

FINASTERIDE- finasteride tablet, film coated **Direct_Rx**

Finasteride

1.1 Monotherapy

Finasteride tablets are indicated for the treatment of symptomatic benign prostatic hyperplasia (BPH) in men with an enlarged prostate to:

- Improve symptoms
- Reduce the risk of acute urinary retention
- Reduce the risk of the need for surgery including transurethral resection of the prostate (TURP) and prostatectomy.

1.2 Combination with Alpha-Blocker

Finasteride tablets administered in combination with the alpha-blocker doxazosin is indicated to reduce the risk of symptomatic progression of BPH (a confirmed ≥ 4 point increase in American Urological Association (AUA) symptom score).

1.3 Limitations of Use

Finasteride tablets are not approved for the prevention of prostate cancer.

Finasteride tablets may be administered with or without meals.

2.1 Monotherapy

The recommended dose of finasteride tablet is one tablet (5 mg) taken once a day [see CLINICAL STUDIES (14.1)].

2.2 Combination with Alpha-Blocker

The recommended dose of finasteride tablet is one tablet (5 mg) taken once a day in combination with the alpha-blocker doxazosin [see CLINICAL STUDIES (14.2)].

5 mg blue colored, round, biconvex, film-coated tablets, marked "F5" on one side and plain on other side.

Finasteride tablets are contraindicated in the following:

Hypersensitivity to any component of this medication.

Pregnancy. Finasteride use is contraindicated in females when they are or may potentially be pregnant. Because of the ability of Type II 5 α -reductase inhibitors to inhibit the conversion of testosterone to 5 α -dihydrotestosterone (DHT), finasteride may cause abnormalities of the external genitalia of a male fetus of a pregnant female who receives finasteride. If this drug is used during pregnancy, or if pregnancy occurs while taking this drug, the pregnant female should be apprised of the potential hazard to the male fetus. [See also WARNINGS AND PRECAUTIONS (5.3), USE IN SPECIFIC POPULATIONS (8.1), and HOW SUPPLIED/STORAGE AND HANDLING (16).] In female rats, low doses of finasteride administered during pregnancy have produced abnormalities of the external genitalia in male offspring.

5.1 Effects on Prostate Specific Antigen (PSA) and the Use of PSA in Prostate Cancer

Detection

In clinical studies, finasteride tablets reduced serum PSA concentration by approximately 50% within six months of treatment. This decrease is predictable over the entire range of PSA values in patients with symptomatic BPH, although it may vary in individuals.

For interpretation of serial PSAs in men taking finasteride tablets, a new PSA baseline should be established at least six months after starting treatment and PSA monitored periodically thereafter. Any confirmed increase from the lowest PSA value while on finasteride tablets may signal the presence of prostate cancer and should be evaluated, even if PSA levels are still within the normal range for men not taking a 5 α -reductase inhibitor. Non-compliance with finasteride tablets therapy may also affect PSA test results. To interpret an isolated PSA value in patients treated with finasteride tablets for six months or more, PSA values should be doubled for comparison with normal ranges in untreated men. These adjustments preserve the utility of PSA to detect prostate cancer in men treated with finasteride tablets.

Finasteride tablets may also cause decreases in serum PSA in the presence of prostate cancer.

The ratio of free to total PSA (percent free PSA) remains constant even under the influence of finasteride tablets. If clinicians elect to use percent free PSA as an aid in the detection of prostate cancer in men undergoing finasteride therapy, no adjustment to its value appears necessary.

5.2 Increased Risk of High-Grade Prostate Cancer

Men aged 55 and over with a normal digital rectal examination and PSA \leq 3.0 ng/mL at baseline taking finasteride 5 mg/day in the 7-year Prostate Cancer Prevention Trial (PCPT) had an increased risk of Gleason score 8 to 10 prostate cancer (finasteride 1.8% vs placebo 1.1%). [See INDICATIONS AND USAGE (1.3) and ADVERSE REACTIONS (6.1).] Similar results were observed in a 4-year placebo-controlled clinical trial with another 5 α -reductase inhibitor (dutasteride, AVODART) (1% dutasteride vs 0.5% placebo). 5 α -reductase inhibitors may increase the risk of development of high-grade prostate cancer. Whether the effect of 5 α -reductase inhibitors to reduce prostate volume, or study-related factors, impacted the results of these studies has not been established.

5.3 Exposure of Females- Risk to Male Fetus

Finasteride tablets is contraindicated in pregnant females and in females who may potentially be pregnant and not indicated for use in females. Based on animal studies and the mechanism of action, finasteride tablets may cause abnormal development of external genitalia in a male fetus if administered to a pregnant female. Females who are pregnant or may potentially be pregnant should not handle crushed or broken finasteride tablets. Finasteride tablets are coated and will prevent contact with the active ingredient during normal handling, provided that the tablets have not been broken or crushed. If a pregnant female comes in contact with crushed or broken finasteride tablets, the contact area should be washed immediately with soap and water. [See Contraindications (4), Use in Specific Populations (8.1), Clinical Pharmacology (12.1 and 12.3), and How Supplied/Storage and Handling (16).]

5.4 Pediatric Patients and Females

Finasteride tablets is not indicated for use in pediatric patients [see USE IN SPECIFIC

POPULATIONS (8.4) and CLINICAL PHARMACOLOGY (12.3)] or females [see also WARNINGS AND PRECAUTIONS (5.3), USE IN SPECIFIC POPULATIONS (8.1), CLINICAL PHARMACOLOGY (12.3), and HOW SUPPLIED/ STORAGE AND HANDLING (16)].

5.5 Effect on Semen Characteristics

Treatment with finasteride tablets for 24 weeks to evaluate semen parameters in healthy male volunteers revealed no clinically meaningful effects on sperm concentration, mobility, morphology, or pH. A 0.6 mL (22.1%) median decrease in ejaculate volume with a concomitant reduction in total sperm per ejaculate was observed. These parameters remained within the normal range and were reversible upon discontinuation of therapy with an average time to return to baseline of 84 weeks.

5.6 Consideration of Other Urological Conditions

Prior to initiating treatment with finasteride tablets, consideration should be given to other urological conditions that may cause similar symptoms. In addition, prostate cancer and BPH may coexist.

Patients with large residual urinary volume and/or severely diminished urinary flow should be carefully monitored for obstructive uropathy. These patients may not be candidates for finasteride therapy.

6.1 Clinical Trials Experience

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

4-Year Placebo-Controlled Study (A Long-Term Efficacy and Safety Study)

In A Long-Term Efficacy and Safety Study, 1,524 patients treated with finasteride tablets and 1,516 patients treated with placebo were evaluated for safety over a period of 4 years. The most frequently reported adverse reactions were related to sexual function. 3.7% (57 patients) treated with finasteride tablets and 2.1% (32 patients) treated with placebo discontinued therapy as a result of adverse reactions related to sexual function, which are the most frequently reported adverse reactions.

Table 1 presents the only clinical adverse reactions considered possibly, probably or definitely drug related by the investigator, for which the incidence on finasteride was $\geq 1\%$ and greater than placebo over the 4 years of the study. In years 2 to 4 of the study, there was no significant difference between treatment groups in the incidences of impotence, decreased libido and ejaculation disorder.

Table 1: Drug-Related Adverse Experiences

Year 1

(%)

Years 2, 3 and 4*(%)

Finasteride

Placebo

Finasteride

Placebo

Impotence

8.1
3.7
5.1
5.1

Decreased Libido

6.4
3.4
2.6
2.6

Decreased Volume of Ejaculate

3.7
0.8
1.5
0.5

Ejaculation Disorder

0.8
0.1
0.2
0.1

Breast Enlargement

0.5
0.1
1.8
1.1

Breast Tenderness

0.4
0.1
0.7
0.3

Rash

0.5
0.2
0.5
0.1

*Combined Years 2 to 4

N = 1,524 and 1,516, finasteride vs placebo, respectively

Phase III Studies and 5-Year Open Extensions

The adverse experience profile in the 1-year, placebo-controlled, Phase III studies, the 5-

year open extensions, and A Long-Term Efficacy and Safety Study were similar.

Medical Therapy of Prostatic Symptoms (MTOPS) Study

In the MTOPS study, 3,047 men with symptomatic BPH were randomized to receive finasteride tablets 5 mg/day (n=768), doxazosin 4 or 8 mg/day (n=756), the combination of finasteride tablets 5 mg/day and doxazosin 4 or 8 mg/day (n=786), or placebo (n=737) for 4 to 6 years. [See CLINICAL STUDIES (14.2).]

The incidence rates of drug-related adverse experiences reported by $\geq 2\%$ of patients in any treatment group in the MTOPS Study are listed in Table 2.

The individual adverse effects which occurred more frequently in the combination group compared to either drug alone were:

asthenia, postural hypotension, peripheral edema, dizziness, decreased libido, rhinitis, abnormal ejaculation, impotence and abnormal sexual function (see Table 2). Of these, the incidence of abnormal ejaculation in patients receiving combination therapy was comparable to the sum of the incidences of this adverse experience reported for the two monotherapies.

Combination therapy with finasteride and doxazosin was associated with no new clinical adverse experience.

Four patients in MTOPS reported the adverse experience breast cancer. Three of these patients were on finasteride only and one was on combination therapy. [See Long-Term Data.]

The MTOPS Study was not specifically designed to make statistical comparisons between groups for reported adverse experiences. In addition, direct comparisons of safety data between the MTOPS study and previous studies of the single agents may not be appropriate based upon differences in patient population, dosage or dose regimen, and other procedural and study design elements.

Adverse Experience Placebo

(N=737)

(%) Doxazosin

4 mg or

8 mg*

(N=756)

(%) Finasteride

(N=768)

(%) Combination

(N=786)

(%)

Table 2:

Incidence $\geq 2\%$ in One or More Treatment Groups

Drug-Related Clinical Adverse Experiences in MTOPS

Body as a whole

Asthenia

7.1

15.7

5.3

16.8
Headache
2.3
4.1
2.0
2.3
Cardiovascular
Hypotension
0.7
3.4
1.2
1.5
Postural Hypotension
8.0
16.7
9.1
17.8
Metabolic and Nutritional
Peripheral Edema
0.9
2.6
1.3
3.3
Nervous
Dizziness
8.1
17.7
7.4
23.2
Libido Decreased
5.7
7.0
10.0
11.6
Somnolence
1.5
3.7
1.7
3.1
Respiratory
Dyspnea
0.7
2.1
0.7
1.9
Rhinitis
0.5
1.3
1.0
2.4

Urogenital

Abnormal Ejaculation

2.3

4.5

7.2

14.1

Gynecomastia

0.7

1.1

2.2

1.5

Impotence

12.2

14.4

18.5

22.6

Sexual Function Abnormal

0.9

2.0

2.5

3.1

* Doxazosin dose was achieved by weekly titration (1 to 2 to 4 to 8 mg). The final tolerated dose (4 mg or 8 mg) was administered at end-Week 4. Only those patients tolerating at least 4 mg were kept on doxazosin. The majority of patients received the 8-mg dose over the duration of the study.

Long-Term Data

High-Grade Prostate Cancer

The PCPT trial was a 7-year randomized, double-blind, placebo-controlled trial that enrolled 18,882 men ≥ 55 years of age with a normal digital rectal examination and a PSA ≤ 3.0 ng/mL. Men received either finasteride 5 mg or placebo daily. Patients were evaluated annually with PSA and digital rectal exams. Biopsies were performed for elevated PSA, an abnormal digital rectal exam, or the end of study. The incidence of Gleason score 8 to 10 prostate cancer was higher in men treated with finasteride (1.8%) than in those treated with placebo (1.1%) [see INDICATIONS AND USAGE (1.3) and WARNINGS AND PRECAUTIONS (5.2)]. In a 4-year placebo-controlled clinical trial with another 5 α -reductase inhibitor (dutasteride, AVODART), similar results for Gleason score 8 to 10 prostate cancer were observed (1% dutasteride vs 0.5% placebo).

No clinical benefit has been demonstrated in patients with prostate cancer treated with finasteride tablets.

Breast Cancer

During the 4- to 6-year placebo- and comparator-controlled MTOPS study that enrolled 3,047 men, there were 4 cases of breast cancer in men treated with finasteride but no cases in men not treated with finasteride. During the 4-year, placebo-controlled A Long-Term Efficacy and Safety Study that enrolled 3,040 men, there were 2 cases of breast cancer in placebo-treated men but no cases in men treated with finasteride. During the 7-year placebo-controlled Prostate Cancer Prevention Trial (PCPT) that enrolled 18,882

men, there was 1 case of breast cancer in men treated with finasteride, and 1 case of breast cancer in men treated with placebo. The relationship between long-term use of finasteride and male breast neoplasia is currently unknown.

Sexual Function

There is no evidence of increased sexual adverse experiences with increased duration of treatment with finasteride tablets. New reports of drug-related sexual adverse experiences decreased with duration of therapy.

6.2 Postmarketing Experience

The following additional adverse events have been reported in postmarketing experience with finasteride tablets. Because these events 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:

- hypersensitivity reactions, such as pruritus, urticaria, and angioedema (including swelling of the lips, tongue, throat, and face)
- testicular pain
- hematospermia
- sexual dysfunction that continued after discontinuation of treatment, including erectile dysfunction, decreased libido and ejaculation disorders (e.g. reduced ejaculate volume). These events were reported rarely in men taking finasteride tablets for the treatment of BPH. Most men were older and were taking concomitant medications and/or had co-morbid conditions. The independent role of finasteride tablets in these events is unknown.
- male infertility and/or poor seminal quality were reported rarely in men taking finasteride tablets for the treatment of BPH. Normalization or improvement of poor seminal quality has been reported after discontinuation of finasteride. The independent role of finasteride tablets in these events is unknown.
- depression
- male breast cancer.

