

PRODUCT MONOGRAPH

ANDROCUR[®]

cyproterone acetate tablets

50 mg

ANDROCUR[®] DEPOT

cyproterone acetate injection

100 mg/mL

Antiandrogen

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Table of Contents

PART I: HEALTH PROFESSIONAL INFORMATION.....	3
SUMMARY PRODUCT INFORMATION	3
INDICATIONS AND CLINICAL USE.....	3
CONTRAINDICATIONS	3
WARNINGS AND PRECAUTIONS.....	4
ADVERSE REACTIONS.....	7
DRUG INTERACTIONS.....	8
DOSAGE AND ADMINISTRATION.....	9
OVERDOSAGE	10
ACTION AND CLINICAL PHARMACOLOGY	10
DOSAGE FORMS, COMPOSITION AND PACKAGING	11
PART II: SCIENTIFIC INFORMATION	12
PHARMACEUTICAL INFORMATION.....	12
CLINICAL TRIALS.....	12
DETAILED PHARMACOLOGY	16
TOXICOLOGY	20
REFERENCES	24
PART III: CONSUMER INFORMATION.....	27

ANDROCUR[®]

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PART I: HEALTH PROFESSIONAL INFORMATION

SUMMARY PRODUCT INFORMATION

Table 1 – Product Information Summary

Route of Administration	Dosage Form, Strength	Clinically Relevant Nonmedicinal Ingredients
Oral	Tablet, 50 mg	<i>For a complete listing see DOSAGE FORMS, COMPOSITION AND PACKAGING section.</i>
Intramuscular	Intramuscular solution, 100 mg/mL	<i>For a complete listing see DOSAGE FORMS, COMPOSITION AND PACKAGING section.</i>

INDICATIONS AND CLINICAL USE

ANDROCUR (cyproterone acetate) is indicated for:

- the palliative treatment of patients with advanced prostatic carcinoma

CONTRAINDICATIONS

- Patients who are hypersensitive to this drug or to any ingredient in the formulation or component of the container. For a complete listing, see the **DOSAGE FORMS, COMPOSITION AND PACKAGING** section.
- Liver disease and hepatic dysfunction.
- Renal insufficiency.
- Dubin Johnson syndrome, Rotor syndrome
- Previous or existing liver tumors (only if these are not due to metastases from carcinoma of the prostate)
- Wasting diseases (with the exception of inoperable carcinoma of the prostate)
- Severe chronic depression
- Existing thromboembolic processes

WARNINGS AND PRECAUTIONS

General

Concomitant Alcohol: Alcohol may reduce the antiandrogenic effect of ANDROCUR in hypersexuality. The relevance of this in prostatic carcinoma is not known; however, it would be prudent to inform the patients that the use of alcohol during ANDROCUR therapy is not advisable.

Physical Performance: Patients should be informed that fatigue and lassitude are common in the first few weeks of therapy, but usually become much less pronounced from the third month on. Marked lassitude and asthenia necessitate special care when driving or operating machinery.

Other Conditions: Like all oily solutions, ANDROCUR DEPOT must be injected intramuscularly. Experience shows that the short-lasting reactions (urge to cough, coughing fits, respiratory distress) which occur in rare cases during or immediately after the injection of oily solutions can be avoided by injecting the solution extremely slowly.

Carcinogenesis and Mutagenesis

Cyproterone acetate showed a potential to initiate and/or promote liver tumor formation in rodents. Very rare cases of benign and malignant liver tumors have been observed in patients receiving ANDROCUR.

Meningioma: Cerebral meningiomas (single and multiple) have been reported in association with long-term use (several years) of ANDROCUR. If a patient treated with ANDROCUR is diagnosed with a meningioma, ANDROCUR must be stopped.

Antiandrogen Withdrawal Syndrome: In some patients with metastatic prostate cancer, antiandrogens (steroidal or nonsteroidal) may promote, rather than inhibit, the growth of prostate cancer. A decrease in PSA and/or clinical improvement following the discontinuation of antiandrogens has been reported. It is recommended that patients prescribed an antiandrogen, who have PSA progression, should have the antiandrogen discontinued immediately and be monitored for 6 to 8 weeks for a withdrawal response prior to any decision to proceed with other prostate cancer therapy.

Gynecomastia: Benign nodules (hyperplasia) of the breast have been reported; these generally subside 1 to 3 months after discontinuation of therapy and/or after a reduction of dosage. The reduction of dosage should be weighed against the risk of inadequate tumor control.

Endocrine and Metabolism

Adrenocortical Function: Suppression of adrenocortical function tests have occurred in patients receiving ANDROCUR and preclinical data also suggest a possible suppression due to the corticoid-like effect of ANDROCUR.

Reduced response to endogenous ACTH was noted by metyrapone test; furthermore, reduced ACTH and cortisol blood levels determined by the Mattingly method were also found.

It is therefore recommended that adrenocortical function tests should be monitored periodically by serum cortisol assay.

Diabetes: ANDROCUR may impair carbohydrate metabolism. Parameters of carbohydrate metabolism, fasting blood glucose and glucose tolerance tests, should be examined carefully in all patients and particularly in all diabetics before and regularly during therapy with ANDROCUR.

Strict medical supervision is necessary if the patient suffers from diabetes, because the requirement for oral antidiabetics or insulin can change during ANDROCUR treatment.

Metabolic Effects: Fluid retention, hypercalcemia and changes in plasma lipid profile may occur. Accordingly, ANDROCUR should be used with caution in patients with cardiac disease.

Nitrogen Balance: A negative nitrogen balance is usual at the start of therapy, but does generally correct itself within 3 months of continued therapy.

Hematologic

Hematology: Hypochromic anemia has been observed rarely during therapy with ANDROCUR. Regular hematological assessment is recommended.

Thromboembolism: Clinical investigations have shown that when ANDROCUR is used alone it has a minor effect on blood clotting factors. However, when ANDROCUR was combined with ethinyl estradiol, changes were found in increased coagulation capability.

The occurrence of thromboembolic events has been reported in patients using ANDROCUR, although a causal relationship has not been established. Patients with previous arterial or venous thrombotic / thromboembolic events (eg, deep venous thrombosis, pulmonary embolism, myocardial infarction) or with a history of cerebrovascular accidents or with advanced malignancies are at increased risk of further thromboembolic events.

ANDROCUR should be discontinued at the first sign of thrombophlebitis or thromboembolism, and the patient should be carefully re-evaluated if manifestations of thrombotic disorder occur: thrombophlebitis, cerebrovascular complications, retinal thrombosis, or pulmonary embolism.

In patients with inoperable carcinoma of the prostate, presenting with a history of thromboembolic processes or suffering from sickle cell anemia or from severe diabetes with vascular changes, a careful risk:benefit evaluation must be carried out in each individual case before ANDROCUR is prescribed.

Hepatic/Biliary/Pancreas

Liver Function: Direct hepatic toxicity, including jaundice, hepatitis, and hepatic failure, which has been fatal in some cases, has been reported in patients treated with 200 to 300 mg ANDROCUR. Most reported cases are in men with prostatic cancer. Toxicity is dose-related and develops, usually, several months after treatment has begun. Liver function tests should be performed pretreatment, at regular intervals during treatment, and whenever any symptoms or

signs suggestive of hepatotoxicity occur. If hepatotoxicity is confirmed, ANDROCUR should normally be withdrawn, unless the hepatotoxicity can be explained by another cause, eg, metastatic disease, in which case cyproterone acetate should be continued only if the perceived benefit outweighs the risk.

In very rare cases, benign and malignant liver tumors which may lead to life-threatening intra-abdominal hemorrhage have been observed after the use of ANDROCUR. If severe upper abdominal complaints, liver enlargement, or signs of intra-abdominal hemorrhage occur, a liver tumor should be included in the differential-diagnostic considerations.

Psychiatric

Depression: ANDROCUR therapy has occasionally been associated with an increased incidence of depressive mood changes, especially during the first 6 to 8 weeks of therapy. Similar mood changes have also been seen following surgical castration and are considered to be due to androgen deprivation. Patients with tendencies to depressive reaction should be carefully observed.

Respiratory

Shortness of Breath: A sensation of shortness of breath was commonly reported in patients treated with 300 mg/day ANDROCUR. Patients with pre-existing pulmonary dysfunction are most likely to be affected.

Sexual Function/Reproduction

Inhibition of Spermatogenesis: The sperm count and the volume of ejaculate are reduced at oral doses of 50 to 300 mg per day. Infertility is usual, and there may be azoospermia after 8 weeks of therapy, which is associated with atrophy of seminiferous tubules.

Follow-up examinations on discontinuation of therapy have shown these changes to be reversible.

Spermatogenesis usually reverts to its previous level about 3 to 5 months after stopping ANDROCUR, or in some patients, after up to 20 months. Production of abnormal spermatozoa during ANDROCUR therapy has been observed; their relationship to abnormal fertilization or malformed embryos is not known.

Skin

ANDROCUR therapy may cause a reduction of sebum production leading to dryness of the skin and transient patchy loss of body hair.

Special Populations

Pregnant Women

Treatment with ANDROCUR is not indicated for use in women.

Nursing Women

Treatment with ANDROCUR is not indicated for use in women.

Pediatrics

ANDROCUR is not recommended for use in children and adolescents below 18 years of age.

ANDROCUR must not be given before the conclusion of puberty since an unfavorable influence on longitudinal growth and the still unstabilized axes of endocrine function cannot be ruled out.

ADVERSE REACTIONS

Adverse Drug Reaction Overview

The adverse events associated most frequently with the use of ANDROCUR are those related to the hormonal effects of the drug. These reactions usually disappear upon discontinuation of therapy or reduction of dose: decreased libido, breast enlargement, breast tenderness, benign nodular hyperplasia of the breast, galactorrhea, gynecomastia, abnormal spermatozoa, impotence, and inhibition of spermatogenesis.

The most serious adverse drug reactions (ADRs) in patients receiving ANDROCUR are hepatic toxicity, benign and malignant liver tumors which may lead to intra-abdominal hemorrhage, and thromboembolic events.

As with other antiandrogenic treatments, long-term androgen deprivation with ANDROCUR may lead to osteoporosis.

Other adverse events which have been reported are listed below:

Cardiovascular System: hypotension, tachycardia, heart failure, syncope, myocardial infarct, hemorrhage, cerebrovascular accident, cardiovascular disorder, retinal vascular disorder, embolus, pulmonary embolism, superficial and deep thrombophlebitis, thrombosis, retinal vein thrombosis, phlebitis, vascular headache, shock.

Gastrointestinal System: constipation, diarrhea, indigestion, anorexia, nausea, vomiting, cholestatic jaundice, cirrhosis of liver, hepatic coma, hepatitis, hepatoma, hepatomegaly, jaundice, liver carcinoma (for further information see **WARNINGS AND PRECAUTIONS - Hepatic/Biliary/Pancreas**), liver failure (for further information see **WARNINGS AND PRECAUTIONS - Hepatic/Biliary/Pancreas**), abnormal liver function test, liver necrosis, pancreatitis, glossitis.

Hematology: increased fibrinogen, decreased prothrombin, thrombocytopenia, anemia (for further information see **WARNINGS AND PRECAUTIONS - Hematologic**), hemolytic anemia, hypochromic anemia, normocytic anemia, leukopenia, leukocytosis.

Metabolism: negative nitrogen balance, decreased response to ACTH, hyperglycemia, lowered cortisol, hypercalcemia, increased SGOT, increased SGPT, increased creatinine, hyponatremia, edema, weight gain, weight loss, diabetes mellitus.

Musculoskeletal System: myasthenia, osteoporosis.

Central Nervous System: fatigue, lassitude, weakness, hot flashes, increased sweating, aphasia, coma, depression, dizziness, encephalopathy, hemiplegia, personality disorder, psychotic depression, abnormal gait, headache, temporary restlessness.

Cerebral meningiomas (single and multiple) have been reported in association with long-term use (several years) of ANDROCUR (see **WARNINGS AND PRECAUTIONS - Carcinogenesis and Mutagenesis**).

Respiratory System: asthma, increased cough, dyspnea, hyperventilation, respiratory disorder, shortness of breath on effort (see **WARNINGS AND PRECAUTIONS - Respiratory**), lung fibrosis.

Skin: eczema, urticaria, erythema nodosum, exfoliative dermatitis, rash, maculopapular rash, dryness of the skin, pruritus, alopecia, hirsutism, skin discoloration, photosensitivity reactions, scleroderma.

Sensory System: ear disorder, optic atrophy, optic neuritis, abnormality of accommodation, abnormal vision, blindness, retinal disorder.

