Six-month gonadotropin releasing hormone (GnRH) agonist depots provide efficacy, safety, convenience, and comfort

E David Crawford

Jason M Phillips

University of Colorado Health
Sciences Center, Aurora, CO, USA

Abstract: Two different 6-month GnRH agonist depot formulations approved for palliative treatment of advanced and metastatic prostate cancer in the United States – leuprolide acetate 45 mg and triptorelin pamoate 22.5 mg – provide patients with efficacy and safety comparable to those of existing 1-, 3-, and 4-month GnRH agonist depots. However, the 6-month formulations can increase patient convenience, comfort, and compliance by reducing the number of physician visits and injections required. At the conclusion of their pivotal trials, the 6-month formulations demonstrated efficacy rates in achieving chemical castration (serum testosterone ≤ 50 ng/dL) that ranged between 93% and 99%. As with existing GnRH agonist depot formulations, hot flashes represented the most common adverse event reported in trials of 6-month leuprolide acetate or triptorelin. As such, these products may prove useful not only for their labeled indication, but also as adjuncts to other treatments such as radical prostatectomy, radiotherapy, and chemotherapy. We recommend further research, including head-to-head trials between the 6-month GnRH depots, to refine our understanding of these products.

Keywords: prostate cancer, leuprolide, leuprolide, triptorelin, 6-month depot, testosterone

Evolution of anti-androgenic therapies

In the United States and Europe, prostate cancer represents the most commonly occurring nonskin cancer in men, among whom it causes more deaths than any cancer other than lung cancer.¹ For men with advanced prostate cancer, testosterone suppression – most often achieved by the administration of a gonadotropin hormone-releasing hormone (GnRH) analog – remains the standard palliative treatment.² In fact, approximately 90% of prostate cancer tumors respond to initial androgen deprivation, thereby improving patients' quality of life and longevity.³

However, early GnRH agonists required patients to perform daily subcutaneous or intramuscular injections,⁴ which could cause pain, injection site reactions, and compliance challenges. Over the past decade, advances in drug delivery systems have given rise to 1-, 3-, 4-, 6-, and 12-month delivery systems for GnRH agonists.² Along with facilitating physician use, these longer-term formulations have greatly improved patient compliance with therapy.⁵

The recent approval of triptorelin pamoate 22.5 mg (Trelstar®, Watson Pharmaceuticals, Inc, Corona, CA) gives US physicians and patients a second 6-month depot formulation to consider, along with leuprolide acetate (LA) 45 mg (Eligard®, Sanofi-aventis, Paris, France), approved in 2004. Based on large-scale Phase III trials, these products appear to offer safety and efficacy equivalent to those of shorter-term depot formulations.^{2,6} However, 6-month depots allow patients who travel, or who have

Correspondence: E David Crawford
University of Colorado Health Sciences
Center, Mail Stop #F710, PO Box 6510,
Aurora, CO 80045, USA
Tel +1 7208480183
Email david.crawford@uchsc.edu

other difficulties being evaluated and treated every 3 months, to double the treatment interval.

The concept of androgen deprivation originated in studies published in 1941 in which Huggins and Hodges first noted the relationship between orchiectomy and prostate cancer. Specifically, they showed how androgen blockade caused dramatic and significant clinical remissions of metastatic prostate cancer.⁷

Today, oncologists utilize androgen suppression as monotherapy for selective patients with localized prostate cancer, and in conjunction with radiotherapy for patients with locally advanced disease or intermediate to high-risk localized disease.⁸ Suppressing circulating testosterone levels is also the most widely used palliative treatment for patients with metastatic disease.⁹

Researchers first isolated and described gonadotropin releasing hormone (GnRH) in 1971, leading to the discovery of the first clinically used synthetic GnRH analog, leuprolide acetate (also called leuprorelin), in 1973.^{10,11} Leuprolide acetate's chemical structure differs from that of GnRH in that it substitutes D-leucine for glycine.¹² This difference gives leuprorelin enhanced binding affinity for the anterior pituitary receptor and increased resistance to degradation by peptidases, resulting in a longer half-life than naturally occurring GnRH and 80 times more potency.¹³

The anterior pituitary gland secretes the gonadotropins luteinizing hormone (LH) and follicle-stimulating hormone (FSH) in response to stimulation by GnRH.^{14,15} In males, FSH supports spermatogenesis, while LH stimulates testosterone production in the testes. Prostate cancer cell growth depends on the presence of testosterone.¹⁴ As such, GnRH agonist therapy causes an initial, transient increase in testosterone levels, but long-term use of such drugs suppresses testosterone production.^{14,16,17}

Triptorelin was first synthesized in the early 1970s.^{18,19} Compared with naturally occurring GnRH, triptorelin substitutes D-tryptophan for glycine at position 6, which provides biological potency superior to the native decapeptide while increasing resistance to proteolytic enzymes.¹⁴ Radioligand binding and inositol phosphate production assays have shown that triptorelin is 131 times more potent than the natural GnRH.¹⁵ Additionally, triptorelin appears to degrade more slowly than natural GnRH.²⁰ Like leuprorelin, use of triptorelin initially stimulates the pituitary gland, causing a temporary increase in testosterone production. Continuous administration of triptorelin downregulates the pituitary GnRH receptors, however, thereby inhibiting secretion of gonadotropins.²⁰

All GnRH agonists possess a short elimination half-life.¹⁸ Specifically, in a pharmacokinetic (PK) study in which investigators gave healthy men intravenous 0.5 mg triptorelin acetate, the drug had a geometric mean terminal elimination half-life of 2.8 hours.²¹ In a similar PK study in which participants received a 1 mg nondepot injection of leuprolide acetate either intravenously or subcutaneously, mean elimination half-life was 2.9 hours and 3.6 hours, respectively.²²

For patients receiving the 6-month triptorelin pamoate formulation, those with renal or hepatic impairment, require no dosage adjustments.²¹ However, no studies to date have evaluated the pharmacokinetics of leuprolide acetate in patient populations with compromised kidney or liver function.⁸

GnRH agonists' short elimination half-life initially meant that patients had to inject such medications daily. However, incorporating triptorelin into a biocompatible, biodegradable copolymer (poly-D,L-lactide-co-glycolide) as microgranules has enabled the creation of 1-, 3-, and 6-month depot formulations of the drug.⁶ Similar advances have facilitated development of long-term leuprolide acetate depot formulations.^{23,24} The first such product, a monthly 7.5 mg leuprolide acetate depot formulation, earned US Food and Drug Administration (FDA) approval for treatment of advanced prostate cancer in 1989.²⁵ Three- and 4-month formulations earned approval in 2002 and 2003, respectively, for the same indication. In 2004, the FDA approved a 6-month 45 mg depot leuprolide acetate formulation.

Additionally, a 12-month leuprolide acetate implant earned FDA approval for locally advanced or metastatic prostate cancer in 2000 but was discontinued by its manufacturer in 2007.^{26,27} Some reviewers postulate that this implant failed commercially because it required a minor surgical procedure and rigorous follow-up with patients.⁸

Comparing and contrasting 6-month formulations

Pivotal trials of 6-month triptorelin pamoate and leuprolide acetate formulations approved in the United States share many design similarities, although subtle differences exist as well. Both trials were 1-year, multicenter, open-label investigations in which investigators evaluated the safety and efficacy of two injections spaced 6 months apart and found high efficacy rates and low rates of serious adverse events (SAEs). Table 1 describes the study design of the two trials.

Table I Study designs of 6-month GnRH depot Phase III trials^{2,6}

Study	Drug	Patients	Stage	Injections
A 12-month clinical study of LA-2585 (45.0 mg): a new 6-month subcutaneous delivery system for leuprolide acetate for the treatment of prostate cancer	Leuprolide 45 mg	n = 111	Stage I to 4NxMx	Day 1, Week 24
Triptorelin 6-month formulation in the management of patients with locally advanced and metastatic prostate cancer: an open-label, noncomparative, multicenter, Phase III study	Triptorelin 22.5 mg	n = 120	Stage T3 to 4NxMx, TxNxMx, or TxNxM1	Day 1, Week 24

Leuprolide acetate

The leuprolide acetate trial that led to US approval sought as its primary endpoint a decrease in total serum testosterone to ≤ 50 ng/dL on at least two consecutive measurements taken 1 week apart.² For this study, researchers enrolled 111 patients with a histological or cytological diagnosis of prostate cancer greater than stage T1, a World Health Organization (WHO) performance score of 0–2, and a life expectancy ≥ 1 year. Mean patient age was 78.2 ± 7.5 years, with nearly half of patients between 70 and 79. Approximately 40% of patients had T2 or M+, 17% had T3, and 4% had stage T1. Investigators administered subcutaneously 45 mg LA at baseline and on day 168 to all patients. Investigators tested patients' blood samples for testosterone, LH, prostate-specific antigen (PSA), and total acid phosphatase at initial screening, at baseline, and at regular intervals throughout the study. They also took additional blood samples from 27 patients for PK analysis of serum LA.

Of the original 111 patients, 103 patients completed the study, of which 73 had no missing data points. All patients received the first study injection, and 106 received the second. Within 8 hours of the first injection, mean LH increased from 6.9 ± 0.3 mIU/mL to 37.9 ± 2.4 mIU/mL. By day 7, however, mean LH had decreased below baseline. It continued to drop through the first 19 weeks of treatment, to a level of 0.1 ± 0.01 mIU/mL at day 133. Mean LH remained at this level until the second injection occurred, on day 168. On day 169, mean LH rose to 0.2 ± 0.02 mIU/mL, holding relatively constant through the duration of the study. At month 12, mean LH measured 0.2 ± 0.14 mIU/mL.²

Baseline testosterone averaged 351.4 ± 28.6 , 352.8 ± 8.0 , 367.5 ± 26.0 , and 385.1 ± 24.0 ng/dL for patients with T1, T2, C, or M+ disease stage, respectively. Mirroring the initial LH increase, mean testosterone rose to 588.6 ± 23.9 ng/dL by day 2. By day 28, 108 of the 111 patients (97%) had reached castrate testosterone levels commonly defined as testosterone < 50 ng/dL.¹¹ However, some authors advise

using plasma testosterone levels that approximate the level achieved through surgical castration, 20 ng/mL.^{28,29} At day 28, 92 patients had testosterone levels < 20 ng/dL. Using testosterone < 50 ng/dL as the definition of castration, patients took an average of 21.2 days to reach testosterone suppression. No patients reported clinically relevant flare reactions to the initial testosterone increase.

Baseline PSA measured ≥ 4 ng/mL in 83/110 patients (75.5%) and averaged 39.8 ± 21.5 ng/mL. Throughout the study, mean PSA declined, to the point that at month 12, only 4/103 patients then participating had increased PSA. Patients reported no changes in bone pain, urinary symptoms, and urinary pain at any point during the study.

Triptorelin

Investigators for the US pivotal trial of a 6-month triptorelin pamoate depot enrolled 120 patients with histologically proven prostate cancer stage T3 to 4NxMx, TxNxMx, or TxNxM1 (where T = tumor, N = node, and M = metastasis) or rising serum PSA after failed local therapy. Patients were also required to have serum testosterone ≥ 5 nmol/L (144 ng/dL) and life expectancy > 18 months.⁶ Researchers calculated the sample size required to demonstrate achievement and maintenance of castrate testosterone levels in 95% of patients with a 2-sided 95% confidence lower limit of 88.7%.

All patients underwent intramuscular triptorelin injections on day 1 and at week 24. Investigators drew blood samples for testosterone assessments at baseline, then monthly thereafter. To assess serum PSA, investigators drew samples on day 1 and at months 3, 6, 9, and 12. For safety assessments, investigators took blood samples at baseline, month 6, and month 12.

Ultimately, 115 patients completed the study. Mean patient age was 71.1 ± 8.5 years. More than 60% of patients were Caucasian. More than half (51.6%) of patients had T3N0Mx or T3NxMx cancer. The latter category accounted for 45.8% of the total patient population. Conversely, a total of 22.5% had either T4NxM1 (5.0%) or T4NxMx (17.5%).

Baseline serum PSA averaged 19.1 µg/L and testosterone levels averaged 17.8 nmol/L.

Study investigators also reported concomitant baseline diagnoses occurring in at least 10% of patients. In this regard, 61.7% of the study population had hypertension; 14.2% had hypercholesterolemia; and 10.0% had diabetes mellitus.

Regarding efficacy, 97.5% (95% confidence interval [CI]: 92.9%–99.5%) of the intent-to-treat population (120 patients) achieved castrate serum testosterone levels (≤ 50 ng/dL) by day 29.⁶ Additionally, 93.0% (95% CI: 86.8%–97.0%) or 107 of the 115 who completed the study maintained castration from months 2 through 12. Also at month 12, 98.3% (113/115) patients had castrate serum testosterone levels. In keeping with these results, median relative decreases in serum PSA from baseline were 97% and 96% at months 6 and 12, respectively.

Discussion

Both of these US trials investigated the efficacy and safety of 6-month GnRH agonists in more than 100 patients with mean ages between 71.1 (triptorelin trial) and 73.6 years (US leuprolide acetate trial) with life expectancy of ≥ 1 year. Inclusion criteria differed between the two studies. For example, the triptorelin trial included patients with PSA relapses after failed local therapies. Specifically, 28.3% in the triptorelin trial had PSA relapse after either radical prostatectomy or radiotherapy. In contrast, the US leuprolide acetate trial included no PSA failures after local therapy.

As for main inclusion criteria, the US leuprolide acetate trial required patients to have a histological or cytological diagnosis of prostate cancer greater than stage 1 and a WHO performance score of 0 to 2. However, the triptorelin trial included patients with locally advanced or metastatic prostate cancer but did not specify any WHO status.

Efficacy

In assessing the drugs' clinical performance, both studies reported on the proportion of patients who reached serum testosterone levels ≤ 50 ng/dL at various time intervals. Although each trial presented its results somewhat differently, a consistent picture emerges, with overall efficacy rates well above 90%. Table 2 summarizes the results of the castration efficacy of the two studies.

Specifically, after 1 month of treatment, 99% (108/109) of patients participating in the US leuprolide acetate trial achieved castration levels as well as 97% of the trial's 111-patient intent-to-treat population. Similarly, 97.5% of patients

Table 2 Comparison of results leuprolide acetate with triptorelin pamoate^{2,6}

	Leuprolide	Triptorelin
Castration efficacy*		
Day 29**	97.0%	97.5%
Month 12	99.0%	98.3%
Castration time	21.2 days	18.8 days
PSA decreases		
Month 12	90.2%	96.0%
Adverse events		
Hot flashes	57.6%	71.7%

Notes: *Castration < 50 ng/dL; **Intent to treat population.

Abbreviation: PSA, prostate specific antigen.

in the triptorelin study's intent-to-treat population (120 patients) achieved chemical castration at 1 month.

At month 12, 99% (102/103) of patients in the US leuprolide acetate trial had castrate testosterone levels as did 98.3% (113/115) of patients in the triptorelin trial. Mean time to castration in these trials was 21.2 days and 18.8 days (the latter calculated in a subset of 15 patients), respectively. Generally, daily and extended-release leuprolide acetate injections induce serum castrate testosterone in 3 to 4 weeks.^{27,30–32}

The leuprolide acetate trial also reported how many of their subjects reached serum testosterone levels ≤ 20 ng/dL. In the US leuprolide acetate trial, this figure was 88%. These data were not available for the triptorelin study.

Taken together, the efficacy rates for these two studies compare favorably to efficacy rates achieved by shorter-term GnRH agonist depot formulations. In particular, end-of-study castration (≤ 50 ng/dL) rates in published studies of existing 1-, 3-, and 4-month LA formulations were 100%, 100%, and 98%, respectively, in studies ranging from 6 to 8 months.^{32–34} Also, at the conclusion of these shorter-term GnRH analog studies, 94%, 94%, and 90% of patients had achieved testosterone levels ≤ 20 ng/dL. The castration rates for these studies after 1 month of therapy ranged between 94% and 98%. In a Phase III study of a 1-month, 3.75 mg LA depot formulation, 96.8% of patients sustained testosterone levels ≤ 50 ng/dL from month 1 through the study's 6-month duration.³⁵

At completion (final day) of the respective studies, the percentage of 1, 3, and 6-month triptorelin treated patients who were at castrate levels ranged between 97%–99%. Table 3 compares the results from the 1, 3, and 6-month depots.³⁶

Ultimately, the 6-month depot achieved similar results to the currently approved 1- and 3-month triptorelin formulations both on day 29 and from months 2 through 9. Using

Table 3 Comparison of 6-month vs shorter depots leuprolide acetate and triptorelin pamoate^{2,6,37–40}

	Day 29	Study completion
Leuprolide		
1-month depot	94.0%	100.0%
3-month depot	98.0%	100.0%
6-month depot	99.0%	99%
Triptorelin		
1-month depot	92.7%	99%
3-month depot	97.7%	97%
6-month depot	97.5%	98%

day 29 data, the 6-month depot achieved 97.5% (95% CI: 92.8%–99.5%), the 1-month depot achieved 92.7% (95% CI: 87.6%–96.2%), and the 3-month depot achieved 97.7% (95% CI: 94.1%–99.4%).

