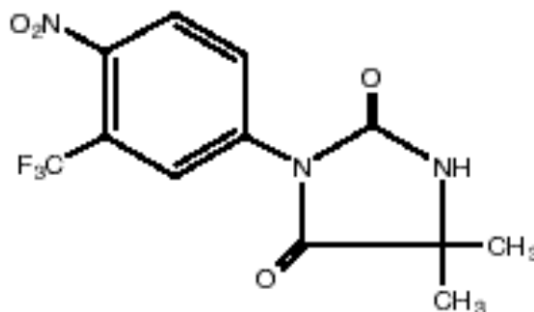


Rx only

NILANDRON[®]
(nilutamide)
Tablets

DESCRIPTION

NILANDRON[®] tablets contain nilutamide, a nonsteroidal, orally active antiandrogen having the chemical name 5,5-dimethyl-3-[4-nitro-3-(trifluoromethyl)phenyl]-2,4-imidazolidinedione with the following structural formula:



Nilutamide is a microcrystalline, white to practically white powder with a molecular weight of 317.25. Its molecular formula is C₁₂H₁₀F₃N₃O₄.

It is freely soluble in ethyl acetate, acetone, chloroform, ethyl alcohol, dichloromethane, and methanol. It is slightly soluble in water [$<0.1\%$ W/V at 25°C (77°F)]. It melts between 153°C and 156°C (307.4°F and 312.8°F).

Each NILANDRON tablet contains 150 mg of nilutamide. Other ingredients in NILANDRON tablets are corn starch, lactose, povidone, docusate sodium, magnesium stearate, and talc.

CLINICAL PHARMACOLOGY

Mechanism of Action

Prostate cancer is known to be androgen sensitive and responds to androgen ablation. In animal studies, nilutamide has demonstrated antiandrogenic activity without other hormonal (estrogen, progesterone, mineralocorticoid, and glucocorticoid) effects. In vitro, nilutamide blocks the effects of testosterone at the androgen receptor level. In vivo, nilutamide interacts with the androgen receptor and prevents the normal androgenic response.

Pharmacokinetics

Absorption:

Analysis of blood, urine, and feces samples following a single oral 150-mg dose of [¹⁴C]-nilutamide in patients with metastatic prostate cancer showed that the drug is rapidly and completely absorbed and that it yields high and persistent plasma concentrations.

Distribution:

After absorption of the drug, there is a detectable distribution phase. There is moderate binding of the drug to plasma proteins and low binding to erythrocytes. The binding is nonsaturable except in the case of alpha-1-glycoprotein, which makes a minor contribution to the total concentration of proteins in the plasma. The results of binding studies do not indicate any effects that would cause nonlinear pharmacokinetics.

Metabolism:

The results of a human metabolism study using ¹⁴C-radiolabelled tablets show that nilutamide is extensively metabolized and less than 2% of the drug is excreted unchanged in urine after 5 days. Five metabolites have been isolated from human urine. Two metabolites display an asymmetric center, due to oxidation of a methyl group, resulting in the formation of D- and L-isomers. One of the metabolites was shown, in vitro, to possess 25 to 50% of the pharmacological activity of the parent drug, and the D-isomer of the active metabolite showed equal or greater potency compared to the L-isomer. However, the pharmacokinetics and the pharmacodynamics of the metabolites have not been fully investigated.

Elimination:

The majority (62%) of orally administered [¹⁴C]-nilutamide is eliminated in the urine during the first 120 hours after a single 150-mg dose. Fecal elimination is negligible, ranging from 1.4% to 7% of the dose after 4 to 5 days. Excretion of radioactivity in urine likely continues beyond 5 days. The mean elimination half-life of nilutamide determined in studies in which subjects received a single dose of 100-300 mg ranged from 38.0 to 59.1 hours with most values between 41 and 49 hours. The elimination of at least one metabolite is generally longer than that of unchanged nilutamide (59-126 hours). During multiple dosing of 150 mg nilutamide (given as 3 x 50 mg) twice a day, steady state was reached within 2 to 4 weeks for most patients, and mean steady state AUC₀₋₁₂ was 110% higher than the AUC_{0-∞} obtained from the first 150 mg dose. These data and in vitro metabolism data suggest that, upon multiple dosing, metabolic enzyme inhibition may occur for this drug.

