogesterone Acetate* | Control† |
| 5 years | -5.38 % n=33 | 0.43 % n=105 | -5.16 % n=21 | 0.19 % n=65 | -6.12 % n=34 | -0.27 % n=106 |
| 7 years | -3.13 % n=12 | 0.53 % n=60 | -1.34 % n=7 | 0.94 % n=39 | -5.38 % n=13 | -0.11 % n=63 |

* The treatment group consisted of women who received Medroxyprogesterone Acetate Injectable Suspension, USP for 5 years and were then followed for 2 years post-use (total time in study of 7 years).

† The control group consisted of women who did not use hormonal contraception and were followed for 7 years.

14.3 Bone Mineral Density Changes in Adolescent Females (12–18 years of age)

The impact of Medroxyprogesterone Acetate Injectable Suspension, USP (150 mg) use for up to 240 weeks (4.6 years) was evaluated in an open-label non-randomized clinical study in 389 adolescent

females (12–18 years). Use of Medroxyprogesterone Acetate Injectable Suspension, USP was associated with a significant decline from baseline in BMD.

Partway through the trial, drug administration was stopped (at 120 weeks). The mean number of injections per Medroxyprogesterone Acetate Injectable Suspension, USP user was 9.3. The decline in BMD at total hip and femoral neck was greater with longer duration of use (see Table 5). The mean decrease in BMD at 240 weeks was more pronounced at total hip (-6.4%) and femoral neck (-5.4%) compared to lumbar spine (-2.1%).

In general, adolescents increase bone density during the period of growth following menarche, as seen in the untreated cohort. However, the two cohorts were not matched at baseline for age, gynecologic age, race, BMD and other factors that influence the rate of acquisition of bone mineral density.

Table 5. Mean Percent Change from Baseline in BMD in Adolescents Receiving ≥4 Injections per 60-week Period, by Skeletal Site and Cohort

| Duration of Treatment | Medroxyprogesterone Acetate Injectable Suspension, USP (150 mg IM) | | Unmatched, Untreated Cohort | |
|-------------------------|--|---------------|-----------------------------|---------------|
| | N | Mean % Change | N | Mean % Change |
| Total Hip BMD | | | | |
| Week 60 (1.2 years) | 113 | -2.75 | 166 | 1.22 |
| Week 120 (2.3 years) | 73 | -5.40 | 109 | 2.19 |
| Week 240 (4.6 years) | 28 | -6.40 | 84 | 1.71 |
| Femoral Neck BMD | | | | |
| Week 60 | 113 | -2.96 | 166 | 1.75 |
| Week 120 | 73 | -5.30 | 108 | 2.83 |
| Week 240 | 28 | -5.40 | 84 | 1.94 |
| Lumbar Spine BMD | | | | |
| Week 60 | 114 | -2.47 | 167 | 3.39 |
| Week 120 | 73 | -2.74 | 109 | 5.28 |
| Week 240 | 27 | -2.11 | 84 | 6.40 |

BMD recovery post-treatment in adolescent women

Longer duration of treatment and smoking were associated with less recovery of BMD following the last injection of Medroxyprogesterone Acetate Injectable Suspension, USP. Table 6 shows the extent of recovery of BMD up to 60 months post-treatment for adolescent women who received Medroxyprogesterone Acetate Injectable Suspension, USP for two years or less compared to more than two years. Post-treatment follow-up showed that, in women treated for more than two years, only lumbar spine BMD recovered to baseline levels after treatment was discontinued. Subjects treated with Medroxyprogesterone Acetate Injectable Suspension, USP for more than two years did not recover to their baseline BMD level at femoral neck and total hip even up to 60 months post-treatment. Adolescent women in the untreated cohort gained BMD throughout the trial period (data not shown).