The success rates observed in studies of 6-month GnRH analog depots also correlates well with the low failure rates observed in studies of 1- and 3-month formulations of leuprolide acetate and goserelin acetate. In these studies, 2% to 15% of patients failed to achieve serum testosterone levels below 50 ng/dL. In addition 13% to 46.4% of patients did not achieve serum testosterone levels below 20 ng/dL.^{37–41}

Furthermore, the US leuprolide acetate trial and the triptorelin trial characterized initial testosterone flares/surges within their patient populations. Such surges have been associated with potentially serious worsening of symptoms such as bone pain, urinary obstruction, spinal cord compression, cardiovascular risk, and possibly death.^{42,43} Administering agents including flutamide, bicalutamide, nilutamide, diethylstilbestrol, ketoconazole, and cyproterone acetate can block some of the consequences of testosterone flares.⁴²

In the US leuprolide acetate study, mean testosterone levels rose 225 ng/dL by 2 days after the first injection, which is similar to testosterone flare levels reported for other GnRH agonist delivery systems.^{27,31} However, this testosterone increase was not associated with greater leuprolide acetate C_{max} , and no patients reported clinically relevant flare reactions to the initial testosterone increase. Accordingly, study authors suggest that higher-dose products carry no greater risk of tumor flare response than do lower-dose, shorter-term products.²

Among 28 patients in this trial selected for PK analysis, 26 completed the study and received both injections. PK data revealed an initial release of LA after each injection, with C_{max} serum values of 82.0 ng/mL 4.4 hours after the first injection and 102.4 ng/mL 4.8 hours after the second injection. Subsequently, serum leuprolide acetate slowly decreased.

During the plateau phase (days 3 to 168), serum leuprolide acetate averaged between 0.2 and 2.0 ng/mL with a 6-month mean value of 0.20 ± 0.14 ng/mL (median: 0.16). A similar pattern emerged after the second injection.

These data essentially mirror PK patterns observed in other leuprolide acetate studies, specifically the 7.5 mg 1-month formulation, a 22.5 mg 3-month formulation, and a 30 mg 4-month formulation. These products reached C_{max} approximately 4.66, 5, and 3.3 hours after injection, respectively, followed by gradual leuprolide acetate declines.^{32,44,45}

Similarly, investigators in the triptorelin trial selected a 15-patient subset in which they assessed triptorelin's pharmacokinetics by measuring serum testosterone levels at baseline and monthly thereafter, as well as on days 2, 3, 5, 8, 15, and 22. Among all treated patients, mean LH levels rose from 7.9 to 38.3 IU 2 hours after the first injection. In keeping with this initial pituitary LH stimulation,⁶ an initial increase in testosterone levels (C_{max} 25.8 nmol/L) occurred after the first injection in the 15 patients analyzed for pharmacodynamic and pharmacokinetic data. However, a rapid and sustained decrease in testosterone levels followed.

The “acute-on-chronic” or microsurge effect refers to agonistic testosterone stimulation occurring in response to serial injections of GnRH agonist depots.⁴⁶ In the triptorelin trial, only 2 of 60 patients whom investigators selected to assess this effect showed an acute-on-chronic increase in testosterone greater than 50 ng/dL 48 hours after the second injection.

Moreover, 2 hours after the second triptorelin injection, investigators observed virtually no LH increases from a fully suppressed level to only 0.1 IU/L. Only two patients failed to achieve complete pituitary desensitization; one of them was severely obese. The second experienced an increase of just 1.1 IU/L.

Consistent with triptorelin's success in maintaining long-term castration, PSA in this study fell by a median of 97% by month 6 and 96% at month 12. Similarly, at month 12 in the

Table 4 Advantages of 6-month depots

Advantage	Results
Fewer frequent injections	Reduced anxiety
Fewer doctor visits	Decreased emotional burden
	Improved flexibility with scheduling
	Improved comfort
	Decreased site reactions
	Decreased cost
	Less missed visits
	Decreased breakthrough (theoretical)

US leuprolide acetate trial, mean PSA overall decreased by 90.2%, and only 4 of 103 patients had increased PSA versus baseline (39.8 ± 1.5 ng/mL).

Testosterone escapes or leaks consist of testosterone elevations above 50 ng/dL in patients on continuous GnRH analog therapy.⁹ In published reports, the long-term rate of such leaks ranges between 4% and 12.5%.^{41,47,48} Perhaps the clearest evidence of the clinical consequences of such leaks comes from a retrospective study which showed that the greater the leak, the shorter a patient's progression-free survival.²⁹

Among the two trials of 6-month GnRH agonists, the triptorelin trial presented the most detailed testosterone escape data. Specifically, eight patients who completed the study failed to maintain castration at all visits between months 2 and 12. Five of these patients had only an isolated testosterone escape with no increase in serum PSA. Investigators considered three of these five events minimal, with one serum testosterone level measuring 67.15 ng/dL at month 2, and two serum testosterone levels measuring 55.91 ng/dL and 56.77 ng/dL at month 6. The other two isolated escapes occurred at month 4 and measured 96.54 ng/dL and 176.37 ng/dL.