Clinical Studies

Nilutamide through its antiandrogenic activity can complement surgical castration, which suppresses only testicular androgens. The effects of the combined therapy were studied in patients with previously untreated metastatic prostate cancer.

In a double-blind, randomized, multicenter study that enrolled 457 patients (225 treated with orchiectomy and NILANDRON, 232 treated with orchiectomy and placebo), the NILANDRON group showed a statistically significant benefit in time to progression and time to death. The results are summarized below.

	NILANDRON	PLACEBO
Median Survival (months)	27.3	23.6
Progression-Free Survival (months)	21.1	14.9
Complete or Partial Regression	41%	24%
Improvement in Bone Pain	54%	37%

INDICATIONS AND USAGE

Metastatic Prostate Cancer

NILANDRON tablets are indicated for use in combination with surgical castration for the treatment of metastatic prostate cancer (Stage D₂).

For maximum benefit, NILANDRON treatment must begin on the same day as or on the day after surgical castration.

CONTRAINDICATIONS

NILANDRON tablets are contraindicated:

- in patients with severe hepatic impairment (baseline hepatic enzymes should be evaluated prior to treatment)

- in patients with severe respiratory insufficiency
- in patients with hypersensitivity to nilutamide or any component of this preparation.

WARNINGS

Interstitial Pneumonitis

Interstitial pneumonitis has been reported in 2% of patients in controlled clinical trials in patients exposed to nilutamide. A small study in Japanese subjects showed that 8 of 47 patients (17%) developed interstitial pneumonitis. Reports of interstitial changes including pulmonary fibrosis that led to hospitalization and death have been reported rarely post-marketing. Symptoms included exertional dyspnea, cough, chest pain, and fever. X-rays showed interstitial or alveolo-interstitial changes, and pulmonary function tests revealed a restrictive pattern with decreased DLco. Most cases occurred within the first 3 months of treatment with NILANDRON, and most reversed with discontinuation of therapy. A routine chest X-ray should be performed prior to initiating treatment with NILANDRON. Baseline pulmonary function tests may be considered. Patients should be instructed to report any new or worsening shortness of breath that they experience while on NILANDRON. **If symptoms occur, NILANDRON should be immediately discontinued until it can be determined if the symptoms are drug related.**

Hepatitis

Rare cases of death or hospitalization due to severe liver injury have been reported post-marketing in association with the use of NILANDRON. Hepatotoxicity in these reports generally occurred within the first 3 to 4 months of treatment. Hepatitis or marked increases in liver enzymes leading to drug discontinuation occurred in 1% of NILANDRON patients in controlled clinical trials.

Serum transaminase levels should be measured prior to starting treatment with NILANDRON, at regular intervals for the first 4 months of treatment, and periodically thereafter. Liver function tests should also be obtained at the first sign or symptom suggestive of liver dysfunction, e.g. nausea, vomiting, abdominal pain, fatigue, anorexia, “flu-like” symptoms, dark urine, jaundice, or right upper quadrant tenderness. If at any time, a patient has jaundice, or their ALT rises above 2 times the upper limit of normal, NILANDRON should be immediately discontinued with close followup of liver function tests until resolution.

Use in Women

NILANDRON has no indication for women, and should not be used in this population, particularly for non-serious or non-life threatening conditions.

Other

Foreign postmarketing surveillance has revealed isolated cases of aplastic anemia in which a causal relationship with NILANDRON could not be ascertained.

PRECAUTIONS

General

Antiandrogen Withdrawal Syndrome

Patients whose disease progresses while being treated with an antiandrogen may experience clinical improvement with discontinuation of the antiandrogen.

Information for Patients

Patients should be informed that NILANDRON tablets should be started on the day of, or on the day after, surgical castration. They should also be informed that they should not interrupt their dosing of NILANDRON or stop taking this medication without consulting their physician.