Table 6: Extent of BMD Recovery (Months Post-Treatment) in Adolescents by Years of Medroxyprogesterone Acetate Injectable Suspension, USP Use (2 Years or Less vs. More than 2 Years)

| Duration of Treatment | 2 years or less | | More than 2 years | |
|-----------------------|-----------------|--------------------|-------------------|--------------------|
| | N | Mean % Change from | N | Mean % Change from |
| | | | | |

| | | baseline | | baseline |
|-------------------------|----|----------|----|----------|
| Total Hip BMD | | | | |
| End of Treatment | 49 | -1.5% | 49 | -6.2% |
| 12 M post-treatment | 33 | -1.4% | 24 | -4.6% |
| 24 M post-treatment | 18 | 0.3% | 17 | -3.6% |
| 36 M post-treatment | 12 | 2.1% | 11 | -4.6% |
| 48 M post-treatment | 10 | 1.3% | 9 | -2.5% |
| 60 M post-treatment | 3 | 0.2% | 2 | -1.0% |
| Femoral Neck BMD | | | | |
| End of Treatment | 49 | -1.6% | 49 | -5.8% |
| 12 M post-treatment | 33 | -1.4% | 24 | -4.3% |
| 24 M post-treatment | 18 | 0.5% | 17 | -3.8% |
| 36 M post-treatment | 12 | 1.2% | 11 | -3.8% |
| 48 M post-treatment | 10 | 2.0% | 9 | -1.7% |
| 60 M post-treatment | 3 | 1.0% | 2 | -1.9% |
| Lumbar Spine BMD | | | | |
| End of Treatment | 49 | -0.9% | 49 | -3.5% |
| 12 M post-treatment | 33 | 0.4% | 23 | -1.1% |
| 24 M post-treatment | 18 | 2.6% | 17 | 1.9% |
| 36 M post-treatment | 12 | 2.4% | 11 | 0.6% |
| 48 M post-treatment | 10 | 6.5% | 9 | 3.5% |
| 60 M post-treatment | 3 | 6.2% | 2 | 5.7% |

14.4 Relationship of fracture incidence to use of Medroxyprogesterone Acetate Injectable Suspension, USP or non-use by women of reproductive age

A retrospective cohort study to assess the association between Medroxyprogesterone Acetate Injectable Suspension, USP and the incidence of bone fractures was conducted in 312,395 female contraceptive users in the UK. The incidence rates of fracture were compared between Medroxyprogesterone Acetate Injectable Suspension, USP users and contraceptive users who had no recorded use of Medroxyprogesterone Acetate Injectable Suspension, USP. The Incident Rate Ratio (IRR) for any fracture during the follow-up period (mean = 5.5 years) was 1.41 (95% CI 1.35, 1.47). It is not known if this is due to Medroxyprogesterone Acetate Injectable Suspension, USP use or to other related lifestyle factors that have a bearing on fracture rate.

In the study, when cumulative exposure to Medroxyprogesterone Acetate Injectable Suspension, USP was calculated, the fracture rate in users who received fewer than 8 injections was higher than that in women who received 8 or more injections. However, it is not clear that cumulative exposure, which may include periods of intermittent use separated by periods of non-use, is a useful measure of risk, as compared to exposure measures based on continuous use.

There were very few osteoporotic fractures (fracture sites known to be related to low BMD) in the study overall, and the incidence of osteoporotic fractures was not found to be higher in Medroxyprogesterone Acetate Injectable Suspension, USP users compared to non-users. Importantly, this study could not determine whether use of Medroxyprogesterone Acetate Injectable Suspension, USP has an effect on fracture rate later in life.

16 HOW SUPPLIED/STORAGE AND HANDLING

Medroxyprogesterone Acetate Injectable Suspension, USP (Medroxyprogesterone Acetate sterile aqueous suspension 150 mg/mL) is supplied in the following strengths and package configurations:

| | | | |
|---------|--|--|--|
| Package | | | |
|---------|--|--|--|

| Package Configuration | Strength | NDC |
|--|-----------|------------------|
| MEDROXYPROGESTERONE ACETATE INJECTABLE SUSPENSION, USP (medroxyprogesterone acetate sterile aqueous suspension 150 mg/mL) | | |
| 1 mL vial | 150 mg/mL | NDC 59762-4537-1 |
| 25 × 1 mL vials | 150 mg/mL | NDC 59762-4537-2 |
| MEDROXYPROGESTERONE ACETATE INJECTABLE SUSPENSION, USP prefilled syringes packaged with 22 gauge × 1 1/2 inch BD SafetyGlide™ Needles | | |
| 1 mL prefilled syringe | 150 mg/mL | NDC 59762-4538-1 |

Vials MUST be stored upright at controlled room temperature 20° to 25°C (68° to 77°F) [see USP].
Store at controlled room temperature 20° to 25°C (68° to 77°F) [see USP].

17 PATIENT COUNSELING INFORMATION

"See FDA-approved patient labeling (Patient Information)."

- Advise patients at the beginning of treatment that their menstrual cycle may be disrupted and that irregular and unpredictable bleeding or spotting results, and that this usually decreases to the point of amenorrhea as treatment with Medroxyprogesterone Acetate Injectable Suspension, USP continues, without other therapy being required.
- Counsel patients that this product does not protect against HIV infection (AIDS) and other sexually transmitted diseases.
- Counsel patients on Warnings and Precautions associated with use of Medroxyprogesterone Acetate Injectable Suspension, USP.
- Counsel patients to use a back-up method or alternative method of contraception when enzyme inducers are used with Medroxyprogesterone Acetate Injectable Suspension, USP.



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Patient Information

(Medroxyprogesterone Acetate Injectable Suspension, USP)

Read this Patient Information carefully before you decide if Medroxyprogesterone Acetate Injectable Suspension, USP is right for you. This information does not take the place of talking with your gynecologist or other healthcare provider who specializes in women's health. If you have any questions about Medroxyprogesterone Acetate Injectable Suspension, USP, ask your healthcare provider. You should also learn about other birth control methods to choose the one that is best for you.

What is the most important information I should know about Medroxyprogesterone Acetate Injectable Suspension, USP?

Medroxyprogesterone Acetate Injectable Suspension, USP can cause serious side effects, including:

- **Use of Medroxyprogesterone Acetate Injectable Suspension, USP may cause you to lose calcium stored in your bone and decrease your bone mass. The longer you use**

Medroxyprogesterone Acetate Injectable Suspension, USP, the greater your loss of calcium from your bones. Your bones may not recover completely when you stop using Medroxyprogesterone Acetate Injectable Suspension, USP.

- **If you use Medroxyprogesterone Acetate Injectable Suspension, USP continuously for a long time (for more than 2 years), it may increase the risk of weak, porous bones (osteoporosis) that could increase the risk of broken bones, especially after menopause.**
- **You should not use Medroxyprogesterone Acetate Injectable Suspension, USP for more than two years unless you cannot use other birth control methods.**
- **It is not known if your risk of developing osteoporosis is greater if you are a teenager or young adult when you start to use Medroxyprogesterone Acetate Injectable Suspension, USP. (See "What are the possible side effects of Medroxyprogesterone Acetate Injectable Suspension, USP?").**

Medroxyprogesterone Acetate Injectable Suspension, USP is intended to prevent pregnancy. Medroxyprogesterone Acetate Injectable Suspension, USP does not protect against HIV infection (AIDS) and other sexually transmitted diseases (STDs).

What is Medroxyprogesterone Acetate Injectable Suspension, USP?

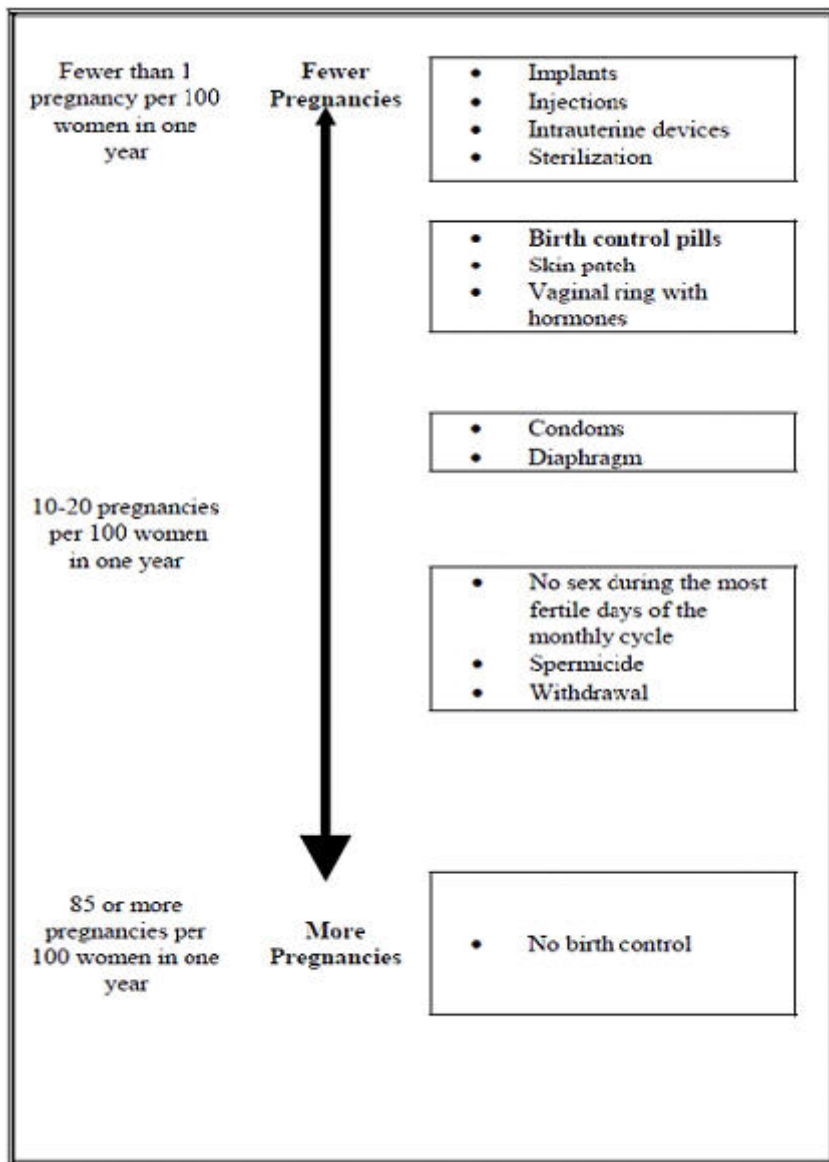
Medroxyprogesterone Acetate Injectable Suspension, USP is a progestin hormone birth control method that is given by injection (a shot) to prevent pregnancy.