Urogenital System: enlarged uterine fibroids, uterine hemorrhage, increased urinary frequency, bladder carcinoma, kidney failure, hematuria, urate crystalluria, urine abnormality.

Other: ascites, allergic reaction, asthenia, chills, fetal chromosome abnormality, death, fever, hernia, malaise, injection site reaction.

Adverse reactions are rarely of sufficient severity to require dosage reduction or discontinuation of treatment.

If reactions are severe, it may be beneficial to reduce the dosage.

DRUG INTERACTIONS

Overview

Although clinical interaction studies have not been performed, since this drug is metabolized by CYP3A4, it is expected that ketoconazole, itraconazole, clotrimazole, ritonavir, and other strong inhibitors of CYP3A4 inhibit the metabolism of cyproterone acetate. On the other hand, inducers of CYP3A4 such as rifampicin, phenytoin, and products containing St. John's wort may reduce the levels of cyproterone acetate.

Based on in vitro inhibition studies, an inhibition of the cytochrome P450 enzymes CYP2C8, 2C9, 2C19, 3A4, and 2D6 is possible at high therapeutic cyproterone acetate doses of 300 mg daily. In addition, cyproterone acetate was also shown to increase the enzymatic activity of CYP1A2 and CYP2E1 in vitro. Caution should be exercised when ANDROCUR is to be co-administered with a substrate of the P450 enzymes.

The risk of statin-associated myopathy or rhabdomyolysis may be increased when those HMGCoA inhibitors (statins) which are primarily metabolized by CYP 3A4 are coadministered with high therapeutic cyproterone acetate doses, since they share the same metabolic pathway.

Drug-Drug Interactions

Interactions with other drugs have not been established.

Drug-Food Interactions

Interactions with food have not been established.

Drug-Herb Interactions

Interactions with herbal products have not been established.

DOSAGE AND ADMINISTRATION

Dosing Considerations

Patients with Hepatic Impairment

The use of ANDROCUR is contraindicated in patients with liver diseases and/or with hepatic dysfunction.

Patients with Renal Impairment

A pharmacokinetic study in patients with renal impairment has not been conducted. As 33% of cyproterone acetate is excreted via the kidney, caution should be taken when ANDROCUR is administered in this patient population.

Recommended Dose and Dosage Adjustment

Oral Tablets: The usual daily initial and maintenance dose of ANDROCUR (cyproterone acetate) is 4 to 6 tablets (200-300 mg) divided into 2 to 3 doses and taken with some liquid after meals.

The maximum daily dose is 300 mg.

Injectable: The usual initial and maintenance dose of ANDROCUR DEPOT is one weekly intramuscular injection of 3 mL (300 mg).

ANDROCUR or ANDROCUR DEPOT therapy should not be interrupted nor the dosage reduced after remission or improvement occurs.

Because of their pharmacokinetic properties, ANDROCUR (oral) and ANDROCUR DEPOT (IM) can be interchanged in the course of long-term treatment. The dosage may be reduced if side effects are intolerable, but should be kept within the oral range of 2 to 6 tablets daily (100-300 mg) or intramuscular injections of 300 mg at weekly intervals, or every two weeks.

Administration

ANDROCUR DEPOT is strictly for intramuscular injection. The injections must be administered very slowly. Special care must be given to avoid intravascular injection.

OVERDOSAGE

For management of a suspected drug overdose, contact your regional Poison Control Centre.

There have been no reports of fatal overdose in man with ANDROCUR (cyproterone acetate). There are no specific antidotes and treatment should be symptomatic. If oral overdose is discovered within two to three hours, gastric lavage can safely be used if indicated.

ACTION AND CLINICAL PHARMACOLOGY

Mechanism of Action

ANDROCUR (cyproterone acetate) is a steroid which clinically demonstrates two distinct properties:

- a) Antiandrogenic: Cyproterone acetate blocks the binding of dihydrotestosterone - the active metabolite of testosterone - to the specific receptors in the prostatic carcinoma cell.
- b) Progestogenic/antigonadotrophic: Cyproterone acetate exerts a negative feed-back on the hypothalamo-pituitary axis, by inhibiting the secretion of LH leading to diminished production of testicular testosterone.

Pharmacokinetics

Absorption

ANDROCUR

The absorption of cyproterone acetate following oral administration is complete. Peak plasma levels are reached 3 to 4 hours after administration. Plasma levels fall rapidly during the first 24 hours as a result of tissue distribution and excretion, and plasma half-life was 38 ± 5 hours.

ANDROCUR DEPOT

Following intramuscular administration, mean maximum blood levels are attained 3.4 days after injection. The mean elimination half-life was found to be 4 days.

Metabolism

The principal metabolite identified was 15 β -hydroxy-cyproterone acetate.

Excretion

Most of the cyproterone acetate is excreted unchanged in the feces (60%) or urine (33%) within 72 hours.

Cyproterone acetate is eliminated with the urine mainly in the form of unconjugated metabolites and with the bile (feces) in the form of glucuronidized metabolites.

DOSAGE FORMS, COMPOSITION AND PACKAGING

ANDROCUR (cyproterone acetate) 50 mg tablets are available in blisters or bottles of 60. Each white to faintly yellowish, round, flat-sided tablet with beveled edges, imprinted one side "BV" in a regular hexagon, other side scored, contains 50 mg of cyproterone acetate and the following excipients: colloidal anhydrous silica, lactose monohydrate, magnesium stearate, maize starch, and povidone 25. The tablet can be divided into equal halves.

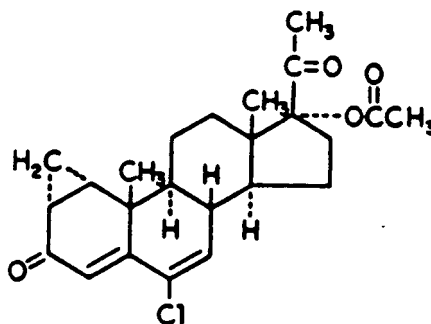
ANDROCUR DEPOT 3 mL (300 mg) ampoules. Each 3 mL ampoule contains cyproterone acetate 100 mg/mL and benzyl benzoate in a castor oil solution.