Clear clinical failures occurred in three patients, as reflected by increases in serum PSA levels. In one of these patients (the obese patient previously mentioned), the first triptorelin injection failed to achieve clinical castration, although the second injection succeeded. The other two patients escaped castration from month 9 (serum testosterone 150 ng/dL) and at month 12 (1213 ng/dL), respectively.

Safety

As with existing GnRH analogs, side effects associated with 6-month depot formulations are generally mild and stem from testosterone suppression.²⁹ Among the two studies, adverse event (AE) reporting rates ranged from 74% in the US leuprolide acetate study (82/111 patients) to 95% in the triptorelin study (115/120 patients). In the latter study, investigators judged 86.7% of these events to be mild.

Hot flashes represented the most commonly reported AE in the two studies, occurring at rates ranging from 57.6% (33.3% mild, 24.3% moderate) of patients in the leuprolide acetate study to 71.7% in the triptorelin study. In studies of existing 1-, 3-, and 4-month LA formulations, hot flashes (mostly mild) also ranked as the most common AE, occurring in 56.7%, 59%, and 78.9% of patients, respectively.³²⁻³⁴

Rates of injection site reactions were fairly similar in both studies of 6-month depots, and the vast majority of

these reactions were considered mild. The lowest rate of injection site reactions, 6.7%, occurred in the triptorelin trial. Conversely, 15.3% of patients in the leuprolide acetate trial experienced injection-site burning.

SAE rates were relatively low in both studies. The triptorelin trial SAE rate was approximately 14% (17 of 120 patients). Only one patient in the US leuprolide acetate trial experienced an SAE, although investigators did not report whether it was related to the study medication.

None of the SAEs in the triptorelin was judged by investigators to be related to study medications. Additionally, investigators in the triptorelin study observed clinically significant treatment-emergent laboratory abnormalities in nine patients. However, only two mild events in the same patient (increased alanine transaminase/ALT and aspartate transaminase/AST) were related to the study drug. These investigators also reported 17 events of hypertension or its worsening, but considered only one to be drug-related. Furthermore, the rate of hypertension events was likely to be high due to 62% baseline hypertension.⁶

Conclusion

For patients with locally advanced or metastatic prostate cancer, hormonal therapy has long been a mainstay of palliative treatment.^{8,16,49,50} Additionally, physicians frequently integrate androgen deprivation with radiotherapy in certain intermediate and high risk patients with localized disease.⁸ In use for more than two decades, GnRH agonists represent the most frequently chosen hormonal therapy for achieving androgen deprivation in patients with prostate cancer.⁵¹⁻⁵⁷

Depot formulations of GnRH agonists have proven preferable to earlier options including daily injections and bilateral orchiectomy for several reasons. These include the reversibility of chemical castration⁵⁸ and, for patients, the ability that injectable GnRH agonists provide to avoid psychological and other comorbidities associated with orchiectomy.⁵⁹

Introduced to the US market in 1989, GnRH agonist depot formulations have evolved to offer treatment intervals of 1, 3, 4, and 6 months, all of which offer similarly acceptable safety and efficacy profiles. Monthly and daily formulations increase the likelihood that patients may delay or miss treatments, which can result in testosterone breakthrough and potentially compromise tumor control and increase symptom progression.⁸ Accordingly, 3- and 4-month depot leuprolide acetate formulations represent the most commonly used hormonal treatments for prostate cancer.⁸ Longer-term formulations give patients the flexibility to choose an

administration schedule that works for them so that patients miss fewer visits, have fewer doctor's visits, and have less frequent injections. This in turn allows for flexibility in scheduling a treatment plan and has the potential to decrease emotional burden and anxiety. Thus, longer acting depots have the potential to decrease injection site reaction, decrease cost, improve comfort, increase compliance and theoretically decrease tumor breakthrough.^{5,32,33,46,47,60–62}

Regarding potential indications for 6-month GnRH agonist depots, research has evaluated shorter-duration GnRH agonists as adjuvant and neoadjuvant treatments for patients undergoing radical prostatectomy and radiation therapy.^{53,63,64} One reviewer predicts that 6-month triptorelin pamoate will prove a valuable adjunct to radiation therapy and chemotherapy because it possesses similar efficacy, but fewer side effects, than adjunctive therapies now commonly available.⁶⁵

Moreover, two clinical trials have shown that depot products often suppress testosterone for longer than the labeled interval.^{66,67} Accordingly, researchers have begun investigating the clinical merits of intermittent androgen deprivation therapy (IADT) with GnRH agonists. Such strategies involve periodic evaluation of serum testosterone levels to guide injection intervals, and to detect nonresponders and testosterone breakthroughs.⁸ Once a patient reaches castrate testosterone levels, the physician suspends GnRH agonist therapy until PSA rises to a set threshold, at which point therapy resumes.³⁵ This approach can help patients avoid morbidity associated with continuous androgen deprivation and may forestall the development of hormone resistance.^{68,69}

The appropriateness of 6-month GnRH agonists has not yet been evaluated in the context of IADT or as adjuvant/neoadjuvant therapy. Also yet to be evaluated is whether or not 6-month LA depots require dosing adjustments in special patient populations such as patients experiencing kidney or liver failure.⁸

Additionally, research has not yet established whether a castration target of serum testosterone below 50 ng/dL or below 20 ng/dL provides clinical benefit. In one report, breakthrough increases above 32 ng/dL predicted shorter survival free of androgen-independent prostate cancer progression.²⁹

Finally, a randomized, controlled trial pitting the 6-month GnRH agonist against each other would further illuminate which product works best in which clinical situations. However, it is unlikely that such a trial will be performed. We look forward to learning more in all the above areas as physicians' and researchers' experience with 6-month depot products accumulates.

Disclosure

ED Crawford has financial interests and/or other relationships with GlaxoSmithKline, Sanofi-aventis, the National Institutes of Health, the University of Colorado Cancer Center, and Watson Pharmaceuticals. J Phillips has nothing to disclose. Editorial assistance provided by Carden Jennings funded by Watson Pharmaceuticals, Inc.

References

1. Prostate Cancer Foundation. Report to the Nation on Prostate Cancer 2004. Available at: <http://www.medscape.org/viewcollection/30050>. Accessed January 6, 2011.
2. Crawford ED, Sartor O, Chu F, Perez R, Karlin G, Garrett JS. A 12-month clinical study of LA-2585 (45.0 mg): a new 6-month subcutaneous delivery system for leuprolide acetate for the treatment of prostate cancer. *J Urol*. 2006;175:533–536.
3. Palmberg C, Koivisto P, Visakorpi T, Tammela T. PSA decline is an independent prognostic marker in hormonally treated prostate cancer. *Eur Urol*. 1999;36:191–196.
4. Cox MC, Scripture CD, Figg WD. Leuprolide acetate given by a subcutaneous extended-release injection: less of a pain? *Expert Rev Anticancer Ther*. 2005;5:605–611.
5. Sharifi R, Bruskewitz RC, Gittleman MC, Graham SD Jr, Hudson PB, Stein B. Leuprolide acetate 22.5 mg 12-week depot formulation in the treatment of patients with advanced prostate cancer. *Clin Ther*. 1996;18:647–657.
6. Lundström EA, Rencken RK, van Wyk JH, et al. Triptorelin 6-month formulation in the management of patients with locally advanced and metastatic prostate cancer: an open-label, non-comparative, multicentre, phase III study. *Clin Drug Investig*. 2009;29:757–765.
7. Huggins C, Hodges C. Studies on prostate cancer: the effect of castration, estrogen and of androgen injection on serum phosphatases in metastatic carcinoma of the prostate. *Cancer Res*. 1941;1:293–297.
8. Sethi R, Sanfilippo N. Six-month depot formulation of leuporelin acetate in the treatment of prostate cancer. *Clin Interv Aging*. 2009;4:259–267.
9. Amo FH. Comparative analysis of six months formulation of LHRH analogues for prostate cancer treatment. *Arch Esp Urol*. 2010;63:275–281.
10. Schally AV, Arimura A, Baba Y, et al. Isolation and properties of the FSH and LH-releasing hormone. *Biochem Biophys Res Commun*. 1971;43:393–399.
11. Fujino M, Fukuda T, Shinagawa S, Kobayashi S, Yamazaki I. Synthetic analogs of luteinizing hormone releasing hormone (LH-RH) substituted in position 6 and 10. *Biochem Biophys Res Commun*. 1974;60:406–413.
12. Monahan MW, Amoss MS, Anderson HA, Vale W. Synthetic analogs of the hypothalamic luteinizing hormone releasing factor with increased agonist or antagonist properties. *Biochemistry*. 1973;12:4616–4620.
13. Chrisp P, Sorkin EM. Leuporelin: a review of its pharmacology and therapeutic use in prostatic disorders. *Drugs Aging*. 1991;1:487–509.
14. Anderson J, Abrahamsson PA, Crawford D, Miller K, Tombal B. javascript: AL_get(this, 'jour', 'BJU Int. '); Management of advanced prostate cancer: can we improve on androgen deprivation therapy? *BJU Int*. 2008;101:1497–501.
15. Millar RP, Troskie BE, Flanagan CA. Comparative receptor binding affinity and inositol phosphate production potency of D-Leu6 and D-Trp6 GnRH agonists on COS-1 cells transfected with the human GnRH receptor. *XIII International Congress of Comparative Endocrinology*. 1997:559–562.
16. Miller K, Anderson J, Abrahamsson P-A. Treatment of prostate cancer with hormonal therapy in Europe. *BJU Int*. 2009;103(Suppl 2):2–6.

17. National Comprehensive Cancer Network. NCCN clinical practice guidelines in oncology: prostate cancer v.2.2009. Available at: http://www.nccn.org/professionals/physician_gls/pdf/prostate.pdf. Accessed December 9, 2009.
18. Lahlou N. [Pharmacokinetics and pharmacodynamics of triptorelin]. *Ann Urol (Paris)*. 2005;39(Suppl 3):78–84. French.
19. Coy DH, Vilchez-Martinez JA, Coy EJ, et al. Analogs of luteinizing hormone-releasing hormone with increased biological activity produced by D-amino acid substitutions in position 6. *J Med Chem*. 1976;19:423–425.
20. Barron JL, Miller RP, Searle D. Metabolic clearance and plasma half-disappearance time of D-TRP6 and exogenous luteinizing hormone-releasing hormone. *J Clin Endocrinol Metab*. 1982;54:1169–1173.
21. Müller FO, Terblanché J, Schall R, et al. Pharmacokinetics of triptorelin after intravenous bolus administration in healthy males and in males with renal or hepatic insufficiency. *Br J Clin Pharmacol*. 1997;44:335–341.
22. Sennello LT, Finley RA, Chu SY, et al. Single-dose pharmacokinetics of leuprolide in humans following intravenous and subcutaneous administration. *J Pharm Sci*. 1986;75:158–160.
23. Periti P, Mazzei T, Mini E. Clinical pharmacokinetics of depot leuprolide. *Clin Pharmacokinet*. 2002;41:485–504.
24. Lee M, Browneller R, Wu Z, Jung A, Ratanawong C, Sharifi R. Therapeutic effects of leuprolide microspheres in prostate cancer. *Adv Drug Deliv Rev*. 1997;28:121–138.
25. Akaza H, Usami M, Koiso K, Kotake T, Aso Y, Nijima T. Long-term clinical study on luteinizing hormone-releasing hormone agonist depot formulation in the treatment of stage D prostatic cancer. The TAP-144-SR Study Group. *Jpn J Clin Oncol*. 1992;22:177–184.
26. Fowler JE, Flanagan M, Gleason DM, Klimberg IW, Gottesman JE, Sharifi R. Evaluation of an implant that delivers leuprolide for 1 year for the palliative treatment of prostate cancer. *Urology*. 2000;55:639–642.
27. Fowler JE Jr, Gottesman JE, Reid CF, Andriole GL Jr, Soloway MS. Safety and efficacy of an implantable leuprolide delivery system in patients with advanced prostate cancer. *J Urol*. 2000;164(3 Pt 1):730–734.
28. Oefelein MG, Feng A, Scolieri MJ, Ricchiutti D, Resnick MI. Reassessment of the definition of castrate levels of testosterone: implications for clinical decision making. *Urology*. 2000;56:1021–1024.
29. Morote J, Orsola A, Planas J, Trilla E, Raventós CX, Cecchini L, Catalán R. Redefining clinically significant castration levels in patients with prostate cancer receiving continuous androgen deprivation therapy. *J Urol*. 2007;178(4 Pt 1):1290–1295.
30. Tunn UW, Bargelloni U, Cosciani S, Fiaccavento G, Guazzieri S, Pagano F. Comparison of LH-RH analogue 1-month depot and 3-month depot by their hormone levels and pharmacokinetic profile in patients with advanced prostate cancer. *Urol Int*. 1998;60(Suppl 1):9–16; discussion 16–17.
31. Sharifi R, Knoll LD, Smith J, Kramolowsky E. Leuprolide acetate (30-mg depot every four months) in the treatment of advanced prostate cancer. *Urology*. 1998;51:271–276.
32. Perez-Marreno R, Chu FM, Gleason D, Loizides E, Wachs B, Tyler RC. A six-month, open-label study assessing a new formulation of leuprolide 7.5 mg for suppression of testosterone in patients with prostate cancer. *Clin Ther*. 2002;24:1902–1914.
33. Chu FM, Jayson M, Dineen MK, Perez R, Harkaway R, Tyler RC. A clinical study of 22.5 mg. La-2550: a new subcutaneous depot delivery system for leuprolide acetate for the treatment of prostate cancer. *J Urol*. 2002;168:1199–1203.
34. Sartor O, Dineen MK, Perez-Marreno R, Chu FM, Carron GJ, Tyler RC. An eight-month clinical study of LA-2575 30.0 mg: a new 4-month, subcutaneous delivery system for leuprolide acetate in the treatment of prostate cancer. *Urology*. 2003;62:319–323.
35. Marberger M, Kaisary AV, Shore ND, et al. Effectiveness, pharmacokinetics, and safety of a new sustained-release leuprolide acetate 3.75-mg depot formulation for testosterone suppression in patients with prostate cancer: a Phase III, open-label, international multicenter study. *Clin Ther*. 2010;32:744–757.
36. Clinical study report. Comparative testosterone pharmacodynamics and therapeutic efficacy of 1- and 3-month formulations of triptorelin pamoate in patients with advanced prostate cancer. DEB-96-TR1-01 (first phase). Lausanne, Switzerland: Debiopharm SA; 1999.
37. Morote J, Esquena S, Abascal JM, et al. Failure to maintain a suppressed level of serum testosterone during long-acting depot luteinizing hormone-releasing hormone agonist therapy in patients with advanced prostate cancer. *Urol Int*. 2006;77:135–138.
38. Sarosdy MF, Schellhammer PF, Soloway MS, et al. Endocrine effects, efficacy and tolerability of a 10.8-mg depot formulation of goserelin acetate administered every 13 weeks to patients with advanced prostate cancer. *BJU Int*. 1999;83:801–806.
39. Oefelein MG, Cornum R. Failure to achieve castrate levels of testosterone during luteinizing hormone releasing hormone agonist therapy: the case for monitoring serum testosterone and a treatment decision algorithm. *J Urol*. 2000;164(3 Pt 1):726–729.
40. Wechsel HW, Zerbib M, Pagano F, Coptcoat MJ. Randomized open labelled comparative study of the efficacy, safety and tolerability of leuprolide acetate 1M and 3M depot in patients with advanced prostatic cancer. *Eur Urol*. 1996;30(Suppl 1):7–14.
41. Jocham D. Leuprolide 3-month depot in the treatment of advanced and metastatic prostate cancer: long-term follow-up results. *Urol Int*. 1998;60(Suppl 2):18–24; discussion 35.
42. Thompson IM. Flare Associated with LHRH-Agonist Therapy. *Rev Urol*. 2001;3(Suppl 3):S10–S14.
43. Bubley GJ. Is the flare phenomenon clinically significant? *Urology*. 2001;58(2 Suppl 1):5–9.
44. Eligard 225 mg: leuprolide acetate for injectable suspension [package insert]. Sanofi-aventis; 2004.
45. Eligard 30 mg: leuprolide acetate for injectable suspension [package insert]. Sanofi-aventis; 2004.
46. Sharifi R, Browneller R; Leuprolide Study Group. Serum testosterone suppression and potential for agonistic stimulation during chronic treatment with monthly and 3-month depot formulations of leuprolide acetate for advanced prostate cancer. *J Urol*. 2002;168:1001–1004.
47. Khan MS, O'Brien A. An evaluation of pharmacokinetics and pharmacodynamics of leuprolide acetate 3M-depot in patients with advanced and metastatic carcinoma of the prostate. *Urol Int*. 1998;60:33–40.
48. Fontana D, Mari M, Martinelli A, et al. 3-month formulation of goserelin acetate ('Zoladex' 10.8-mg depot) in advanced prostate cancer: results from an Italian, open, multicenter trial. *Urol Int*. 2003;70:316–320.
49. Horwich A, Parker C, Kataja V. Prostate cancer: ESMO clinical recommendations for diagnosis, treatment and follow-up. *Ann Oncol*. 2008;19(Suppl 2):45–46.
50. National Institute for Health and Clinical Excellence. Prostate cancer: diagnosis and treatment. Available at: <http://www.nice.org.uk/nicemedia/live/11924/39687/39687.pdf>. Accessed January 7, 2011.
51. Moretti RM, Monagnani Marelli M, van Groeninghen JC, Motta M, Limonta P. Inhibitory activity of luteinizing hormone-releasing hormone on tumor growth and progression. *Endocr Relat Cancer*. 2003;10:161–167.
52. D'Amico AV, Manola J, Loffredo M, Renshaw AA, Della Croce A, Kantoff PW. 6-month androgen suppression plus radiation therapy vs radiation therapy alone for patients with clinically localized prostate cancer: a randomized controlled trial. *JAMA*. 2004;292:821–827.
53. Gleave ME, Goldenberg SL, Chin JL, et al. Randomized comparative study of 3 versus 8-month neoadjuvant hormonal therapy before radical prostatectomy: biochemical and pathological effects. *J Urol*. 2001;166:500–5006; discussion 506–507.
54. Smith MR. Androgen deprivation therapy for prostate cancer: new concepts and concerns. *Curr Opin Endocrinol Diabetes Obes*. 2007;14:247–254.
55. De Jong IJ, Eaton A, Bladou F. LHRH agonists in prostate cancer: frequency of treatment, serum testosterone measurement and castrate level: consensus opinion from a roundtable discussion. *Curr Med Res Opin*. 2007;23:1077–1080.
56. Waxman JH, Wass JA, Hendry WF, et al. Treatment with gonadotrophin releasing hormone analogue in advanced prostatic cancer. *Br Med J (Clin Res Ed)*. 1983;286:1309–1312.

57. Mcleod DG. Hormonal therapy: historical perspective to future directions. *Urology*. 2003;61(2 Suppl 1):3–7.
58. Bergquist C, Nillius SJ, Bergh T, Skarin G, Wide L. Inhibitory effects on gonadotrophin secretion and gonadal function in men during chronic treatment with a potent stimulatory luteinizing hormone-releasing hormone analogue. *Acta Endocrinol (Copenh)*. 1979;91:601–608.
59. Vogelzang NJ, Chodak GW, Soloway MS, et al. Goserelin versus orchiectomy in the treatment of advanced prostate cancer: final results of a randomized trial. Zoladex Prostate Study Group. *Urology*. 1995;46:220–226.
60. Sharifi R, Soloway M; Leuprolide Study Group. Clinical study of leuprolide depot formulation in the treatment of advanced prostate cancer. *J Urol*. 1990;143:68–71.
61. Sharifi R, Gulley JL, Dahut WL. Androgen deprivation therapy for prostate cancer. *JAMA*. 2005;294:238–244.
62. Kienle E, Lübben G; German Leuprorelin Study Group. Efficacy and safety of leuprorelin acetate depot for prostate cancer. *Urol Int*. 1996;56(Suppl 1):23–30.
63. Persad R. Leuprorelin acetate in prostate cancer: a European update. *Int J Clin Pract*. 2002;56:389–396.
64. Hellerstedt BA, Pienta KJ. The current state of hormonal therapy for prostate cancer. *Cancer J Clin*. 2002;52:154–179.
65. Whelan P. Triptorelin embonate: a 6-month formulation for prostate cancer. *Expert Opin Pharmacother*. 2010;11:2929–2932.
66. Pathak AS, Pacificar JS, Shapiro CE, Williams SG. Determining dosing intervals for luteinizing hormone releasing hormone agonists based on serum testosterone levels: a prospective study. *J Urol*. 2007;177: 2132–2135; discussion 2135.
67. Greil S, Robinson EA, Singal B, Kleer E. Efficacy over time of LHRH analogs in the treatment of PCa—a prospective analysis using serum testosterone to determine dosing intervals. *Urology*. 2009;73: 631–634.
68. Shore ND, Crawford ED. Intermittent androgen deprivation therapy: redefining the standard of care? *Rev Urol*. 2010;12:1–11.
69. Calais da Silva FEC, Bono AV, Whelan P, et al. Intermittent androgen deprivation for locally advanced and metastatic prostate cancer: results from a randomised phase 3 study of the South European Urooncological Group. *Eur Urol*. 2009;55:1269–1277.

Cancer Management and Research

Publish your work in this journal

Cancer Management and Research is an international, peer-reviewed open access journal focusing on cancer research and the optimal use of preventative and integrated treatment interventions to achieve improved outcomes, enhanced survival and quality of life for the cancer patient. The journal welcomes original research, clinical & epidemiological

Submit your manuscript here: <http://www.dovepress.com/cancer-management-and-research-journal>

studies, reviews & evaluations, guidelines, expert opinion & commentary, case reports & extended reports. The manuscript management system is completely online and includes a very quick and fair peer-review system, which is all easy to use. Visit <http://www.dovepress.com/testimonials.php> to read real quotes from published authors.

Dovepress