

Temsirolimus - Metastatic Renal Cell Carcinoma

(This form must be completed before the first dose is dispensed.)

1. Patient Profile

- * Surname:
- * Given Name:
- * OHIN: * Chart Number:
- * Postal Code:
- * Height (cm): * Weight (kg):
- * BSA (m²): * Gender: Male Female Other
- * Date of Birth:
Day Month Year
- * Site:
- * Attending Physician (MRP- Most Responsible Physician):
- Requested Prior Approval Yes * Patient on Clinical Trial Yes No
- Other (specify):
- Specify Arm:
 Standard of care arm Experimental arm
 Blinded / Unknown

Request prior approval for enrolment

- * Justification for Funding
-

2. Eligibility Criteria

The patient must meet the following criteria:

Patient has poor risk¹ metastatic renal cell carcinoma, independent of histology, and is being treated with temsirolimus in the first line setting Yes

3. Funded Dose

- Temsirolimus 25mg IV weekly until disease progression

4. Notes

1. Poor risk is defined using a modification of criteria from Mekhail et al (J Clin Oncol. 2005; 23:832-41). Patients must have at least 3 of the following features:
 - a. high lactate dehydrogenase (> 1.5 times upper limit of normal);
 - b. low hemoglobin (< lower limit of normal);
 - c. high corrected serum calcium (> 10mg/dL);
 - d. time from initial diagnosis to first treatment is less than 12 months;
 - e. poor performance status (Karnofsky performance status < 80);
 - f. metastases in multiple organ sites (e.g. lung, liver, retroperitoneal lymph nodes, etc.)

5. Supporting Documents

To ensure reimbursement of your claim, both the completed enrolment form and a copy of the required documentation (where applicable) must be submitted through CCO e-Claims.

Signature of Attending Physician (MRP-Most Responsible Physician):

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Day Month Year