PART II: SCIENTIFIC INFORMATION

PHARMACEUTICAL INFORMATION

Drug Substance

Proper name:	Cyproterone acetate
Chemical name:	6-chloro-17 α -hydroxy-1 α , 2 α -methylene-pregna-4, 6-diene-3, 20-dione-acetate
Molecular formula:	C ₂₄ H ₂₉ ClO ₄
Molecular weight:	416.95
Structural formula:	



Physicochemical properties:	White to faintly yellow micronized powder. Insoluble in water, very freely soluble in chloroform and dioxane. Melting range is 206-213°C.
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CLINICAL TRIALS

A total of 24 studies have been conducted with ANDROCUR (cyproterone acetate) in patients requiring palliative treatment for advanced prostatic carcinoma. Worldwide, more than 1,000 patients have participated in these studies, which included several large multicentre trials in addition to the important comparative multicentre trial conducted by the European Cancer Oncology Group. North American experience has been accumulated in the U.S. by Drs. Scott (Johns Hopkins Hospital, Baltimore), Geller (Mercy Hospital & Medical Center, San Diego), and by Drs. Wein and Murphy (Hospital of the University of Pennsylvania, Philadelphia).

Study Demographics and Trial Design

Patients and Stage of Disease

As shown in [Table 2](#), more than 90% of the patients treated with ANDROCUR had stage C advanced prostatic carcinoma, or stage D1 or D2 prostatic carcinoma with metastasis.

Table 2: Patients

Stage	No. of Patients
A or B	18
C	174
C or D	502
D	349
Not specified	39
Total	1082

The majority of patients (75%) had had no therapy prior to treatment with ANDROCUR. A large group of patients had received various types of estrogen therapy, but had proven to be refractory or unable to tolerate the drug. A few patients had undergone an orchiectomy or had received radiation therapy ([Table 3](#)).

Table 3: Previous Therapy

Previous Therapy	No. of Patients
None	809
Orchiectomy	76
Estrogen	253
Radiation	16

Dosage and Administration

The oral route of administration of ANDROCUR was employed in 910 patients (84%), while 172 patients received ANDROCUR DEPOT, an oily solution containing 100 mg/mL cyproterone acetate. The standard dose of the latter was one weekly IM injection of 300 mg. As shown in the table below ([Table 4](#)), the daily oral dose varied considerably from study to study and from patient to patient. However, most patients were treated with doses ranging from 200 to 300 mg/day. In orchiectomized patients, the daily dose was generally reduced by about 50% to a range of 100 to 200 mg/day orally or the frequency of ANDROCUR DEPOT injections was reduced to one every 2 weeks.

Table 4: Dose of ANDROCUR or ANDROCUR DEPOT

Entity	Route	Dose	No. of Patients
ANDROCUR	Oral	100 mg/day	15
		200 mg/day	197
		250 mg/day	135
		300 mg/day	114
		100-300 mg/day	449
ANDROCUR DEPOT	IM	300 mg/week	172

Only 32 patients (3%) received concomitant drug therapy with ANDROCUR. No other patients received concomitant drugs, but 521 patients (48%) underwent an orchiectomy ([Table 5](#)).

Table 5: Concomitant Therapy

Concomitant Therapy	No. of Patients
None	529
Estrogen (DES 0.1 mg)	32
Orchiectomy	521

Study Results

Effect on Serum Testosterone and Prostatic Acid Phosphatase (PAP)

Table 6: Effect on Serum Testosterone and Prostatic Acid Phosphatase (PAP)

Parameter	No. of Studies	Result
Serum testosterone	7	70-90% reduction
Prostatic acid phosphatase	11	Normalization in 90% of responding patients

The effect of ANDROCUR on serum testosterone was monitored in 7 studies ([Table 6](#)). Serum testosterone was rapidly reduced following daily oral doses of 200 to 300 mg, with castrate levels being achieved within 1 to 4 weeks. The reduction was usually in the order of 70% to 90%; the greatest percent reduction occurred when ANDROCUR was combined with estrogen.

Results of PAP evaluations consistently showed a normalization of values within a very short time in responding patients. Similarly, when there were signs of progressing metastasis, PAP values again deviated from normal levels.

Effect on Primary Tumor

The effect of ANDROCUR on the primary tumor was assessed in a total of 678 patients. Of these, 489 were previously untreated; the primary tumor was reduced in 318 of these (65%) and was stabilized in another 69 (14%). Thus, the overall positive response rate in this group was 79% ([Table 7](#)).

A significant, though smaller, percentage (59%) of estrogen-refractory patients also exhibited a positive result.

Table 7: Effect on Primary Tumor

Patient Group	Number	Response of Primary Tumor		Total With Positive Effect
		Reduced	Stabilized	
Previously untreated	489	318 (65%)	69 (14%)	387 (79%)
Estrogen refractory	189	112 (59%)	-	112 (59%)

Effect on Metastasis

As shown in [Table 8](#), metastasis was reduced in 31% of 216 evaluable patients who had not previously been treated, but in only 13% of the evaluable estrogen-refractory patients. The progression of metastases appeared to be time-dependent. Despite reduced serum testosterone levels, metastases progressed over a period of several months to years, even in patients who were initially stabilized. The major cause of death during therapy with ANDROCUR was the progression of metastases and not the primary tumors.

Table 8: Effect on Metastases

Patient Group	Number	Response of Metastases		Total with Positive Effect
		Reduced	Stabilized	
Previously untreated	216	67 (31%)	82 (39%)	149 (70%)
Estrogen refractory	71	10 (13%)	7 (10%)	17 (23%)

Effect on Pain

[Table 9](#) illustrates the incidence of pain relief reported in each of 13 studies. Pain relief was noted in approximately 50 to 80% of patients receiving treatment with ANDROCUR. The effect of ANDROCUR on pain generally paralleled its effect on metastases. As long as metastases remained improved or stabilized, the analgesic requirement was also reduced. Renewed analgesic requirements were frequently indicative of metastatic progression.

Table 9: Pain Relief

Investigator	Incidence of Pain Relief
Dr. Bracci	172/216
Dr. Giuliani	12/16
Dr. Smith	12/25
Dr. Scott	8/10
Dr. Geller	8/10
Dr. Mauermayer	38/58
Dr. Wein	13/24
Dr. Tvetter	2/6
Dr. Di Silverio	13/20
Dr. Ah-Lan	9/16
Dr. Pescatore	12/16
Dr. Hermabessiere	2/4
Dr. Bruchovsky	15/24
Total	316/425 = 74%

Subjective and Objective Responses

A general improvement in the subjective assessment of the quality of life was achieved in 70% of the 367 evaluable patients ([Table 10](#)).