How well does Medroxyprogesterone Acetate Injectable Suspension, USP work?

Your chance of getting pregnant depends on how well you follow the directions for taking your Medroxyprogesterone Acetate Injectable Suspension, USP. The more carefully you follow the directions (such as returning every 3 months for your next injection), the less chance you have of getting pregnant.

In clinical studies, about 1 out of 100 women got pregnant during the first year that they used Medroxyprogesterone Acetate Injectable Suspension, USP.

The following chart shows the chance of getting pregnant for women who use different methods of birth control. Each box on the chart contains a list of birth control methods that are similar in effectiveness. The most effective methods are at the top of the chart. The box on the bottom of the chart shows the chance of getting pregnant for women who do not use birth control and are trying to get pregnant.



How should I take Medroxyprogesterone Acetate Injectable Suspension, USP?

- Medroxyprogesterone Acetate Injectable Suspension, USP is given by your healthcare provider as a shot into your muscle (intramuscular injection). The shot is given in your buttock or upper arm 1 time every 3 months. At the end of the 3 months, you will need to return to your healthcare provider for your next injection in order to continue your protection against pregnancy.
- **To make sure that you are not pregnant before you take Medroxyprogesterone Acetate Injectable Suspension, USP, the first injection should be given only:**
 - during the first 5 days of a normal menstrual period, or
 - within the first 5 days after giving birth, **if you are not breastfeeding**, or
 - at the 6th week after giving birth, **if you are feeding your baby only breastmilk.**
- Medroxyprogesterone Acetate Injectable Suspension, USP may be given at other times than those listed above, but you will likely need to have a pregnancy test first to show that you are not pregnant.
- During treatment with Medroxyprogesterone Acetate Injectable Suspension, USP, you should see your healthcare provider every year for a blood pressure check and other healthcare needs.

Who Should Not Use Medroxyprogesterone Acetate Injectable Suspension, USP?

Do not use Medroxyprogesterone Acetate Injectable Suspension, USP if you:

- are pregnant or think you might be pregnant
- have bleeding from your vagina that has not been explained

- have breast cancer now or in the past, or think you have breast cancer
- have had a stroke
- ever had blood clots in your arms, legs or lungs
- have problems with your liver or liver disease
- are allergic to Medroxyprogesterone Acetate Injectable Suspension, USP medroxyprogesterone acetate or any of the other ingredients. See the end of this leaflet for a complete list of ingredients in Medroxyprogesterone Acetate Injectable Suspension, USP.

What should I tell my healthcare provider before taking Medroxyprogesterone Acetate Injectable Suspension, USP?

