# BC Cancer Protocol Summary for Androgen Deprivation Therapy for Prostate Cancer

| Protocol Code | GUPADT        |
|---------------|---------------|
| Tumour Group  | Genitourinary |

**Contact Physician** 

Dr. Mira Keyes

## ELIGIBILITY:

Patients must have:

- Localized prostate cancer with recurrence risks or biochemical relapses
- Locally advanced or metastatic prostate adenocarcinoma

#### TESTS:

- Baseline: PSA, testosterone
- If clinically indicated: PSA, testosterone
- For patients taking flutamide, every 3 months: total bilirubin, ALT, alkaline phosphatase

#### TREATMENT:

| Drug   | Dose and BC Cancer Administration Guideline  |  |
|--|--|--|
| goserelin long acting  |  |  |
| (ZOLADEX)  | <ul> <li>3.6 mg subcutaneous every month, or</li> </ul>  |  |
| (ZOLADEX LA)   | <ul> <li>10.8 mg subcutaneous every 3 months</li> </ul>  |  |
| OR   |  |  |
| leuprolide long acting   |  |  |
|  | <ul> <li>7.5 mg IM every month, or</li> </ul>  |  |
| (LUPRON DEPOT)   | <ul> <li>22.5 mg IM every 3 months, or</li> </ul>  |  |
|  | <ul> <li>30 mg IM every 4 months, or</li> </ul>  |  |
|  |  |  |
| (ELIGARD)  | <ul> <li>7.5 mg subcutaneous every month, or</li> </ul>  |  |
|  | <ul> <li>22.5 mg subcutaneous every 3 months, or</li> </ul>  |  |
|  | <ul> <li>30 mg subcutaneous every 4 months, or</li> </ul>  |  |
|  | 45 mg subcutaneous every 6 months  |  |
| OR   |  |  |
| degarelix  | <ul> <li>Starting dose: 240 mg subcutaneous* (as two injections of<br/>120 mg) on day 1, followed by</li> </ul>                          |  |
|  | <ul> <li>Maintenance dose: 80 mg subcutaneous (as a single<br/>injection) every month, starting one month after starting dose</li> </ul> |  |
| *Injections to be given in the abdominal region. To reduce incidence of injection site |  |  |

\*Injections to be given in the abdominal region. To reduce incidence of injection site reactions: inject slowly; leave needle in place for 30 seconds after injection and then withdraw needle slowly.

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Revised:

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**Treatment duration:** depends on the indication. May continue in castrate resistant disease.

| Drug           | Dose        | BC Cancer Administration Guideline |
|----------------|-------------|------------------------------------|
| bicalutamide** | 50 mg daily | PO                                 |
| OR             |             |                                    |
| flutamide      | 250 mg TID  | PO                                 |

#### If required, add on oral anti-androgen agent:

\*\* Preferred anti-androgen

- Degarelix does not induce a testosterone surge or clinical flare, therefore antiandrogen use with degarelix is not required.
- To block clinical flare to LHRH agonist, oral anti-androgen agent should be started at least 1 to 2 weeks prior to first LHRH agonist dose.

#### Oral anti-androgen treatment duration:

- To block clinical flare to LHRH agonist: 3 to 4 weeks.
- Other indications: May be continued for as long as patient is treated with LHRH agonist.

### **PRECAUTIONS:**

- 1. **Disease flare** can occur during the first few days of LHRH agonist therapy when the testosterone level may be elevated. Patients should receive an antiandrogen for 3 to 4 weeks with the initial dose.
- 2. **Androgen deprivation** may cause hot flashes, impotence, gynecomastia, erythema and irritation of the injection site. Muscle weakness and weight gain. Adverse CNS effects occur in 3% or more patients, including dizziness, pain, headache and paresthesias. Mood changes. Prolonged suppression of testosterone may occur in the elderly (over 75 years) and with the longer lasting preparations.
- 3. **Increased risks of cardiac events:** Androgen deprivation therapy may increase cardiovascular risk in men with prostate cancer. Physicians should assess cardiovascular risk and manage as per clinical practice guidelines.
- 4. **Osteoporosis:** Androgen deprivation therapy may cause an increased risk of osteoporosis and fractures. Physicians should assess osteoporosis risk and manage as per clinical practice guidelines.
- 5. **Diarrhea:** discontinue flutamide if diarrhea develops.

# Call Dr. Mira Keyes or tumour group delegate at (604) 877-6000 or 1-800-663-3333 with any problems or questions regarding this treatment program.

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#### References

- Loblaw DA, Virgo KS, Nam R, et al. Initial hormonal management of androgen-sensitive metastatic, recurrent, or progressive prostate cancer: 2006 update of an American Society of Clinical Oncology practice guideline. J Clin Oncol 2007;25(12):1596-605.
- 2. Cornford P, van den Bergh RCN, Briers E, et al. EAU-EANM-ESTRO-ESUR-SIOG guidelines on prostate cancer. Part II-2020 update: treatment of relapsing and metastatic prostate cancer. Eur Urol 2021;79(2):263-282.
- 3. Lowrance WT, Breau RH, Chou R, et al. Advanced prostate cancer: AUA/ASTRO/SUO Guideline PART I. J Urol 2021;205(1):14-21.
- Bekelman JE, Rumble RB, Chen RC, et al. Clinically localized prostate cancer: ASCO clinical practice guideline endorsement of an American Urological Association/American Society for Radiation Oncology/Society of Urologic Oncology guideline. J Clin Oncol 2018;36(32):3251-3258.
- 5. Eastham JA, Auffenberg GB, Barocas DA, et al. Clinically localized prostate cancer: AUA/ASTRO guideline. Part III: principles of radiation and future directions. J Urol 2022;208(1):26-33.