Because of the possibility of interstitial pneumonitis, patients should also be told to report immediately any dyspnea or aggravation of pre-existing dyspnea.

Because of the possibility of hepatitis, patients should be told to consult with their physician should nausea, vomiting, abdominal pain, or jaundice occur.

Because of the possibility of an intolerance to alcohol (facial flushes, malaise, hypotension) following ingestion of NILANDRON, it is recommended that intake of alcoholic beverages be avoided by patients who experience this reaction. This effect has been reported in about 5% of patients treated with NILANDRON.

In clinical trials, 13% to 57% of patients receiving NILANDRON reported a delay in adaptation to dark, ranging from seconds to a few minutes, when passing from a lighted area to a dark area. This effect sometimes does not abate as drug treatment is continued. Patients who experience this effect should be cautioned about driving at night or through tunnels. This effect can be alleviated by the wearing of tinted glasses.

Drug Interactions

In vitro, nilutamide has been shown to inhibit the activity of liver cytochrome P-450 isoenzymes and, therefore, may reduce the metabolism of compounds requiring these systems.

Consequently, drugs with a low therapeutic margin, such as vitamin K antagonists, phenytoin, and theophylline, could have a delayed elimination and increases in their serum half-life leading to a toxic level. The dosage of these drugs or others with a similar metabolism may need to be modified if they are administered concomitantly with nilutamide. For example, when vitamin K antagonists are administered concomitantly with nilutamide, prothrombin time should be carefully monitored and, if necessary, the dosage of vitamin K antagonists should be reduced.

Carcinogenesis, Mutagenesis, Impairment of Fertility

Administration of nilutamide to rats for 18 months at doses of 0, 5, 15, or 45 mg/kg/day produced benign Leydig cell tumors in 35% of the high-dose male rats (AUC exposures in high-dose rats were approximately 1 - 2 times human AUC exposures with therapeutic doses). The increased incidence of Leydig cell tumors is secondary to elevated luteinizing hormone (LH) concentrations resulting from loss of feedback inhibition at the pituitary. Elevated LH and testosterone concentrations are not observed in castrated men receiving NILANDRON. Nilutamide had no effect on the incidence, size, or time of onset of any spontaneous tumor in rats.

Nilutamide displayed no mutagenic effects in a variety of in vitro and in vivo tests (Ames test, mouse micronucleus test, and two chromosomal aberration tests).

In reproduction studies in rats, nilutamide had no effect on the reproductive function of males and females, and no lethal, teratogenic, or growth-suppressive effects on fetuses were found. The maximal dose at which nilutamide did not affect reproductive function in either sex or have an effect on fetuses

was estimated to be 45 mg/kg orally (AUC exposures in rats approximately 1-2 times human therapeutic AUC exposures).

Pregnancy

Animal reproduction studies have not been conducted with nilutamide. It is also not known whether nilutamide can cause fetal harm when administered to a pregnant woman or can affect reproductive capacity. Nilutamide should be given to a pregnant woman only if clearly needed.

Pediatric Use

Safety and effectiveness in pediatric patients have not been determined.

Animal Pharmacology and Toxicology

Administration of NILANDRON to beagle dogs resulted in drug-related deaths at dose levels that produce AUC exposures in dogs much lower than the AUC exposures of men receiving the therapeutic doses of 150 and 300 mg/day. Nilutamide-induced toxicity in dogs was cumulative with progressively lower doses producing death when given for longer durations. Nilutamide given to dogs at 60 mg/kg/day (1-2 times human AUC exposure) for 1 month produced 100% mortality. Administration of 20 and 30 mg/kg/day nilutamide (1/2-1 times human AUC exposure) for 6 months resulted in 20% and 70% mortality in treated dogs. Administration to dogs of 3, 6, and 12 mg/kg/day nilutamide (1/10-1/2 human AUC exposure) for 1 year resulted in 8%, 33%, and 50% mortality, respectively. A **“no-effect level” for nilutamide-induced mortality in dogs was not identified.** Pathology data from the one-year oral toxicity study suggest that the deaths in dogs were secondary to liver toxicity. Marked-to-massive hepatocellular swelling and vacuolization were observed in affected dogs. Liver toxicity in dogs was not consistently associated with elevations of liver enzymes.

Administration of nilutamide to rats at a dose level of 45 mg/kg/day (AUC exposure in rats 1-2 times human therapeutic AUC exposures) for 18 months increased the incidence of lung pathology (granulomatous inflammation and chronic alveolitis).

The hepatic and pulmonary adverse effects observed in nilutamide-treated animals and men are similar to effects observed with another nitroaromatic compound, nitrofurantoin. Nilutamide and nitrofurantoin are both metabolized in vitro to nitroanion free-radicals by microsomal NADPH-cytochrome P450 reductase in the lungs and liver of rats and humans.

ADVERSE REACTIONS

Clinical Trial Experience

The following adverse experiences were reported during a multicenter clinical trial comparing NILANDRON + surgical castration versus placebo + surgical castration. The most frequently reported (greater than 5%) adverse experiences during treatment with NILANDRON tablets in combination with surgical castration are listed below. For comparison, adverse experiences seen with surgical castration and placebo are also listed.

Adverse Experience	NILANDRON + surgical castration (N=225) % All	Placebo + surgical castration (N=232) % All
Cardiovascular System		
Hypertension	5.3	2.6
Digestive System		
Nausea	9.8	6.0
Constipation	7.1	3.9
Endocrine System		
Hot flushes	28.4	22.4
Metabolic and Nutritional System		
Increased AST	8.0	3.9
Increased ALT	7.6	4.3
Nervous System		
Dizziness	7.1	3.4
Respiratory System		
Dyspnea	6.2	7.3
Special Senses		
Impaired adaptation to dark	12.9	1.3
Abnormal vision	6.7	1.7
Urogenital System		
Urinary tract infection	8.0	9.1

The overall incidence of adverse experiences was 86% (194/225) for the NILANDRON group and 81% (188/232) for the placebo group.

The following adverse experiences were reported during a multicenter clinical trial comparing NILANDRON + leuprolide versus placebo + leuprolide. The most frequently reported (greater than 5%) adverse experiences during treatment with NILANDRON tablets in combination with leuprolide are listed below. For comparison, adverse experiences seen with leuprolide and placebo are also listed.

Adverse Experience	NILANDRON + leuprolide (N=209) % All	Placebo + leuprolide (N=202) % All
Body as a Whole		
Pain	26.8	27.7
Headache	13.9	10.4
Asthenia	19.1	20.8
Back pain	11.5	16.8
Abdominal pain	10.0	5.4
Chest pain	7.2	4.5
Flu syndrome	7.2	3.0
Fever	5.3	6.4
Cardiovascular System		
Hypertension	9.1	9.9
Digestive System		
Nausea	23.9	8.4
Constipation	19.6	16.8

Anorexia	11.0	6.4
Dyspepsia	6.7	4.5
Vomiting	5.7	4.0
Endocrine System		
Hot flushes	66.5	59.4
Impotence	11.0	12.9
Libido decreased	11.0	4.5
Hemic and Lymphatic System		
Anemia	7.2	6.4
Metabolic and Nutritional System		
Increased AST	12.9	13.9
Peripheral edema	12.4	17.3
Increased ALT	9.1	8.9
Musculoskeletal System		
Bone Pain	6.2	5.0
Nervous System		
Insomnia	16.3	15.8
Dizziness	10.0	11.4
Depression	8.6	7.4
Hypesthesia	5.3	2.0
Respiratory System		
Dyspnea	10.5	7.4
Upper respiratory infection	8.1	10.9
Pneumonia	5.3	3.5
Skin and Appendages		
Sweating	6.2	3.0
Body hair loss	5.7	0.5
Dry skin	5.3	2.5
Rash	5.3	4.0
Special Senses		
Impaired adaptation to dark	56.9	5.4
Chromatopsia	8.6	0.0
Impaired adaptation to light	7.7	1.0
Abnormal vision	6.2	4.5
Urogenital System		
Testicular atrophy	16.3	12.4
Gynecomastia	10.5	11.9
Urinary tract infection	8.6	21.3
Hematuria	8.1	7.9
Urinary tract disorder	7.2	10.4
Nocturia	6.7	6.4

The overall incidence of adverse experiences is 99.5% (208/209) for the NILANDRON group and 98.5% (199/202) for the placebo group.