The following additional adverse event related to sexual dysfunction that continued after discontinuation of treatment has been reported in postmarketing experience with finasteride at lower doses used to treat male pattern baldness. Because the event is reported voluntarily from a population of uncertain size, it is not always possible to reliably estimate its frequency or establish a causal relationship to drug exposure:

- orgasm disorders

7.1 Cytochrome P450-Linked Drug Metabolizing Enzyme System

No drug interactions of clinical importance have been identified. Finasteride does not appear to affect the cytochrome P450-linked drug metabolizing enzyme system. Compounds that have been tested in man have included antipyrine, digoxin, propranolol, theophylline, and warfarin and no clinically meaningful interactions were found.

7.2 Other Concomitant Therapy

Although specific interaction studies were not performed, finasteride tablets was

concomitantly used in clinical studies with acetaminophen, acetylsalicylic acid, α -blockers, angiotensin-converting enzyme (ACE) inhibitors, analgesics, anti-convulsants, beta-adrenergic blocking agents, diuretics, calcium channel blockers, cardiac nitrates, HMG-CoA reductase inhibitors, nonsteroidal anti-inflammatory drugs (NSAIDs), benzodiazepines, H₂ antagonists and quinolone anti-infectives without evidence of clinically significant adverse interactions.

8.1 Pregnancy

Risk Summary

Finasteride tablets are contraindicated in pregnant females and not indicated for use in females. Based on animal studies and the mechanism of action, finasteride tablets may cause abnormal development of external genitalia in a male fetus if administered to a pregnant female [see Warnings and Precautions (5.3) and Clinical Pharmacology (12.1)].

In an embryo-fetal development study in rats, there was a dose-dependent increase in hypospadias that occurred in 3.6 to 100% of male offspring of pregnant rats administered oral finasteride during the period of major organogenesis at doses approximately 0.1 to 86 times the maximum recommended human dose (MRHD) of 5 mg/day (based on AUC at animal doses of 0.1 to 100 mg/kg/day). Decreased prostatic and seminal vesicular weights, delayed preputial separation and transient nipple development were also observed in male offspring at oral maternal doses approximately 0.03 times the MRHD (based on AUC at animal dose of 0.03 mg/kg/day), along with decreased anogenital distance in male offspring at oral maternal doses approximately 0.003 times the MRHD (based on AUC at animal dose of 0.003 mg/kg/day).

Finasteride tablet is a Type II 5 α -reductase inhibitor that prevents conversion of testosterone to 5 α -dihydrotestosterone (DHT), a hormone necessary for normal development of male genitalia. If this drug is used during pregnancy, or if the patient becomes pregnant while taking this drug, the patient should be apprised of the potential hazard to the male fetus.

Abnormal male genital development is an expected consequence when conversion of testosterone to 5 α -dihydrotestosterone (DHT) is inhibited by 5 α -reductase inhibitors. These outcomes are similar to those reported in male infants with genetic 5 α -reductase deficiency. Females could be exposed to finasteride through contact with crushed or broken finasteride tablets or semen from a male partner taking finasteride tablets. With regard to finasteride exposure through the skin, finasteride tablets are coated and will prevent skin contact with finasteride during normal handling if the tablets have not been crushed or broken. Females who are pregnant or may potentially be pregnant should not handle crushed or broken finasteride tablets because of possible exposure of a male fetus. With regard to potential finasteride exposure through semen, three studies have been conducted that measured finasteride concentrations in semen in men receiving finasteride tablets 5 mg/day. In these studies the highest amount of finasteride in semen was estimated to be 50-to 100-fold less than the dose of finasteride (5 μ g) that had no effect on circulating DHT levels in men [see Data and Clinical Pharmacology (12.3)].

Data

Human Data

In 2 studies of healthy subjects (n=69) receiving finasteride tablets 5 mg/day for 6 to 24 weeks, finasteride concentrations in semen ranged from undetectable (<0.1 ng/mL) to

10.54 ng/mL. In an earlier study using a less sensitive assay, finasteride concentrations in semen of 16 subjects receiving finasteride tablets 5 mg/day ranged from undetectable (<1.0 ng/mL) to 21 ng/mL. Using the highest semen level measured and assuming 100% absorption would be up to 105 ng per day, which is 50-to 100-fold less than the dose of finasteride (5 µg) that had no effect on circulating DHT levels in men [see Clinical Pharmacology (12.3)].

Animal Data

In an embryo-fetal development study, pregnant rats received finasteride during the period of major organogenesis (gestation days 6 to 17). At maternal doses of oral finasteride approximately 0.1 to 86 times the maximum recommended human dose (MRHD) of 5 mg/day (based on AUC at animal doses of 0.1 to 100 mg/kg/day) there was a dose-dependent increase in hypospadias that occurred in 3.6 to 100% of male offspring. Exposure multiples were estimated using data from nonpregnant rats. Days 16 to 17 of gestation is a critical period in male fetal rats for differentiation of the external genitalia. At oral maternal doses approximately 0.03 times the MRHD (based on AUC at animal dose of 0.03 mg/kg/day), male offspring had decreased prostatic and seminal vesicular weights, delayed preputial separation and transient nipple development. Decreased anogenital distance occurred in male offspring of pregnant rats that received approximately 0.003 times the MRHD (based on AUC at animal dose of 0.003 mg/kg/day). No abnormalities were observed in female offspring at any maternal dose of finasteride.

No developmental abnormalities were observed in the offspring of untreated females mated with finasteride treated male rats that received approximately 61 times the MRHD (based on AUC at animal dose of 80 mg/kg/day). Slightly decreased fertility was observed in male offspring after administration of about 3 times the MRHD (based on AUC at animal dose of 3 mg/kg/day) to female rats during late gestation and lactation. No effects on fertility were seen in female offspring under these conditions.

No evidence of male external genital malformations or other abnormalities were observed in rabbit fetuses exposed to finasteride during the period of major organogenesis (gestation days 6 to 18) at maternal oral doses up to 100 mg/kg /day, (finasteride exposure levels were not measured in rabbits). However, this study may not have included the critical period for finasteride effects on development of male external genitalia in the rabbit.

The fetal effects of maternal finasteride exposure during the period of embryonic and fetal development were evaluated in the rhesus monkey (gestation days 20 to 100), in a species and development period more predictive of specific effects in humans than the studies in rats and rabbits. Intravenous administration of finasteride to pregnant monkeys at doses as high as 800 ng/day (estimated maximal blood concentration of 1.86 ng/mL or about 143 times the highest estimated exposure of pregnant females to finasteride from semen of men taking 5 mg/day) resulted in no abnormalities in male fetuses. In confirmation of the relevance of the rhesus model for human fetal development, oral administration of a dose of finasteride (2 mg/kg/day or approximately 18,000 times the highest estimated blood levels of finasteride from semen of men taking 5 mg/day) to pregnant monkeys resulted in external genital abnormalities in male fetuses. No other abnormalities were observed in male fetuses and no finasteride-related abnormalities were observed in female fetuses at any dose.