Before taking Medroxyprogesterone Acetate Injectable Suspension, USP, tell your healthcare provider if you have:

- risk factors for weak bones (osteoporosis) such as bone disease, use alcohol or smoke regularly, anorexia nervosa, or a strong family history of osteoporosis
- irregular or lighter than usual menstrual periods
- breast cancer now or in the past, or think you have breast cancer
- a family history of breast cancer
- an abnormal mammogram (breast X-ray), fibrocystic breast disease, breast nodules or lumps, or bleeding from your nipples
- kidney problems
- high blood pressure
- had a stroke
- had blood clots in your arms, legs or lungs
- migraine headaches
- asthma
- epilepsy (convulsions or seizures)
- diabetes
- depression or a history of depression
- any other medical conditions

If you are breastfeeding or plan to breastfeed, Medroxyprogesterone Acetate Injectable Suspension, USP can pass into your breast milk. Talk to your healthcare provider about the best way to feed your baby if you take Medroxyprogesterone Acetate Injectable Suspension, USP.

Tell your healthcare provider about all of the medicines you take, including prescription and nonprescription medicines, vitamins, and herbal supplements.

Medroxyprogesterone Acetate Injectable Suspension, USP and certain other medicines may affect each other, causing serious side effects.

Sometimes the doses of other medicines may need to be changed while you are taking Medroxyprogesterone Acetate Injectable Suspension, USP.

Some medicines may make Medroxyprogesterone Acetate Injectable Suspension, USP less effective at preventing pregnancy, including those listed below.

Especially tell your healthcare provider if you take:

- medicine to help you sleep
- bosentan
- medicine for seizures
- griseofulvin
- an antibiotic
- medicine for HIV (AIDS)
- St. John's wort

Know the medicines you take. Keep a list of your medicines with you to show your healthcare provider or pharmacist before you first start taking Medroxyprogesterone Acetate Injectable Suspension, USP or when you get a new medicine.

Follow your healthcare provider's instructions about using a back-up method of birth control if you are taking medicines that may make Medroxyprogesterone Acetate Injectable Suspension, USP less effective.

What are the possible side effects of Medroxyprogesterone Acetate Injectable Suspension, USP?

Medroxyprogesterone Acetate Injectable Suspension, USP can cause **serious** side effects, including:

- Effect on the bones: See "What is the most important information I should know about Medroxyprogesterone Acetate Injectable Suspension, USP?". Teenage years are the most important years to gain bone strength. The decrease in calcium in your bones is of most concern if you are a teenager or have the following problems:
 - bone disease
 - an eating disorder (anorexia nervosa)
 - a strong family history of osteoporosis
 - you take a drug that can lower the amount of calcium in your bones (drugs for epilepsy or steroid drugs)
 - you drink a lot of alcohol (more than 2 drinks a day)
 - you smoke

If you need a birth control method for more than 2 years, your healthcare provider may switch you to another birth control method instead of using Medroxyprogesterone Acetate Injectable Suspension, USP. If you continue using Medroxyprogesterone Acetate Injectable Suspension, USP, your healthcare provider may ask you to have a bone test, especially if you have other risks for weak bones.

When Medroxyprogesterone Acetate Injectable Suspension, USP is stopped, your bones may start to regain calcium. However, in a study of teenage girls who used Medroxyprogesterone Acetate Injectable Suspension, USP for more than 2 years, their hip bones did not completely recover by 5 years after they stopped using Medroxyprogesterone Acetate Injectable Suspension, USP. Taking calcium and Vitamin D and exercising daily may lessen the loss of calcium from your bones.

- increased risk of breast cancer. Studies of women who have used different forms of contraception found that women under 35 years of age who first used Medroxyprogesterone Acetate Injectable Suspension, USP within the previous 4 to 5 years may have a slightly increased risk of developing breast cancer.
- blood clots in your arms, legs, lungs, and eyes
- stroke
- a pregnancy outside of your uterus (ectopic pregnancy). Ectopic pregnancy is a medical emergency that often requires surgery. Ectopic pregnancy can cause internal bleeding, infertility, and even death.
- allergic reactions. Severe allergic reactions have been reported in some women using Medroxyprogesterone Acetate Injectable Suspension, USP.
- loss of vision or other eye problems
- migraine headaches
- depression
- convulsions or seizures
- liver problems

Call your healthcare provider right away if you have:

- sharp chest pain, coughing up blood, or sudden shortness of breath (indicating a possible clot in the lung)
- sudden severe headache or vomiting, dizziness or fainting, problems with your eyesight or speech, weakness, or numbness in an arm or leg (indicating a possible stroke)

- severe pain or swelling in the calf (indicating a possible clot in the leg)
- sudden blindness, partial or complete (indicating a possible clot in the blood vessels of the eye)
- unusually heavy vaginal bleeding
- severe pain or tenderness in the lower abdominal area
- persistent pain, pus, or bleeding at the injection site
- yellowing of the eyes or skin
- hives or difficulty breathing

The most common side effects of Medroxyprogesterone Acetate Injectable Suspension, USP include:

- irregular vaginal bleeding, such as lighter or heavier menstrual bleeding, or continued spotting
- weight gain. You may experience weight gain while you are using Medroxyprogesterone Acetate Injectable Suspension, USP. About two-thirds of the women who used Medroxyprogesterone Acetate Injectable Suspension, USP in the clinical trials reported a weight gain of about 5 pounds during the first year of use. You may continue to gain weight after the first year. Women who used Medroxyprogesterone Acetate Injectable Suspension, USP for 2 years gained an average of 8 pounds over those 2 years.
- abdominal pain
- headache
- weakness
- tiredness
- nervousness
- dizziness

Tell your healthcare provider if you have any side effect that bothers you or does not go away.

These are not all the possible side effects of Medroxyprogesterone Acetate Injectable Suspension, USP. 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 at 1- 800-FDA-1088.

What other information should I know before choosing Medroxyprogesterone Acetate Injectable Suspension, USP?

- **Pregnancy.** When you take Medroxyprogesterone Acetate Injectable Suspension, USP every 3 months, your chance of getting pregnant is very low. You could miss a period or have a light period and not be pregnant. If you miss 1 or 2 periods and think you might be pregnant, see your healthcare provider as soon as possible. You should not use Medroxyprogesterone Acetate Injectable Suspension, USP if you are pregnant. However, Medroxyprogesterone Acetate Injectable Suspension, USP taken by accident during pregnancy does not seem to cause birth defects.
- **Nursing Mothers.** Although Medroxyprogesterone Acetate Injectable Suspension, USP can be passed to the nursing baby in the breast milk, no harmful effects on babies have been found. Medroxyprogesterone Acetate Injectable Suspension, USP does not stop the breasts from producing milk, so it can be used by nursing mothers. However, to minimize the amount of Medroxyprogesterone Acetate Injectable Suspension, USP that is passed to the baby in the first weeks after birth, you should wait until your baby is 6 weeks old before you start using Medroxyprogesterone Acetate Injectable Suspension, USP for birth control.

How will Medroxyprogesterone Acetate Injectable Suspension, USP change my periods?

- **Change in normal menstrual cycle.** The side effect reported most frequently by women who use Medroxyprogesterone Acetate Injectable Suspension, USP for birth controls is a change in their normal menstrual cycle. During the first year of using Medroxyprogesterone Acetate Injectable Suspension, USP, you might have one or more of the following changes:
 - irregular or unpredictable bleeding or spotting
 - an increase or decrease in menstrual bleeding

- **no bleeding at all.** In clinical studies of Medroxyprogesterone Acetate Injectable Suspension, USP, 55% of women reported no menstrual bleeding (amenorrhea) after one year of use and 68% of women reported no menstrual bleeding after two years of use.
- **Missed period.** During the time you are using Medroxyprogesterone Acetate Injectable Suspension, USP for birth controls, you may skip a period, or your periods may stop completely. If you have been receiving your shot of Medroxyprogesterone Acetate Injectable Suspension, USP regularly every 3 months, then you are probably not pregnant. However, if you think that you may be pregnant, see your healthcare provider.

Unusually heavy or continuous bleeding is not a usual effect of Medroxyprogesterone Acetate Injectable Suspension, USP and if this happens you should see your healthcare provider right away.

With continued use of Medroxyprogesterone Acetate Injectable Suspension, USP, bleeding usually decreases and many women stop having periods completely. When you stop using Medroxyprogesterone Acetate Injectable Suspension, USP your menstrual period will usually, in time, return to its normal cycle.

What if I want to become pregnant?

Because Medroxyprogesterone Acetate Injectable Suspension, USP is a long-acting birth control method, it takes some time after your last shot for its effect to wear off. Most women who try to get pregnant after using Medroxyprogesterone Acetate Injectable Suspension, USP get pregnant within 18 months after their last shot. The length of time you use Medroxyprogesterone Acetate Injectable Suspension, USP has no effect on how long it takes you to become pregnant after you stop using it.

General Information about Medroxyprogesterone Acetate Injectable Suspension, USP

Medicines are sometimes prescribed for conditions that are not mentioned in patient information leaflets. This leaflet summarizes the most important information about Medroxyprogesterone Acetate Injectable Suspension, USP. If you would like more information, talk with your healthcare provider. You can ask your healthcare provider for information about Medroxyprogesterone Acetate Injectable Suspension, USP that is written for healthcare providers.

What are the ingredients in Medroxyprogesterone Acetate Injectable Suspension, USP?

Active ingredient: medroxyprogesterone acetate

Inactive ingredients: polyethylene glycol 3350, polysorbate 80, sodium chloride, methylparaben, propylparaben, and water for injection. When necessary, pH is adjusted with sodium hydroxide or hydrochloric acid, or both.



GREENSTONE® BRAND

Distributed by

Greenstone LLC

Peapack, NJ 07977

LAB-0152-7.0

September 2011

PRINCIPAL DISPLAY PANEL - 1 mL Vial

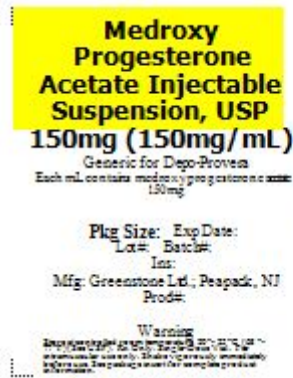
NDC 68788-9233-1

GREENSTONE® BRAND

**medroxyprogesterone
acetate injectable
suspension, USP**

150 mg in 1 mL

Intramuscular Use Only



Medroxy Progesterone Acetate
Injectable Suspension, USP 150 mg
(150 mg/mL)
Qty: 10
Lot: B04

Medroxy Progesterone Acetate
Injectable Suspension, USP 150 mg
(150 mg/mL)
Qty: 10
Lot: B04

Medroxy Progesterone Acetate
Injectable Suspension, USP 150 mg
(150 mg/mL)
Qty: 10
Lot: B04

Medroxy Progesterone Acetate
Injectable Suspension, USP 150 mg
(150 mg/mL)
Qty: 10
Lot: B04

Log

Client

Billing

Patient

MEDROXYPROGESTERONE ACETATE

medroxyprogesterone acetate injection, suspension

Product Information

| | | | |
|-------------------------|-------------------------|--------------------|--------------------------------|
| Product Type | HUMAN PRESCRIPTION DRUG | Item Code (Source) | NDC:68788-9233(NDC:59762-4537) |
| Route of Administration | INTRAMUSCULAR | | |

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|--|---------------------|----------------|
| medroxyprogesterone acetate (UNII: C2QI4IOI2G) (medroxyprogesterone - UNII:HSU1C9YRES) | medroxyprogesterone | 150 mg in 1 mL |

Inactive Ingredients

| Ingredient Name | Strength |
|---|-----------------|
| Polyethylene glycol 3350 (UNII: G2M7P15E5P) | 28.9 mg in 1 mL |
| Polysorbate 80 (UNII: 6OZP39ZG8H) | 2.41 mg in 1 mL |
| Sodium chloride (UNII: 451W47IQ8X) | 8.68 mg in 1 mL |
| Methylparaben (UNII: A2I8C7HI9T) | 1.37 mg in 1 mL |
| Propylparaben (UNII: Z8IX2SC10H) | 0.15 mg in 1 mL |
| Water (UNII: 059QF0KO0R) | |
| sodium hydroxide (UNII: 55X04QC32I) | |
| hydrochloric acid (UNII: QTT17582CB) | |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|---------------------|----------------------|--------------------|
| 1 | NDC:68788-9233-1 | 1 mL in 1 VIAL | | |

Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|--------------------|--|----------------------|--------------------|
|--------------------|--|----------------------|--------------------|

| | | | |
|------------------------|-----------|------------|--|
| NDA authorized generic | NDA020246 | 10/29/1992 | |
|------------------------|-----------|------------|--|

Labeler - Preferred Pharmaceuticals, Inc (791119022)

Establishment

| Name | Address | ID/FEI | Business Operations |
|--------------------------------|---------|-------------|---------------------|
| Preferred Pharmaceuticals, Inc | | 79 1119 022 | relabel |

Revised: 9/2011

Preferred Pharmaceuticals, Inc