Some frequently occurring adverse experiences, for example hot flushes, impotence, and decreased libido, are known to be associated with low serum androgen levels and known to occur with medical or surgical castration alone. Notable was the higher incidence of visual disturbances (variously described as impaired adaptation to darkness, abnormal vision, and colored vision), which led to treatment discontinuation in 1% to 2% of patients.

Interstitial pneumonitis occurred in one (<1%) patient receiving NILANDRON in combination with surgical castration and in seven patients (3%) receiving NILANDRON in combination with leuprolide and one patient receiving placebo in combination with leuprolide. Overall, it has been reported in 2% of

patients receiving NILANDRON. This included a report of interstitial pneumonitis in 8 of 47 patients (17%) in a small study performed in Japan.

In addition, the following adverse experiences were reported in 2 to 5% of patients treated with NILANDRON in combination with leuprolide or orchiectomy.

Body as a Whole: Malaise (2%).

Cardiovascular System: Angina (2%), heart failure (3%), syncope (2%).

Digestive System: Diarrhea (2%), gastrointestinal disorder (2%), gastrointestinal hemorrhage (2%), melena (2%).

Metabolic and Nutritional System: Alcohol intolerance (5%), edema (2%), weight loss (2%).

Musculoskeletal System: Arthritis (2%).

Nervous System: Dry mouth (2%), nervousness (2%), paresthesia (3%).

Respiratory System: Cough increased (2%), interstitial lung disease (2%), lung disorder (4%), rhinitis (2%).

Skin and Appendages: Pruritus (2%).

Special Senses: Cataract (2%), photophobia (2%).

Laboratory Values: Haptoglobin increased (2%), leukopenia (3%), alkaline phosphatase increased (3%), BUN increased (2%), creatinine increased (2%), hyperglycemia (4%).

To report SUSPECTED ADVERSE REACTIONS, contact Concordia Pharmaceuticals Inc. at 1-877-370-1142 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

OVERDOSAGE

One case of massive overdosage has been published. A 79-year-old man attempted suicide by ingesting 13 g of nilutamide (i.e., 43 times the maximum recommended dose). Despite immediate gastric lavage and oral administration of activated charcoal, plasma nilutamide levels peaked at 6 times the normal range 2 hours after ingestion. There were no clinical signs or symptoms or changes in parameters such as transaminases or chest X-ray. Maintenance treatment (150 mg/day) was resumed 30 days later.

In repeated-dose tolerance studies, doses of 600 mg/day and 900 mg/day were administered to 9 and 4 patients, respectively. The ingestion of these doses was associated with gastrointestinal disorders, including nausea and vomiting, malaise, headache, and dizziness. In addition, a transient elevation in hepatic enzyme levels was noted in one patient.

Since nilutamide is protein bound, dialysis may not be useful as treatment for overdose. As in the management of overdosage with any drug, it should be borne in mind that multiple agents may have been taken. If vomiting does not occur spontaneously, it should be induced if the patient is alert. General supportive care, including frequent monitoring of the vital signs and close observation of the patient, is indicated.

DOSAGE AND ADMINISTRATION

The recommended dosage is 300 mg once a day for 30 days, followed thereafter by 150 mg once a day. NILANDRON tablets can be taken with or without food.

HOW SUPPLIED

NILANDRON 150 mg tablets are supplied in boxes of 30 tablets. Each box contains 3 child-resistant, PVC, aluminum foil-backed blisters of 10 tablets (NDC 59212-111-14). Each white, biconvex, cylindrical (10 mm in diameter) tablet has a triangular logo on one side and an internal reference number (168D) on the other.

Store at 25°C (77°F); excursions permitted between 15-30°C (59-86°F) [see USP Controlled Room Temperature]. Protect from light. Keep out of reach of children.

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Manufactured for:
Concordia Pharmaceuticals Inc.
St. Michael, Barbados BB11005

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