The objective evaluations of remissions shown in [Table 10](#) were based on ECOG criteria. The best results were obtained when ANDROCUR was used in combination with orchiectomy. One study revealed that more than 1/3 of the patients treated with ANDROCUR achieved a complete or partial

remission for 3-5 years. The Canadian study found that a complete or partial remission was still evident in 75% of the patients after one year of treatment.

Table 10: Subjective and Objective Responses

Subjective Responses			
No. Evaluable Patients		No. Improved ^a	
367		255 (70%)	
Objective Responses (ECOG Criteria)			
Treatment	Patient Group	No. of Patients	No. With Complete or Partial Remissions
ANDROCUR	Previously untreated	270	134 (50%)
ANDROCUR	Estrogen-refractory	77	31 (44%)
ANDROCUR/ Orchiectomy	Previously untreated and/or estrogen-refractory	274	154 (60%)

a Based on criteria of general improvement in quality of life (ie, weight gain, pain relief, etc.)

Survival Rate

Table 11: Survival Rate

Investigator	No. of Patients	Stage	Duration of Treatment	Survival	
				ANDROCUR	Estrogen
Dr. Mauermayer	58	C or D	2 - 5 years	38/58 (70%)	-
Dr. Wein	55	A (7)	4 years	39/55 (70%)	-
		C (25)			
		D (23)			
Dr. Bracci	216	C or D	5 years	138/216 (64%)	-
Dr. Di Silverio	20	D	up to 38 months	3/20 (15%)	-
Dr. Giuliani	68	C	5 years	30/68 (44%)	31%
Dr. Giuliani	38	D	3 years	10/38 (27%)	10%
Dr. Jacobi	51	C or D	2 years	18/40 (45%)	-
Dr. Pavone	103	C or D	3.5 - 5 years	42/103 (41%)	41%
Dr. Bruchovsky	29	D	9 - 15 months	23/29 (80%)	-

As shown in [Table 11](#) above, 5-year survival rates ranged from 41% to 64%. The 3-year rate for stage D patients was 27% and 1- to 2-year rates varied from a low of 15% up to a high of 80%. These survival rates generally represented an improvement over results previously obtained with estrogen therapy.

DETAILED PHARMACOLOGY

Animal Pharmacology

Antiandrogenic Effects

Cyproterone acetate at doses of 10 or 50 mg/kg inhibits the effects of endogenously produced and exogenously administered androgens at the prostate by means of competitive inhibition.

In mice and dogs, cyproterone acetate induces a dose-dependent atrophy of the accessory sex glands, the prostate, seminal vesicles, and preputial glands.

Spermatogenesis is inhibited in a dose-related manner; however, the atrophy in the Leydig cells are slight.

In the rat the start of puberty is prevented or delayed. Cyproterone acetate inhibits the physiological closure of the epiphyseal cartilages and bone maturation.

It impairs the function of the sebaceous glands, and the thickness of the epidermis decreases.

The treatment of pregnant animals with cyproterone acetate leads to developmental disturbances in male fetuses. Testosterone-dependent differentiation processes are affected: signs of feminization of varying degrees of severity develop.

Progestogenic and Antigonadotrophic Effect

On subcutaneous injections a total dose of 0.003 mg cyproterone acetate is about 100 times stronger than progesterone in the maintenance of pregnancy (Clauberg test). Like all potent progestogens, cyproterone acetate has antigonadotrophic properties which can be demonstrated in the parabiosis test, the testicular inhibition test in infantile rats, and by the inhibition of ovulation.

Pharmacokinetic Studies in Animals

Pharmacokinetic studies have been carried out in a number of animal species (rats, rabbits, dogs, and monkeys) using either methylene-¹⁴C- or carboxy-¹⁴C-labelled cyproterone acetate.

Cyproterone acetate is absorbed at most dose levels tested except in high doses. Peak plasma levels are usually obtained within 1 to 4 hours of oral dosing. Because of its lipophilic character, cyproterone acetate is taken up and concentrated in the liver and fatty tissues in all animal species. Cyproterone acetate is not hydrolysed, and mainly cyproterone acetate and the metabolite 15 β -hydroxycyproterone acetate are found in the tissues and in plasma. The elimination half-life of cyproterone acetate is slow in most species (1-2 days), in a ratio of 4:6 with urine and feces; an exception is the dog, which excretes cyproterone acetate in 1 to 3 days. On repeated daily dosing, cyproterone acetate shows limited rise, and plasma levels can be taken as a reliable index of the concentrations of cyproterone acetate in the body. Cyproterone acetate passes the placental barrier, but only reaches the fetus in low concentrations. The pharmacokinetics, biotransformation, and metabolic spectra of cyproterone acetate are similar in man and the rhesus monkey.

Human Pharmacology

Antiandrogenic Effect

The following actions which are associated with the antiandrogenic effects have been described in man: reduction of sexual drive; inhibition of spermatogenesis; palliative effect in prostatic carcinoma; inhibition of sebaceous gland activity; suppression of signs of androgenization in women; inhibition of premature genital development in children; and other associated symptoms.

Progestogenic and Antigonadotrophic Effect

Cyproterone acetate in man is also a potent progestogen and has an antigonadotrophic effect. It intervenes with the hypothalamo-pituitary pathway, causing an inhibition of increased secretion of LH, and a decrease in gonadal testicular androgens.

Thus, unlike pure antiandrogens, cyproterone acetate does not cause a compensatory increase in androgen secretion.

Other Endocrine Effects

No distinct influence on the 17-ketosteroids, 17-ketogenic steroids or on total estrogens in the 24-hour urine has been observed in male patients. On fluorometric determination of urinary cortisol, the value apparently increases because the cyproterone acetate eliminated with the urine is also measured. Simultaneously, cyproterone acetate also reduces the reaction of the adrenal cortex to exogenous ACTH in patients; the baseline cortisol and ACTH values may also be reduced.

Pharmacokinetics

A bioavailability study was performed in 5 male volunteer subjects receiving a single oral dose of 50 mg ¹⁴C-cyproterone acetate tablets.

Results of the study showed that cyproterone acetate is absorbed slowly, but completely (100%), from the gastrointestinal tract. The maximum plasma level was reached 3 to 4 hours after ingestion. The mean plasma levels were 700 nmol/L (=290µg/L) cyproterone acetate or, including the radioactivity of metabolites, 960 nmol/L (=400µg/L) cyproterone acetate equivalent.

The plasma levels fell quickly up to 24 hours after administration because of extensive tissue distribution. The half-life of cyproterone acetate in plasma was calculated as 38 ± 5 hours (see [Figure 1](#)).

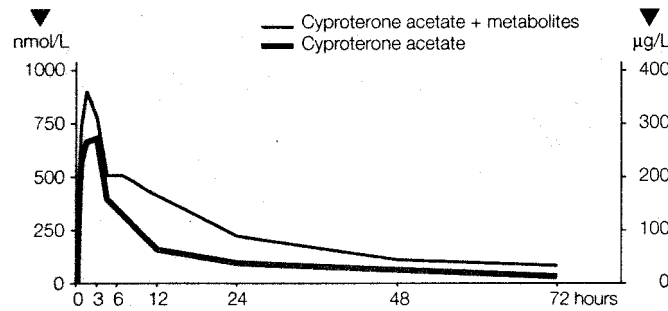


Figure 1: Relationship of Unchanged Cyproterone Acetate to the Total ¹⁴C-labelled Substance (Cyproterone Acetate + Metabolites) in the Plasma of a Male Subject Following Oral Administration of 50 mg ¹⁴C-cyproterone Acetate

On oral administration cyproterone acetate was eliminated with a half-life of 38 ± 2 hours. After 10 days, $33 \pm 6\%$ of the dose could be recovered in the urine and $60 \pm 8\%$ in the feces (see [Figure 2](#)).

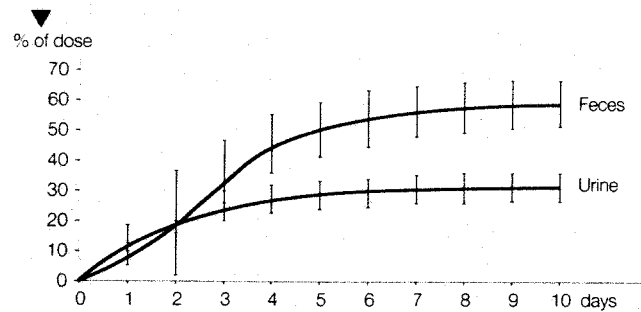


Figure 2: Elimination (% of Dose) Following Oral Administration of 50 mg ¹⁴C-cyproterone Acetate in Male Subjects. Mean Values \pm Standard Deviation (n=5)

The intramuscular injection of 300 mg radioactivity labelled cyproterone acetate in a castor oil solution corresponding to ANDROCUR DEPOT was administered to male patients.

The maximum plasma level was reached 82 ± 21 hours after administration. Half the maximum value could be seen less than 24 hours after administration; the values did not fall below this level until about 10 days after dosing (see [Figure 3](#)).

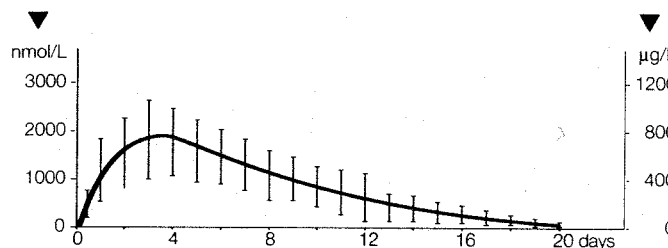


Figure 3: Plasma Levels (Cyproterone Acetate Equivalents/mL) Following IM injection of 300 mg Cyproterone Acetate in Oily Solution in Male Patients. Mean Values \pm Standard Deviation (n=11)

The elimination half-life of the cyproterone acetate released from the depot was 38 ± 14 hours which is the same as measured under oral administration.

A steady-state study was also carried out in 5 patients who received 300 mg of ANDROCUR DEPOT on a weekly basis. Determinations of the cyproterone acetate concentrations were carried out after the first, third, and fifth injections. Table 12 summarizes the results of the study.

In man, cyproterone acetate is eliminated in the urine mainly in the form of unconjugated metabolites and in the bile in the form of glucuronized metabolites; the main metabolite was 15β -OH cyproterone acetate.

Table 12: Pharmacokinetic Parameters After One and Several Intramuscular Injections of 300 mg Cyproterone Acetate in an Oil Solution (ANDROCUR DEPOT) in 5 Patients (Mean Values \pm SD).

Parameter	1st Injection	3rd Injection	5th Injection
t_{\max} (d)	1.8 ± 0.4	2.4 ± 0.5	3.0 ± 1.0
C_{\max} (ng/mL)	273 ± 54	387 ± 111	406 ± 57
$t_{1/2}$ (d)	4.4 ± 1.9	4.1 ± 1.3	3.9 ± 1.3

TOXICOLOGY

ANDROCUR (cyproterone acetate) has been found at low doses of 2 to 10 mg/kg to cause liver abnormalities in dogs and rats in the form of proliferative liver changes including increased liver weight, liver cell hypertrophy with an increase in the smooth endoplasmic reticulum, and a rise in the serum glutamic pyruvic transaminase (SGPT). At high doses of 50 to 100 mg/kg, nodular hepatic hyperplasia and hepatomas have also been observed.

Recognized first-line tests of genotoxicity gave negative results when conducted with CPA. However, further tests showed that CPA was capable of producing adducts with DNA (and an increase in DNA repair activity) in liver cells from rats and monkeys and also in freshly isolated human hepatocytes.

This DNA-adduct formation occurred at exposures that might be expected to occur in the recommended dose regimens for CPA. One in vivo consequence of CPA treatment was the increased incidence of focal, possible preneoplastic, liver lesions in which cellular enzymes were altered in female rats.

The clinical relevance of these findings is presently uncertain. Clinical experience to date would not support an increased incidence of hepatic tumors in man.

Acute Toxicity

The LD_{50} after single application of cyproterone acetate was as follows:

Table 13: LD₅₀ After Single Application of Cyroterone Acetate

Animal Species	Oral (mg/kg)	Subcutaneous (mg/kg)	Intraperitoneal (mg/kg)	Intramuscular (mg/kg)
Mouse	>6000	>5000	>4000	-
Rat	>4000	1500	1000	-
Dog	>3000	-	-	>100 (approx.)

On the basis of the above LD₅₀ values, cyproterone acetate can be considered practically nontoxic following single dose administration. The maximum intramuscular doses were also tolerated without symptoms in the dog, with exception of local tolerance manifestation.