8.2 Lactation

Risk Summary

Finasteride tablets are not indicated for use in females.

8.3 Females and Males of Reproductive Potential

Infertility

Females

Finasteride tablets are not indicated for use in females.

Males

Treatment with finasteride tablets for 24 weeks to evaluate semen parameters in healthy male volunteers revealed no clinically meaningful effects on sperm concentration, mobility, morphology, or pH. A 0.6 mL (22.1%) median decrease in ejaculate volume with a concomitant reduction in total sperm per ejaculate was observed. These parameters remained within the normal range and were reversible upon discontinuation of therapy with an average time to return to baseline of 84 weeks [see Warnings and Precautions (5.5)].

There have been postmarketing reports of male infertility and/or poor seminal quality; normalization or improvement of seminal quality has been reported after discontinuation of finasteride [see Adverse Reactions (6.2)].

8.4 Pediatric Use

Finasteride tablets are not indicated for use in pediatric patients.

Safety and effectiveness in pediatric patients have not been established.

8.5 Geriatric Use

Of the total number of subjects included in A Long-Term Efficacy and Safety Study, 1,480 and 105 subjects were 65 and over and 75 and over, respectively. No overall differences in safety or effectiveness were observed between these subjects and younger subjects, and other reported clinical experience has not identified differences in responses between the elderly and younger patients. No dosage adjustment is necessary in the elderly [see CLINICAL PHARMACOLOGY (12.3) and CLINICAL STUDIES (14)] .

8.6 Hepatic Impairment

Caution should be exercised in the administration of finasteride tablets in those patients with liver function abnormalities, as finasteride is metabolized extensively in the liver [see CLINICAL PHARMACOLOGY (12.3)].

8.7 Renal Impairment

No dosage adjustment is necessary in patients with renal impairment [see CLINICAL PHARMACOLOGY (12.3)] .

Patients have received single doses of finasteride tablets up to 400 mg and multiple doses of finasteride tablets up to 80 mg/day for three months without adverse effects. Until further experience is obtained, no specific treatment for an overdose with finasteride tablets can be recommended. Significant lethality was observed in male and female mice at single oral doses of 1,500 mg/m² (500 mg/kg) and in female and male

rats at single oral doses of 2,360 mg/m² (400 mg/kg) and 5,900 mg/m² (1,000 mg/kg), respectively.

Finasteride, a synthetic 4-azasteroid compound, is a specific inhibitor of steroid Type II 5 α -reductase, an intracellular enzyme that converts the androgen testosterone into 5 α -dihydrotestosterone (DHT).

Finasteride is 4-azaandrost-1-ene-17-carboxamide, N-(1, 1-dimethylethyl)-3-oxo-, (5 α , 17 β)-. The empirical formula of finasteride is C₂₃H₃₆N₂O₂ and its molecular weight is 372.55. Its structural formula is:

[Structure]

Finasteride USP is a white crystalline powder with a melting point near 250°C. It is freely soluble in chloroform and in lower alcohol solvents, but is practically insoluble in water.

Finasteride tablets, USP for oral administration are film-coated tablets that contain 5 mg of finasteride and the following inactive ingredients: lactose monohydrate, microcrystalline cellulose, pregelatinized starch (maize), sodium starch glycolate, lauroylmacrogol 32 Glycer, magnesium stearate, hypromellose, titanium dioxide, polyethylene glycol, and FD & C blue #2/indigo carmine aluminium lake.

12.1 Mechanism of Action

The development and enlargement of the prostate gland is dependent on the potent androgen, 5 α -dihydrotestosterone (DHT). Type II 5 α -reductase metabolizes testosterone to DHT in the prostate gland, liver and skin. DHT induces androgenic effects by binding to androgen receptors in the cell nuclei of these organs.

Finasteride is a competitive and specific inhibitor of Type II 5 α -reductase with which it slowly forms a stable enzyme complex.

Turnover from this complex is extremely slow ($t_{1/2}$ ~ 30 days). This has been demonstrated both in vivo and in vitro. Finasteride has no affinity for the androgen receptor. In man, the 5 α -reduced steroid metabolites in blood and urine are decreased after administration of finasteride.

12.2 Pharmacodynamics

In man, a single 5-mg oral dose of finasteride tablets produces a rapid reduction in serum DHT concentration, with the maximum effect observed 8 hours after the first dose. The suppression of DHT is maintained throughout the 24-hour dosing interval and with continued treatment. Daily dosing of finasteride tablets at 5 mg/day for up to 4 years has been shown to reduce the serum DHT concentration by approximately 70%. The median circulating level of testosterone increased by approximately 10 to 20% but remained within the physiologic range. In a separate study in healthy men treated with finasteride 1 mg per day (n=82) or placebo (n=69), mean circulating levels of testosterone and estradiol were increased by approximately 15% as compared to baseline, but these remained within the physiologic range.

In patients receiving finasteride tablets 5 mg/day, increases of about 10% were observed in luteinizing hormone (LH) and follicle-stimulating hormone (FSH), but levels remained within the normal range. In healthy volunteers, treatment with finasteride

tablets did not alter the response of LH and FSH to gonadotropin-releasing hormone indicating that the hypothalamic-pituitary-testicular axis was not affected.

In patients with BPH, finasteride tablets have no effect on circulating levels of cortisol, prolactin, thyroid-stimulating hormone, or thyroxine.

No clinically meaningful effect was observed on the plasma lipid profile (i.e., total cholesterol, low density lipoproteins, high density lipoproteins and triglycerides) or bone mineral density.

Adult males with genetically inherited Type II 5 α -reductase deficiency also have decreased levels of DHT. Except for the associated urogenital defects present at birth, no other clinical abnormalities related to Type II 5 α -reductase deficiency have been observed in these individuals. These individuals have a small prostate gland throughout life and do not develop BPH.

In patients with BPH treated with finasteride (1 to 100 mg/day) for 7 to 10 days prior to prostatectomy, an approximate 80% lower DHT content was measured in prostatic tissue removed at surgery, compared to placebo; testosterone tissue concentration was increased up to 10 times over pretreatment levels, relative to placebo. Intraprostatic content of PSA was also decreased.

In healthy male volunteers treated with finasteride tablets for 14 days, discontinuation of therapy resulted in a return of DHT levels to pretreatment levels in approximately 2 weeks. In patients treated for three months, prostate volume, which declined by approximately 20%, returned to close to baseline value after approximately three months of discontinuation of therapy.

