

Use this checklist to determine if a patient meets the restrictions for funding in the **hospital setting**. For more details, refer to [Section H](#) of the Pharmaceutical Schedule. For community funding, see the [Special Authority Criteria](#).

**PREScriBER**

Name: .....

Ward: ..... NHI: .....

**Lenvatinib**

**INITIATION – thyroid cancer**

Re-assessment required after 6 months

**Prerequisites** (tick boxes where appropriate)

- Patient is currently on treatment with lenvatinib and met all remaining criteria prior to commencing treatment
- or
- The patient has locally advanced or metastatic differentiated thyroid cancer
- and
- Patient must have symptomatic progressive disease prior to treatment
- or
- Patient must have progressive disease at critical anatomical sites with a high risk of morbidity or mortality where local control cannot be achieved by other measures
- and
- A lesion without iodine uptake in a RAI scan
- or
- Receiving cumulative RAI greater than or equal to 600 mCi
- or
- Experiencing disease progression after a RAI treatment within 12 months
- or
- Experiencing disease progression after two RAI treatments administered within 12 months of each other
- and
- Patient has thyroid stimulating hormone (TSH) adequately suppressed
- and
- Patient is not a candidate for radiotherapy with curative intent
- and
- Surgery is clinically inappropriate
- and
- Patient has an ECOG performance status of 0-2

**CONTINUATION – thyroid cancer**

Re-assessment required after 6 months

**Prerequisites** (tick box where appropriate)

- There is no evidence of disease progression

I confirm that the above details are correct:

Signed: ..... Date: .....

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**PREScriBER**

Name: ..... Name: .....

Ward: ..... NHI: .....

**Lenvatinib** - *continued*

**INITIATION – unresectable hepatocellular carcinoma**

Re-assessment required after 6 months

**Prerequisites** (tick boxes where appropriate)

- Patient has unresectable hepatocellular carcinoma
- and
- Patient has preserved liver function (Childs-Pugh A)
- and
- Transarterial chemoembolisation (TACE) is unsuitable
- and
- Patient has an ECOG performance status of 0-2
- and
- Patient has not received prior systemic therapy for their disease in the palliative setting
- or
- Patient has experienced treatment-limiting toxicity from treatment with atezolizumab with bevacizumab
- and
- No disease progression since initiation of atezolizumab with bevacizumab

**CONTINUATION – unresectable hepatocellular carcinoma**

Re-assessment required after 6 months

**Prerequisites** (tick box where appropriate)

- There is no evidence of disease progression

**INITIATION – renal cell carcinoma**

Re-assessment required after 4 months

**Prerequisites** (tick boxes where appropriate)

- The patient has metastatic renal cell carcinoma
- and
- The disease is of predominant clear-cell histology
- and
- The patient has documented disease progression following one previous line of treatment
- and
- The patient has an ECOG performance status of 0-2
- and
- Lenvatinib is to be used in combination with everolimus
- or
- Patient has received funded treatment with nivolumab for the second line treatment of metastatic renal cell carcinoma
- and
- Patient has experienced treatment limiting toxicity from treatment with nivolumab
- and
- Lenvatinib is to be used in combination with everolimus
- and
- There is no evidence of disease progression

**CONTINUATION – renal cell carcinoma**

Re-assessment required after 4 months

**Prerequisites** (tick box where appropriate)

- There is no evidence of disease progression

I confirm that the above details are correct:

Signed: ..... Date: .....