BC Cancer Protocol Summary for Symptomatic Management of Functional Carcinoid and Neuroendocrine Tumors of the GI Tract Using Octreotide (SANDOSTATIN LAR®)

Protocol Code: UGIOCTLAR

Tumour Group: GI

Contact Physician: GI Systemic Therapy

ELIGIBILITY:
- Functional neuroendocrine tumours of the mid-gut or pancreas (including carcinoid tumours and VIPoma), with symptoms due to ectopic hormone secretion (flushes, diarrhea, wheezing, etc.)
- A BC Cancer Compassionate Access Program (CAP) request with appropriate clinical information for each patient must be approved prior to treatment. CAP approval is valid as long as patient has symptomatic benefits.

EXCLUSIONS:
- Pregnant or lactating women

TESTS:
- No specific tests apart from those required to monitor the underlying disease
- A pretreatment ultrasound of the gall bladder is recommended to rule out the formation of gallstones. Repeat every 6 months while on therapy.

PREMEDICATIONS:
- Not usually required

TREATMENT:

<table>
<thead>
<tr>
<th>Drug</th>
<th>Dose</th>
<th>BC Cancer Administration Guideline</th>
</tr>
</thead>
<tbody>
<tr>
<td>octreotide</td>
<td>20 to 30 mg</td>
<td>Intramuscular (deep intragluteal*) injection</td>
</tr>
<tr>
<td>(SANDOSTATIN LAR®)</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

*May use quadriceps for self-administration

Repeat every four weeks until tumour progression. Treatment may be started while patient is receiving octreotide daily SC injections. These should be continued for two weeks after the first octreotide monthly IM injection. However, initiation of monthly
octreotide LAR® does not necessarily require a subcutaneous octreotide trial and monitoring period.

**DOSE MODIFICATIONS:**
For patients in whom symptoms are not fully controlled within the 4 weeks, the dose of octreotide (SANDOSTATIN LAR®) may be increased incrementally to a maximum of 60 mg every 4 weeks. For patients with breakthrough symptoms, every 3 week dosing may be considered. However, total monthly dosing greater than 60 mg requires CAP approval.

**PRECAUTIONS:**
- **Concomitant diabetes:** Patients on oral hypoglycemics or insulin should be monitored closely for changes in blood glucose levels for several days after the start of octreotide (SANDOSTATIN LAR®) to determine the need for any dosage adjustments.
- **Previous history of gallstones.**
- **Potential for some impairment in thyroid function:** monitor for signs and symptoms of hypothyroidism.

Call the GI Systemic Therapy physician at your regional cancer centre or the GI Systemic Therapy Chair Dr. Janine Davies at (604) 877-6000 or 1-800-670-3322 with any problems or questions regarding this treatment program.

References: