

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: Name:

Ward: NHI:

Bevacizumab

INITIATION – unresectable hepatocellular carcinoma

Re-assessment required after 6 months

Prerequisites (tick boxes where appropriate)

- Patient is currently on treatment with bevacizumab, and met all remaining criteria prior to commencing treatment
- or**
- Patient has locally advanced or metastatic, unresectable hepatocellular carcinoma
 - and**
 - Patient has preserved liver function (Child-Pugh A)
 - and**
 - Transarterial chemoembolisation (TACE) is unsuitable
 - and**
 - Patient has not received prior systemic therapy for the treatment of hepatocellular carcinoma
 - or**
 - Patient received funded lenvatinib before 1 March 2025
 - or**
 - Patient has experienced treatment-limiting toxicity from treatment with lenvatinib
 - and**
 - No disease progression since initiation of lenvatinib
 - and**
 - Patient has an ECOG performance status of 0-2
 - and**
 - To be given in combination with atezolizumab

CONTINUATION – unresectable hepatocellular carcinoma

Re-assessment required after 6 months

Prerequisites (tick box where appropriate)

- No evidence of disease progression

INITIATION – advanced or metastatic ovarian cancer

Re-assessment required after 4 months

Prerequisites (tick boxes where appropriate)

- or**
- The patient has FIGO Stage IV epithelial ovarian, fallopian tube, or primary peritoneal cancer
 - or**
 - The patient has previously untreated advanced (FIGO Stage IIIB or IIIC) epithelial ovarian, fallopian tube, or primary peritoneal cancer
 - and**
 - Debulking surgery is inappropriate
 - or**
 - The cancer is sub-optimally debulked (maximum diameter of any gross residual disease greater than 1cm)
 - and**
 - Bevacizumab to be administered at a maximum dose of 15 mg/kg every three weeks
 - and**
 - 18 weeks concurrent treatment with chemotherapy is planned

I confirm that the above details are correct:

Signed: Date:

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PREScriBER

Name:

Ward: NHI:

Bevacizumab - *continued*

CONTINUATION – advanced or metastatic ovarian cancer

Re-assessment required after 4 months

Prerequisites (tick box where appropriate)

No evidence of disease progression

INITIATION – Recurrent Respiratory Papillomatosis

Re-assessment required after 12 months

Prerequisites (tick boxes where appropriate)

Maximum of 6 doses
and
 The patient has recurrent respiratory papillomatosis
and
 The treatment is for intra-lesional administration

CONTINUATION – Recurrent Respiratory Papillomatosis

Re-assessment required after 12 months

Prerequisites (tick boxes where appropriate)

Maximum of 6 doses
and
 The treatment is for intra-lesional administration
and
 There has been a reduction in surgical treatments or disease regrowth as a result of treatment

INITIATION – Ocular Conditions

Prerequisites (tick boxes where appropriate)

Ocular neovascularisation
or
 Exudative ocular angiopathy

I confirm that the above details are correct:

Signed: Date: