# **BC Cancer Protocol Summary for Neoadjuvant or Adjuvant Therapy** for Breast Cancer Using Tamoxifen

Protocol Code BRAJTAM

**Tumour Group** Breast

Dr. Susan Ellard Contact Physician

#### **ELIGIBILITY**:

- may be given preoperatively in hormone receptor positive breast cancer patients who are unsuitable for immediate surgery or preoperative chemotherapy
- Adjuvant hormonal treatment for breast cancer, initiated up to 10 years after diagnosis and treatment
  - All premenopausal hormone receptor-positive women: upfront tamoxifen for up to a total of 10 years only (eligible if completed 5 yrs of therapy within last 12 months)
- Options for postmenopausal hormone receptor positive invasive breast cancer:
  - Upfront tamoxifen for up to a total of 10 years only (eligible if completed 5 yrs of therapy within last 12 months)
    - Consider aromatase inhibitor (AI) options below if disease higher than T1N0 low grade tumours
    - Any postmenopausal hormone receptor positive invasive breast cancer in patients intolerant to aromatase inhibitors
  - Early switch: 2-3 years of adjuvant Tamoxifen to begin 5 years of hormone blockade (except T1N0 low grade disease)
  - Late switch: 5 years of adjuvant Tamoxifen, followed by up to 5 additional years of letrozole, or 3 additional years of anastrozole or exemestane (except T1N0 low grade) Note: patients who started on letrozole but had to switch to another aromatase inhibitor due to side effects may also continue to 5 years with CAP approval only
- See Cancer Management Guidelines for current guidelines.

#### **EXCLUSIONS:**

- Hormone receptor-negative
- Patients with a history of significant thromboembolic disease

# TESTS:

- Annually: gynecological exam (postmenopausal patients with an intact uterus)
- If clinically indicated (see PRECAUTIONS, below): CBC and diff, platelets, serum cholesterol and triglycerides, bilirubin, alk phos, ALT, GGT, ophthalmologic exam

#### TREATMENT:

# Upfront:

tamoxifen 20 mg PO daily x up to a total of 10 years

# Early Switch:

tamoxifen 20 mg PO daily x 2-3 years, followed by BRAJANAS, BRAJEXE, or BRAJLET for a total treatment time of 5 years.

# Late Switch:

tamoxifen 20 mg PO daily x 5 years, followed by BRAJANAS, BRAJEXE, or BRAJLET for a further 3 years of treatment (up to 5 years if letrozole – see BRAJLET)

## **MODIFICATIONS:**

- 1. Intolerant or serious complications during tamoxifen therapy
  - Post-menopausal patients may be switched to BRAJANAS, BRAJEXE. BRAJLET for a total of 5 years of adjuvant hormonal therapy

#### PRECAUTIONS:

- 1. Myelosuppression: Mild myelosuppression with transient thrombocytopenia may occur rarely. The association with tamoxifen is uncertain.
- 2. Endometrial Cancer: Annual gynecological examinations are recommended. Pelvic complaints, such as unusual vaginal bleeding, require prompt evaluation.
- 3. Ocular Toxicity: Ocular toxicity is rare and may occur after only a few weeks of therapy, although it is more common with prolonged treatment. Ophthalmologic examination is recommended if visual disturbances occur.
- 4. Thromboembolism: Tamoxifen is associated with an increased risk of thromboembolism that is comparable to estrogen replacement therapy.
- 5. Hepatotoxicity: While hepatotoxicity is rare and usually presents as elevated hepatic enzymes, more serious liver abnormalities have been reported.
- 6. Ovulation Induction: Tamoxifen may induce ovulation in pre- and peri-menopausal women. Barrier forms of contraception are recommended.
- 7. Hyperlipidemia: Elevations in cholesterol and triglycerides may occur in patients with pre-existing hyperlipidemias.

Call Dr. Susan Ellard or tumour group delegate at (250) 712-3900 or 1-888-563-7773 with any problems or questions regarding this treatment program.

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## References:

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- Delozier T, Switsers O, Genot JY et al. Delayed adjuvant tamoxifen: ten-year results of a collaborative randomized controlled trial in early breast cancer (TAM-02 trial). Ann Oncol 2000;11:515-9.
- 3. Coombes, RC et al. A randomized trial of exemestane after two to three years of tamoxifen therapy in postmenopausal women with primary breast cancer. N Engl J Med 2004; 350(11):1081-92
- Goss PE, Ingle JN, Martino S, et al. A randomized trial of letrozole in postmenopausal women after five years of tamoxifen therapy for early-stage breast cancer. N Engl J Med 2003;349(19):1793-02.
- 5. Davies C, Pan H, Godwin J, et al. Long-term effects of continuing adjuvant tamoxifen to 10 years versus stopping at 5 years after diagnosis of oestrogen receptor-positive breast cancer: ATLAS, a randomised trial. Lancet 2013;381(9869):805-16.