12.3 Pharmacokinetics

Absorption

In a study of 15 healthy young subjects, the mean bioavailability of finasteride 5-mg tablets was 63% (range 34 to 108%), based on the ratio of area under the curve (AUC) relative to an intravenous (IV) reference dose. Maximum finasteride plasma concentration averaged 37 ng/mL (range, 27 to 49 ng/mL) and was reached 1 to 2 hours postdose. Bioavailability of finasteride was not affected by food.

Distribution

Mean steady-state volume of distribution was 76 liters (range, 44 to 96 liters). Approximately 90% of circulating finasteride is bound to plasma proteins. There is a slow accumulation phase for finasteride after multiple dosing. After dosing with 5 mg/day of finasteride for 17 days, plasma concentrations of finasteride were 47 and 54% higher than after the first dose in men 45 to 60 years old (n=12) and \geq 70 years old (n=12), respectively. Mean trough concentrations after 17 days of dosing were 6.2 ng/mL (range, 2.4 to 9.8 ng/mL) and 8.1 ng/mL (range, 1.8 to 19.7 ng/mL), respectively, in the two age groups. Although steady state was not reached in this study, mean trough plasma concentration in another study in patients with BPH (mean age, 65 years) receiving 5 mg/day was 9.4 ng/mL (range, 7.1 to 13.3 ng/mL; n=22) after over a year of dosing.

Finasteride has been shown to cross the blood brain barrier but does not appear to distribute preferentially to the CSF.

In 2 studies of healthy subjects (n=69) receiving finasteride tablets 5 mg/day for 6 to 24

weeks, finasteride concentrations in semen ranged from undetectable (<0.1 ng/mL) to 10.54 ng/mL. In an earlier study using a less sensitive assay, finasteride concentrations in the semen of 16 subjects receiving finasteride tablets 5 mg/day ranged from undetectable (<1.0 ng/mL) to 21 ng/mL. Thus, based on a 5-mL ejaculate volume, the amount of finasteride in semen was estimated to be 50- to 100-fold less than the dose of finasteride (5 mcg) that had no effect on circulating DHT levels in men [see also USE IN SPECIFIC POPULATIONS (8.1)].

Metabolism

Finasteride is extensively metabolized in the liver, primarily via the cytochrome P450 3A4 enzyme subfamily. Two metabolites, the t-butyl side chain monohydroxylated and monocarboxylic acid metabolites, have been identified that possess no more than 20% of the 5 α -reductase inhibitory activity of finasteride.

Excretion

In healthy young subjects (n=15), mean plasma clearance of finasteride was 165 mL/min (range, 70 to 279 mL/min) and mean elimination half-life in plasma was 6 hours (range, 3 to 16 hours). Following an oral dose of ¹⁴C-finasteride in man (n=6), a mean of 39% (range, 32 to 46%) of the dose was excreted in the urine in the form of metabolites; 57% (range, 51 to 64%) was excreted in the feces.

The mean terminal half-life of finasteride in subjects \geq 70 years of age was approximately 8 hours (range, 6 to 15 hours; n=12), compared with 6 hours (range, 4 to 12 hours; n=12) in subjects 45 to 60 years of age. As a result, mean AUC(0-24 hr) after 17 days of dosing was 15% higher in subjects \geq 70 years of age than in subjects 45 to 60 years of age (p=0.02).

Mean (\pm SD)

Table 3:

Mean (SD) Pharmacokinetic Parameters

in Healthy Young Subjects (n=15)

Bioavailability

63% (34 to 108%)*

Clearance (mL/min)

165 (55)

Volume of Distribution (L)

76 (14)

Half-Life (hours)

6.2 (2.1)

*Range

Pediatric

Finasteride pharmacokinetics have not been investigated in patients <18 years of age.

Finasteride is not indicated for use in pediatric patients [see WARNINGS AND PRECAUTIONS (5.4) , USE IN SPECIFIC POPULATIONS (8.4)].

Gender

Finasteride is not indicated for use in females [see CONTRAINDICATIONS (4) , Warnings

and Precautions (5.3 and 5.4), USE IN SPECIFIC POPULATIONS (8.1) , and HOW SUPPLIED/STORAGE AND HANDLING (16)].

Geriatric

No dosage adjustment is necessary in the elderly. Although the elimination rate of finasteride is decreased in the elderly, these findings are of no clinical significance. [See CLINICAL PHARMACOLOGY (12.3) and USE IN SPECIFIC POPULATIONS (8.5) .]

Mean (\pm SD)

45 to 60 years old (n=12) \geq 70 years old (n=12)

*First-dose values; all other parameters are last-dose values

Table 4:

Mean (SD) Noncompartmental Pharmacokinetic Parameters

After Multiple Doses of 5 mg/day in Older Men

AUC (ng•hr/mL)

389 (98)

463 (186)

Peak Concentration (ng/mL)

46.2 (8.7)

48.4 (14.7)

Time to Peak (hours)

1.8 (0.7)

1.8 (0.6)

Half-Life (hours)*

6.0 (1.5)

8.2 (2.5)

Race

The effect of race on finasteride pharmacokinetics has not been studied.

Hepatic Impairment

The effect of hepatic impairment on finasteride pharmacokinetics has not been studied. Caution should be exercised in the administration of finasteride tablets in those patients with liver function abnormalities, as finasteride is metabolized extensively in the liver.

Renal Impairment

No dosage adjustment is necessary in patients with renal impairment. In patients with chronic renal impairment, with creatinine clearances ranging from 9.0 to 55 mL/min, AUC, maximum plasma concentration, half-life, and protein binding after a single dose of ¹⁴C-finasteride were similar to values obtained in healthy volunteers. Urinary excretion of metabolites was decreased in patients with renal impairment. This decrease was associated with an increase in fecal excretion of metabolites. Plasma concentrations of metabolites were significantly higher in patients with renal impairment (based on a 60% increase in total radioactivity AUC).

However, finasteride has been well tolerated in BPH patients with normal renal function receiving up to 80 mg/day for 12 weeks, where exposure of these patients to metabolites would presumably be much greater.

13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility

Carcinogenesis

No evidence of a tumorigenic effect was observed in a 24-month study in Sprague-Dawley rats receiving doses of finasteride up to 160 mg/kg/day in males and 320 mg/kg/day in females. These doses produced respective systemic exposure in rats of 111 and 274 times those observed in man receiving the recommended human dose of 5 mg/day. All exposure calculations were based on calculated AUC (0-24 hr) for animals and mean AUC(0-24 hr) for man (0.4 mcg•hr/mL).

In a 19-month carcinogenicity study in CD-1 mice, a statistically significant ($p \leq 0.05$) increase in the incidence of testicular Leydig cell adenomas was observed at 228 times the human exposure (250 mg/kg/day). In mice at 23 times the human exposure, estimated (25 mg/kg/day) and in rats at 39 times the human exposure (40 mg/kg/day) an increase in the incidence of Leydig cell hyperplasia was observed. A positive correlation between the proliferative changes in the Leydig cells and an increase in serum LH levels (2- to 3-fold above control) has been demonstrated in both rodent species treated with high doses of finasteride. No drug-related Leydig cell changes were seen in either rats or dogs treated with finasteride for 1 year at 30 and 350 times (20 mg/kg/day and 45 mg/kg/day, respectively) or in mice treated for 19 months at 2.3 times the human exposure, estimated (2.5 mg/kg/day).

Mutagenesis

No evidence of mutagenicity was observed in an in vitro bacterial mutagenesis assay, a mammalian cell mutagenesis assay, or in an in vitro alkaline elution assay. In an in vitro chromosome aberration assay, using Chinese hamster ovary cells, there was a slight increase in chromosome aberrations. These concentrations correspond to 4,000 to 5,000 times the peak plasma levels in man given a total dose of 5 mg. In an in vivo chromosome aberration assay in mice, no treatment-related increase in chromosome aberration was observed with finasteride at the maximum tolerated dose of 250 mg/kg/day (228 times the human exposure) as determined in the carcinogenicity studies.

Impairment of Fertility

In sexually mature male rabbits treated with finasteride at 543 times the human exposure (80 mg/kg/day) for up to 12 weeks, no effect on fertility, sperm count, or ejaculate volume was seen. In sexually mature male rats treated with 61 times the human exposure (80 mg/kg/day), there were no significant effects on fertility after 6 or 12 weeks of treatment; however, when treatment was continued for up to 24 or 30 weeks, there was an apparent decrease in fertility, fecundity and an associated significant decrease in the weights of the seminal vesicles and prostate. All these effects were reversible within 6 weeks of discontinuation of treatment. No drug-related effect on testes or on mating performance has been seen in rats or rabbits. This decrease in fertility in finasteride-treated rats is secondary to its effect on accessory sex organs (prostate and seminal vesicles) resulting in failure to form a seminal plug. The seminal plug is essential for normal fertility in rats and is not relevant in man.

14.1 Monotherapy

Finasteride tablets 5 mg/day was initially evaluated in patients with symptoms of BPH and enlarged prostates by digital rectal examination in two 1-year, placebo-controlled,

randomized, double-blind studies and their 5-year open extensions.

Finasteride tablets was further evaluated in the finasteride tablets A Long-Term Efficacy and Safety Study, a double-blind, randomized, placebo-controlled, 4-year, multicenter study. 3,040 patients between the ages of 45 and 78, with moderate to severe symptoms of BPH and an enlarged prostate upon digital rectal examination, were randomized into the study (1,524 to finasteride, 1,516 to placebo) and 3,016 patients were evaluable for efficacy. 1,883 patients completed the 4-year study (1,000 in the finasteride group, 883 in the placebo group).

Effect on Symptom Score

Symptoms were quantified using a score similar to the American Urological Association Symptom Score, which evaluated both obstructive symptoms (impairment of size and force of stream, sensation of incomplete bladder emptying, delayed or interrupted urination) and irritative symptoms (nocturia, daytime frequency, need to strain or push the flow of urine) by rating on a 0 to 5 scale for six symptoms and a 0 to 4 scale for one symptom, for a total possible score of 34.

Patients in A Long-Term Efficacy and Safety Study had moderate to severe symptoms at baseline (mean of approximately 15 points on a 0 to 34 point scale). Patients randomized to finasteride tablets who remained on therapy for 4 years had a mean (\pm 1 SD) decrease in symptom score of 3.3 (\pm 5.8) points compared with 1.3 (\pm 5.6) points in the placebo group. (See Figure 1.) A statistically significant improvement in symptom score was evident at 1 year in patients treated with finasteride tablets vs placebo (-2.3 vs -1.6), and this improvement continued through Year 4.

Figure 1 Symptom Score in A Long-Term Efficacy and Safety Study

[Finasteride-Fig.1]

Results seen in earlier studies were comparable to those seen in A Long-Term Efficacy and Safety Study. Although an early improvement in urinary symptoms was seen in some patients, a therapeutic trial of at least 6 months was generally necessary to assess whether a beneficial response in symptom relief had been achieved. The improvement in BPH symptoms was seen during the first year and maintained throughout an additional 5 years of open extension studies.

Effect on Acute Urinary Retention and the Need for Surgery

In A Long-Term Efficacy and Safety Study, efficacy was also assessed by evaluating treatment failures. Treatment failure was prospectively defined as BPH-related urological events or clinical deterioration, lack of improvement and/or the need for alternative therapy. BPH-related urological events were defined as urological surgical intervention and acute urinary retention requiring catheterization. Complete event information was available for 92% of the patients. The following table (Table 5) summarizes the results.

Patients (%)*
Event Placebo
N=1503 Finasteride

N=1513 Relative
Risk† 95% CI P
Value†

Table 5: All Treatment Failures in A Long-Term Efficacy and Safety Study

All Treatment Failures

37.1

26.2

0.68

(0.57 to 0.79)

<0.001

Surgical Interventions for BPH

10.1

4.6

0.45

(0.32 to 0.63)

<0.001

Acute Urinary
Retention Requiring
Catheterization

6.6

2.8

0.43

(0.28 to 0.66)

< 0.001

Two consecutive symptom scores ≥ 20

9.2

6.7

Bladder Stone

0.4

0.5

Incontinence

2.1

1.7

Renal Failure

0.5

0.6

UTI
5.7
4.9

Discontinuation due to worsening of BPH, lack of improvement, or to receive other medical treatment
21.8
13.3

*patients with multiple events may be counted more than once for each type of event
†Hazard ratio based on log rank test

Compared with placebo, finasteride tablets was associated with a significantly lower risk for acute urinary retention or the need for BPH-related surgery [13.2% for placebo vs 6.6% for finasteride tablets; 51% reduction in risk, 95% CI: (34 to 63%)]. Compared with placebo, finasteride tablets was associated with a significantly lower risk for surgery [10.1% for placebo vs 4.6% for finasteride tablets; 55% reduction in risk, 95% CI: (37 to 68%)] and with a significantly lower risk of acute urinary retention [6.6% for placebo vs 2.8% for finasteride tablets; 57% reduction in risk, 95% CI: (34 to 72%)]; see Figures 2 and 3.

Figure 2 Percent of Patients Having Surgery for BPH, Including TURP

Figure 3 Percent of Patients Developing Acute Urinary Retention (Spontaneous and Precipitated)
[Finasteride-fig.3]

Effect on Maximum Urinary Flow Rate

In the patients in A Long-Term Efficacy and Safety Study who remained on therapy for the duration of the study and had evaluable urinary flow data, finasteride tablets increased maximum urinary flow rate by 1.9 mL/sec compared with 0.2 mL/sec in the placebo group.

There was a clear difference between treatment groups in maximum urinary flow rate in favor of finasteride tablets by month 4 (1.0 vs 0.3 mL/sec) which was maintained throughout the study. In the earlier 1-year studies, increase in maximum urinary flow rate was comparable to A Long-Term Efficacy and Safety Study and was maintained through the first year and throughout an additional 5 years of open extension studies.

Effect on Prostate Volume

In A Long-Term Efficacy and Safety Study, prostate volume was assessed yearly by magnetic resonance imaging (MRI) in a subset of patients. In patients treated with finasteride tablets who remained on therapy, prostate volume was reduced compared with both baseline and placebo throughout the 4-year study. Finasteride tablets decreased prostate volume by 17.9% (from 55.9 cc at baseline to 45.8 cc at 4 years) compared with an increase of 14.1% (from 51.3 cc to 58.5 cc) in the placebo group ($p < 0.001$). (See Figure 4.)

Results seen in earlier studies were comparable to those seen in A Long-Term Efficacy and Safety Study. Mean prostate volume at baseline ranged between 40 to 50 cc. The reduction in prostate volume was seen during the first year and maintained throughout an additional five years of open extension studies.

Figure 4 Prostate Volume in A Long-Term Efficacy and Safety Study

[finasteride-fig4a]

Prostate Volume as a Predictor of Therapeutic Response

A meta-analysis combining 1-year data from seven double-blind, placebo-controlled studies of similar design, including 4,491 patients with symptomatic BPH, demonstrated that, in patients treated with finasteride tablets, the magnitude of symptom response and degree of improvement in maximum urinary flow rate were greater in patients with an enlarged prostate at baseline.

14.2 Combination with Alpha-Blocker Therapy

The Medical Therapy of Prostatic Symptoms (MTOPS) Trial was a double-blind, randomized, placebo-controlled, multicenter, 4- to 6-year study (average 5 years) in 3,047 men with symptomatic BPH, who were randomized to receive finasteride tablets 5 mg/day ($n=768$), doxazosin 4 or 8 mg/day ($n=756$), the combination of finasteride tablets 5 mg/day and doxazosin 4 or 8 mg/day ($n=786$), or placebo ($n=737$).

All participants underwent weekly titration of doxazosin (or its placebo) from 1 to 2 to 4 to 8 mg/day. Only those who tolerated the 4 or 8 mg dose level were kept on doxazosin (or its placebo) in the study. The participant's final tolerated dose (either 4 mg or 8 mg) was administered beginning at end-Week 4. The final doxazosin dose was administered once per day, at bedtime.

The mean patient age at randomization was 62.6 years (± 7.3 years). Patients were Caucasian (82%), African American (9%), Hispanic (7%), Asian (1%) or Native American ($< 1\%$). The mean duration of BPH symptoms was 4.7 years (± 4.6 years). Patients had moderate to severe BPH symptoms at baseline with a mean AUA symptom score of approximately 17 out of 35 points. Mean maximum urinary flow rate was 10.5 mL/sec (± 2.6 mL/sec). The mean prostate volume as measured by transrectal ultrasound was 36.3 mL (± 20.1 mL).

Prostate volume was ≤ 20 mL in 16% of patients, ≥ 50 mL in 18% of patients and between 21 and 49 mL in 66% of patients.

The primary endpoint was a composite measure of the first occurrence of any of the following five outcomes: a ≥ 4 point confirmed increase from baseline in symptom score, acute urinary retention, BPH-related renal insufficiency (creatinine rise), recurrent urinary tract infections or urosepsis, or incontinence. Compared to placebo, treatment with finasteride tablets, doxazosin, or combination therapy resulted in a reduction in the risk of experiencing one of these five outcome events by 34% ($p=0.002$), 39%

($p < 0.001$), and 67% ($p < 0.001$), respectively. Combination therapy resulted in a significant reduction in the risk of the primary endpoint compared to treatment with finasteride tablets alone (49%; $p \leq 0.001$) or doxazosin alone (46%; $p \leq 0.001$). (See Table 6.)

Table 6: Count and Percent Incidence of Primary Outcome Events by Treatment Group in MTOPS

Treatment Group

Placebo

N=737

Doxazosin

N=756

Finasteride

N=768

Combination

N=786

Total

N=3,047

Event

N (%)

N (%)

N (%)

N (%)

N (%)

AUA 4-point rise 100 (13.6)

59 (7.8)

74 (9.6)

41 (5.2)

274 (9.0)

Acute urinary

retention 18 (2.4) 13 (1.7)

6 (0.8)

4 (0.5)

41 (1.3)

Incontinence 8 (1.1)

11 (1.5)

9 (1.2)

3 (0.4)

31 (1.0)

Recurrent UTI/urosepsis 2 (0.3)

2 (0.3)

0 (0.0)

1 (0.1)

5 (0.2)

Creatinine rise 0 (0.0)

0 (0.0)

0 (0.0)

0 (0.0)
0 (0.0)
Total Events 128 (17.4)
85 (11.2)
89 (11.6)
49 (6.2)
351 (11.5)

The majority of the events (274 out of 351; 78%) was a confirmed ≥ 4 point increase in symptom score, referred to as symptom score progression. The risk of symptom score progression was reduced by 30% ($p=0.016$), 46% ($p<0.001$), and 64% ($p<0.001$) in patients treated with finasteride tablets, doxazosin, or the combination, respectively, compared to patients treated with placebo (see Figure 5). Combination therapy significantly reduced the risk of symptom score progression compared to the effect of finasteride tablets alone ($p<0.001$) and compared to doxazosin alone ($p=0.037$).

Figure 5
Cumulative Incidence of a 4-Point Rise in AUA Symptom Score by Treatment Group

[Finasteride-Fig.5]

Treatment with finasteride tablets, doxazosin or the combination of finasteride tablets with doxazosin, reduced the mean symptom score from baseline at year 4. Table 7 provides the mean change from baseline for AUA symptom score by treatment group for patients who remained on therapy for four years.

Table 7: Change From Baseline in AUA Symptom Score by Treatment Group at Year 4 in MTOPS

Placebo
N=534 Doxazosin
N=582 Finasteride
N=565 Combination
N=598
Baseline Mean (SD)
16.8 (6.0)
17.0 (5.9)
17.1 (6.0)
16.8 (5.8)
Mean Change
AUA Symptom Score (SD)
- 4.9 (5.8)
- 6.6 (6.1)
- 5.6 (5.9)
-7.4 (6.3)
Comparison to

Placebo (95% CI)

- 1.8

(-2.5, -1.1)

- 0.7

(-1.4, 0.0)

-2.5

(-3.2, -1.8)

Comparison to

Doxazosin alone (95% CI)

-0.7

(-1.4, 0.0)

Comparison to

Finasteride alone (95% CI)

-1.8

(-2.5, -1.1)

The results of MTOPS are consistent with the findings of the 4-year, placebo-controlled study A Long-Term Efficacy and Safety Study [see CLINICAL STUDIES (14.1)] in that treatment with finasteride tablets reduces the risk of acute urinary retention and the need for BPH-related surgery. In MTOPS, the risk of developing acute urinary retention was reduced by 67% in patients treated with finasteride tablets compared to patients treated with placebo (0.8% for finasteride tablets and 2.4% for placebo). Also, the risk of requiring BPH-related invasive therapy was reduced by 64% in patients treated with finasteride tablets compared to patients treated with placebo (2.0% for finasteride tablets and 5.4% for placebo).

14.3 Summary of Clinical Studies

The data from these studies, showing improvement in BPH-related symptoms, reduction in treatment failure (BPH-related urological events), increased maximum urinary flow rates, and decreasing prostate volume, suggest that finasteride tablets arrests the disease process of BPH in men with an enlarged prostate.

Finasteride tablets, USP 5 mg are available as blue colored, 7 mm round, biconvex, film coated tablets, marked "F5" on one side and plain on other side. They are supplied as follows:

NDC 67877-288-30, bottles of 30 tablets

NDC 72198-542-90, bottles of 90 tablets

NDC 67877-288-01, bottles of 100 tablets

NDC 67877-288-05, bottles of 500 tablets

NDC 67877-288-10, bottles of 1000 tablets

NDC 67877-288-33, Carton pack of 10 unit-dose tablets

Storage and Handling

Store at room temperatures 20°C to 25°C (68°F to 77°F). [See USP Controlled Room Temperature]. Protect from light and keep container tightly closed.

Females should not handle crushed or broken finasteride tablets when they are pregnant or may potentially be pregnant because of the possibility of absorption of finasteride and the subsequent potential risk to a male fetus [see WARNINGS AND PRECAUTIONS (5.3) and USE IN SPECIFIC POPULATIONS (8.1)].

Advise the patient to read the FDA-approved patient labeling (Patient Information).

17.1 Increased Risk of High-Grade Prostate Cancer

Patients should be informed that there was an increase in high-grade prostate cancer in men treated with 5 α -reductase inhibitors indicated for BPH treatment, including finasteride tablets, compared to those treated with placebo in studies looking at the use of these drugs to prevent prostate cancer [see INDICATIONS AND USAGE (1.3), WARNINGS AND PRECAUTIONS (5.2), and ADVERSE REACTIONS (6.1)].

17.2 Exposure of Females-Risk to Male Fetus

Physicians should inform patients that females who are pregnant or may potentially be pregnant should not handle crushed or broken finasteride tablets because of the possibility of absorption of finasteride and the subsequent potential risk to the male fetus.

Finasteride tablets are coated and will prevent contact with the active ingredient during normal handling, provided that the tablets have not been broken or crushed. If a female who is pregnant or may potentially be pregnant comes in contact with crushed or broken finasteride tablets, the contact area should be washed immediately with soap and water [see CONTRAINDICATIONS (4), WARNINGS AND PRECAUTIONS (5.3), USE IN SPECIFIC POPULATIONS (8.1) and HOW SUPPLIED/STORAGE AND HANDLING (16)].

17.3 Additional Instructions

Physicians should inform patients that the volume of ejaculate may be decreased in some patients during treatment with finasteride tablets.

This decrease does not appear to interfere with normal sexual function. However, impotence and decreased libido may occur in patients treated with finasteride tablets [see Adverse Reactions (6.1)].

Physicians should instruct their patients to promptly report any changes in their breasts such as lumps, pain or nipple discharge. Breast changes including breast enlargement, tenderness and neoplasm have been reported [see Adverse Reactions (6.1)].

Physicians should instruct their patients to read the patient package insert before starting therapy with finasteride tablets and to reread it each time the prescription is renewed so that they are aware of current information for patients regarding finasteride tablets.

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Manufactured by:

Alkem Laboratories Ltd.,

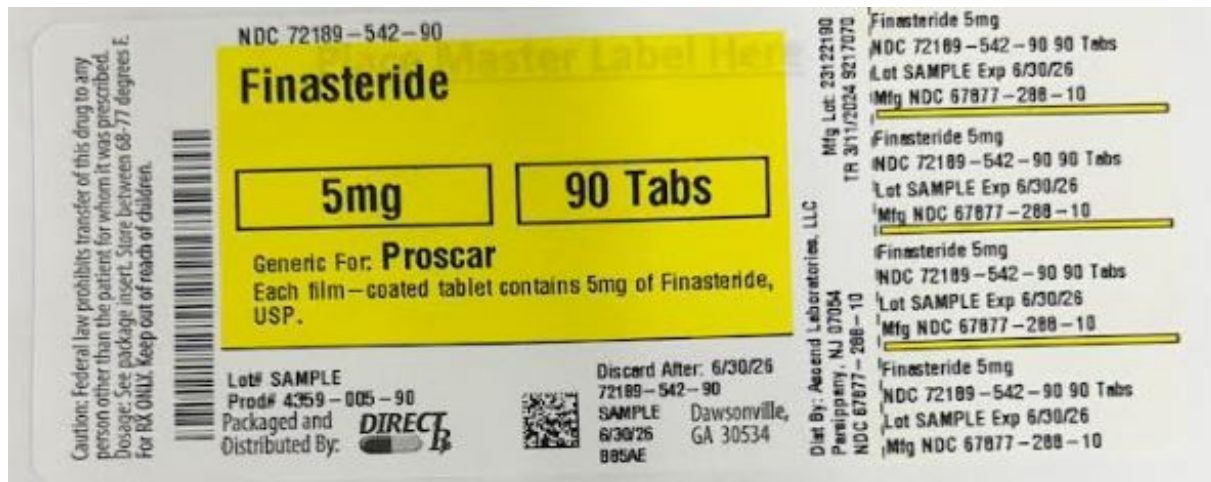
INDIA.

Distributed by:

Ascend Laboratories, LLC

Parsippany, NJ 07054

Revised: November, 2022



FINASTERIDE

finasteride tablet, film coated

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:72189-542(NDC:67877-288)
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
FINASTERIDE (UNII: 57GNO57U7G) (FINASTERIDE - UNII:57GNO57U7G)	FINASTERIDE	5 mg

Inactive Ingredients

Ingredient Name	Strength
STARCH, CORN (UNII: O8232NY3SJ)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
FD&C BLUE NO. 2 (UNII: L06K8R7DQK)	

Product Characteristics

Color	blue	Score	no score
Shape	ROUND	Size	7mm
Flavor		Imprint Code	F5

Contains**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72189-542-90	90 in 1 BOTTLE; Type 0: Not a Combination Product	03/11/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA204304	03/11/2024	

Labeler - Direct_Rx (079254320)**Registrant** - Direct_Rx (079254320)**Establishment**

Name	Address	ID/FEI	Business Operations
Direct_Rx		079254320	repack(72189-542)

Revised: 3/2024

Direct_Rx