Repeated Dose Toxicity

Chronic Toxicity Studies

Table 14: Chronic Toxicity Studies

Animal Species	Dosage and Duration	Mortality and Clinical and Laboratory Observations	Necropsy and Histopathology
Rats 35/sex/dose	0; 10; 50 and 250 mg/kg 78 weeks oral	250 mg/kg: marked increase in mortality rate. 50 and 250 mg/kg: 40-50% decrease in body weight gain. SGPT increase: males 10 and 250 mg/kg; females 50 mg/kg. BUN increase: males 50 and 250 mg/kg. Cholesterol increase: all treated groups.	Dose-related increase in liver weights. Increase thyroid weight except for low dose males. Dose-related decrease in gonads, adrenal, prostate, seminal vesicle, and uterus weights. Histopathology: toxic manifestation in liver and kidneys - less at 10 mg/kg, more extensive at 50 and 250 mg/kg. Changes included: yellow nodules and mottling of liver (including liver cell hyperplasia and liver cell adenomas and endoplasmic inclusion bodies), discolored kidneys with rough surfaces.
Rats 60/sex/dose	0; 0.04; 0.4 and 2 mg/kg 104 weeks oral	No drug-related mortality. Dose-related decrease in body weight gains in males and increase in females. Food consumption reduced and thinning and loss of hair was also noted for high-dose males. Decrease in hemoglobin and erythrocytes at 0.4 and 2 mg/kg. SGOT, SGPT and alkaline phosphatase increased at 2 mg/kg.	2 mg/kg increased incidence of subcutaneous masses and/or nodules; liver discoloration and nodules; atrophy of testes, seminal vesicles, and prostate. Increased incidence of mammary neoplasms (adenomas and adenocarcinomas).
Mice 50/sex/dose	0; 0.04; 0.4 and 2 mg/kg 105 weeks oral	No dose-related mortality. Thinning and loss of hair at 2 mg/kg. Slightly reduced body weight gain at 2 mg/kg.	Slightly increased incidence of skin masses and/or nodules and alopecia. No drug-related inflammatory, degenerative, proliferative and/or neoplastic lesions.

Table 14: Chronic Toxicity Studies

Animal Species	Dosage and Duration	Mortality and Clinical and Laboratory Observations	Necropsy and Histopathology
Dogs Beagle 4/sex/dose	0; 10; 32 and 100 mg/kg 55 weeks oral	No mortality. Excessive lacrimation, retarded pupillary reflex, mild conjunctivitis, hyperemia of gums, abdominal distention, sparsity of hair, and quieted behaviour. Laboratory tests: slightly elevated alkaline phosphatase and SGPT at 100 mg/kg in 2 dogs. Elevated sedimentation rate, slightly reduced lymphocytes with increase in segmented neutrophils and decrease in eosinophils.	Reduced adrenal, testes, and prostate weight for all cyproterone acetate-treated animals. Ovary and uterus weights reduced at 100 mg/kg. Liver weight slightly increased for some dogs. Histopathology: marked adrenal atrophy of zona fasciculata and reticularis, testicular atrophy and absence of spermatogenesis, some Leydig cell hyperplasia, prostatic atrophy, ovarian and uterine atrophy, hyperplasia of mammary gland in males and females.
Rhesus monkey 4 females/ dose	0; 0.04; 0.4 and 40 mg/kg 12 weeks oral	No mortality or behaviour changes. Dose-related alopecia. Raised insulin level above 0.04 mg/kg. Negative influence on coagulation at 0.4 mg/kg and 40 mg/kg. Stimulation of ACTH cells at 0.4 mg and above. Increase in prolactin cells and slight reduction in gonadotrophin cells. Galactorrhea in all treated.	At doses of 0.4 mg/kg and above - diffuse liver cell hypertrophy and an increase in smooth endoplasmic reticulum. At the two highest doses, 2 and 3 animals also had occasional eosinophil cytoplasmic inclusion bodies in the liver cells. In most treated animals small mammary nodules were palpable in the glandular tissue; at 40 mg/kg slight ductus proliferation was also noted.

Reproductive Toxicology

Table 15: Fertility and Reproduction Study

Animal Species	Route and Dosage of Administration	Findings
Rats 24/sex/dose (2 generations)	0; 0.4; 4.0 and 40 mg/kg oral	0.4 mg/kg: No influence by drug on fertility of the P1 and F1 generations. 4 mg/kg: Significant decrease in body weights but no impairment of pre- and postnatal development. 40 mg/kg: Food intake and body weight gain reduced. Although attempted matings were increased, less than 50% of the females had litters. No specific pathological changes were found in the dams, fetuses, or young. Similarly, no malformations were observed.

Mutagenesis

No mutagenic effect of cyproterone acetate was demonstrated in either in vitro (Salmonella typhimurium) or in vivo (micronucleus test in the monkey.)

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PART III: CONSUMER INFORMATION

ANDROCUR®

cyproterone acetate tablets

ANDROCUR® DEPOT

cyproterone acetate injection

This leaflet is Part 3 of a three-part "Product Monograph" published when ANDROCUR was approved for sale in Canada and is designed specifically for Consumers. This leaflet is a summary and will not tell you everything about ANDROCUR. Contact your doctor or pharmacist if you have any questions about the drug.

ABOUT THIS MEDICATION

What the medication is used for:

ANDROCUR is indicated for the palliative treatment of patients with advanced prostate cancer.

What it does:

ANDROCUR is a drug acting against male sex hormones (androgens). Androgens encourage the growth of prostate cancer and ANDROCUR inhibits this effect.

When it should not be used:

Do not take ANDROCUR if you:

- are allergic (hypersensitive) to cyproterone acetate or any of the other ingredients of ANDROCUR;
- have a liver disease or reduced liver function;
- have Dubin-Johnson syndrome or Rotor syndrome that affects the excretion of red blood cell pigment (bilirubin);
- have or have had liver tumors that are not due to the spread of prostate cancer;
- have reduced kidney function;
- have wasting diseases that are not related to prostate cancer;
- suffer from severe chronic depression;
- have conditions that increase your risk for developing blood clots (thromboembolic process).

What the medicinal ingredient is:

Cyproterone acetate

What the nonmedicinal ingredients are:

The other tablet ingredients are colloidal anhydrous silica, corn starch, lactose monohydrate, magnesium stearate, and povidone 25

The other solution ingredients are castor oil for injection, benzyl benzoate.

What dosage forms it comes in:

ANDROCUR is supplied in 2 forms:

Tablet: Each tablet contains 50 mg cyproterone acetate

Solution: Each 3 mL ampoule contains 100 mg/mL cyproterone acetate in oily solution

WARNINGS AND PRECAUTIONS

ANDROCUR may decrease the function of the hormone-producing glands attached to the kidney (adrenal glands). Therefore your doctor may conduct some tests to monitor this effect during treatment with ANDROCUR.

BEFORE you use ANDROCUR talk to your doctor or pharmacist if you have or have had any of the following conditions:

- Blood clots;
- Liver problem;
- Depression;
- Breathing problem (shortness of breath).
- Diabetes. Your doctor may need to adjust your antidiabetic medication. This requires strict supervision during treatment with ANDROCUR.
- Anemia. Your doctor will monitor your red-blood cell count during treatment.

ANDROCUR is not recommended for use in children and adolescents below 18 years of age and before the end of puberty, as it may have a negative effect on growth and the immature hormonal regulation functions.

ANDROCUR is not recommended for use in women.

INTERACTIONS WITH THIS MEDICATION

Drugs that may interact with ANDROCUR include:

- Statins (medicines for reducing blood fats)
- Ketoconazole, itraconazole, clotrimazole (for fungal infections)
- Ritonavir (for viral infections)
- Rifampicin (for tuberculosis)
- Phenytoin (for epilepsy)
- St. John's Wort (herbal remedy for depression)

PROPER USE OF THIS MEDICATION

You should follow the dose prescribed by your doctor.

Usual dose**ANDROCUR Tablet**

200 mg to 300 mg daily taken orally in two or three divided doses with liquid after meals. Maximum daily dose is 300 mg.

ANDROCUR DEPOT Injection

300 mg given slowly into a muscle (intramuscular) once a week.

Overdose

In case of drug overdose, contact a health care practitioner, hospital emergency department or regional Poison Control Centre immediately, even if there are no symptoms.

Missed Dose**ANDROCUR Tablet**

Do not take the missed ANDROCUR tablet (do not take a double dose to make up for the missed tablet). Continue taking the tablets at the regular time.

ANDROCUR DEPOT Injection

Have a new appointment with your doctor in order to make up for the forgotten injection. You should not get a double dose.

SIDE EFFECTS AND WHAT TO DO ABOUT THEM

ANDROCUR can have side effects, like all medicines, but not everybody gets them. For further information about any of these effects, ask a doctor or pharmacist.

If any of the side effects get serious, or if you notice any side effects not listed here, please tell your doctor or pharmacist.

Very frequent side effects:

- Reduced sexual interest (decreased libido)
- Swelling of the breast, breast soreness (gynecomastia)
- Impotence
- Abnormal sperm, low sperm count which may be reversible after ANDROCUR is discontinued.

Other side effects:

- Hair loss or unusual increase in hair growth
- Constipation or diarrhea (loose stools)
- Weight gain
- Tiredness and weakness
- Shortness of breath
- Skin rash, blisters
- Vision change
- Decrease in blood pressure
- Increase in blood sugar
- Depression

Rare but serious side effects:

- Liver toxicity: generally feeling unwell, fever, nausea, vomiting, loss of appetite, itching, yellowing of the skin and eyes, light-colored stools, dark-colored urine
- Life-threatening internal bleeding (intra-abdominal hemorrhage): unusual upper abdominal pains which do not disappear within a short time
- Blood clots: swelling of the calf or leg (blood clots in the leg), chest pain and being short of breath (blood clots in the lung), suddenly feeling weak, loss of coordination, slurred speech (a stroke or blood clots in the brain), temporary blindness (blood clots in the eye)

Bone loss (osteoporosis) and benign brain tumors (cerebral meningioma) have been reported with long term use of ANDROCUR.

SERIOUS SIDE EFFECTS, AND WHAT TO DO ABOUT THEM

Symptom/ Effect	Talk with your doctor or pharmacist		Stop taking drug and call your doctor or pharmacist
	Only if severe	In all cases	
Inability to achieve or maintain an erection	✓		
Liver toxicity, (generally feeling unwell, fever, nausea, vomiting, loss of appetite, itching all over the body, yellowing of the skin or eyes, light colored bowel movements, dark urine) liver inflammation (hepatitis), liver failure			✓
Reduced sexual drive	✓		
Reversible inhibition of sperm production	✓		
Swelling of the calf or leg, chest pain, suddenly feeling weak			✓
Unusual upper abdominal pains which do not disappear within a short time			✓

This is not a complete list of side effects. For any unexpected effects while taking ANDROCUR, contact your doctor or pharmacist.

HOW TO STORE IT

Keep out of the reach and sight of children.

Do not take ANDROCUR after the expiry date which is stated on the pack.

Medicines must not be disposed of via wastewater or household waste. Ask your pharmacist how to dispose of medicine no longer required. These measures will help to protect the environment.

REPORTING SUSPECTED SIDE EFFECTS

Canada Vigilance Program

You can report any suspected adverse reactions associated with the use of health products to the Canada Vigilance Program by 1 of the following 3 ways:

- Report online at www.healthcanada.gc.ca/medeffect
- Call toll-free at 1-866-234-2345
- Complete a Canada Vigilance Report Form and:
 - Fax toll free to 1-866-678-6789
 - Mail to: Canada Vigilance Program
Health Canada
Postal Locator 0701D
Ottawa, ON K1A 0K9

Postage paid labels, Canada Vigilance Report Form and the adverse reaction reporting guidelines are available on the MedEffect™ Canada website at www.Healthcanada.gc.ca/medeffect.

NOTE: Should you require information related to the management of the side effect, please contact your health professional. The Canada Vigilance Program does not provide medical advice.

Bayer Inc.

You can report any suspected adverse reactions associated with the use of health products to Bayer Inc. by:

- Toll-free telephone: 1-800-265-7382
- Email: canada.medinfo@bayer.com
- Regular mail: Bayer Inc.
77 Belfield Road
Toronto, Ontario
M9W 1G6
Canada

NOTE: Should you require information related to the management of the side effect, please contact your health professional. Bayer Inc. does not provide medical advice.

MORE INFORMATION

For more information, please contact your health professional or pharmacist first, or Bayer Medical Information at 1-800-265-7382 or canada.medinfo@bayer.com.

This document plus the full product monograph, prepared for health professionals can be found at <http://www.bayer.ca> or by contacting the sponsor at the above mentioned phone number and email address.

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