I confirm that the above details are correct:

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

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.

	PATIENT:
me:	Name:
rd:	NHI:
vacizur	ab
-assessm	unresectable hepatocellular carcinoma ent required after 6 months s (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
	O Patient has preserved liver function (Child-Pugh A)
	Transarterial chemoembolisation (TACE) is unsuitable
	O Patient has not received prior systemic therapy for the treatment of hepatocellular carcinoma or
	O Patient received funded lenvatinib before 1 March 2025  or
	Patient has experienced treatment-limiting toxicity from treatment with lenvatinib
	O No disease progression since initiation of lenvatinib
	nd Patient has an ECOG performance status of 0-2
	Patient has an ECOG performance status of 0-2
	O Patient has an ECOG performance status of 0-2
	O Patient has an ECOG performance status of 0-2  nd O To be given in combination with atezolizumab
DNTINUA'	Patient has an ECOG performance status of 0-2  To be given in combination with atezolizumab  ION – unresectable hepatocellular carcinoma ent required after 6 months
ONTINUA' assessmerequisite	Patient has an ECOG performance status of 0-2  To be given in combination with atezolizumab  ION – unresectable hepatocellular carcinoma ent required after 6 months s (tick box where appropriate)
ONTINUA' assessmerequisite	Patient has an ECOG performance status of 0-2  To be given in combination with atezolizumab  ION – unresectable hepatocellular carcinoma ent required after 6 months
ONTINUA assessm erequisite	Patient has an ECOG performance status of 0-2  To be given in combination with atezolizumab  ION – unresectable hepatocellular carcinoma ent required after 6 months s (tick box where appropriate)
ONTINUA: -assessmerequisite O No ITIATION -assessm	Patient has an ECOG performance status of 0-2  To be given in combination with atezolizumab  ION – unresectable hepatocellular carcinoma ent required after 6 months s (tick box where appropriate)  evidence of disease progression  - advanced or metastatic ovarian cancer
ONTINUA assessmerequisite  O No  ITIATIONassessmerequisite	Patient has an ECOG performance status of 0-2  To be given in combination with atezolizumab  ION – unresectable hepatocellular carcinoma ent required after 6 months s (tick box where appropriate)  evidence of disease progression  advanced or metastatic ovarian cancer ent required after 4 months s (tick boxes where appropriate)  The patient has FIGO Stage IV epithelial ovarian, fallopian tube, or primary peritoneal cancer
ONTINUA assessmerequisite  O No  ITIATIONassessmerequisite	Patient has an ECOG performance status of 0-2  To be given in combination with atezolizumab  ION – unresectable hepatocellular carcinoma ent required after 6 months s (tick box where appropriate) evidence of disease progression  advanced or metastatic ovarian cancer ent required after 4 months s (tick boxes where appropriate)  The patient has FIGO Stage IV epithelial ovarian, fallopian tube, or primary peritoneal cancer  The patient has previously untreated advanced (FIGO Stage IIIB or IIIC) epithelial ovarian, fallopian tube, or primary
ONTINUA assessmerequisite  O No  ITIATIONassessmerequisite	Patient has an ECOG performance status of 0-2  To be given in combination with atezolizumab  ION – unresectable hepatocellular carcinoma ent required after 6 months s (tick box where appropriate)  evidence of disease progression  advanced or metastatic ovarian cancer ent required after 4 months s (tick boxes where appropriate)  The patient has FIGO Stage IV epithelial ovarian, fallopian tube, or primary peritoneal cancer
ONTINUA assessmerequisite  O No  ITIATIONassessmerequisite	Patient has an ECOG performance status of 0-2  To be given in combination with atezolizumab  ION – unresectable hepatocellular carcinoma ent required after 6 months is (tick box where appropriate)  evidence of disease progression  advanced or metastatic ovarian cancer ent required after 4 months is (tick boxes where appropriate)  The patient has FIGO Stage IV epithelial ovarian, fallopian tube, or primary peritoneal cancer or the peritoneal cancer entrement of the patient has previously untreated advanced (FIGO Stage IIIB or IIIC) epithelial ovarian, fallopian tube, or primary peritoneal cancer or Debulking surgery is inappropriate
ONTINUA assessmerequisite  O No  ITIATIONassessmerequisite	Patient has an ECOG performance status of 0-2  To be given in combination with atezolizumab  ION – unresectable hepatocellular carcinoma ent required after 6 months is (tick box where appropriate)  evidence of disease progression  advanced or metastatic ovarian cancer ent required after 4 months is (tick boxes where appropriate)  The patient has FIGO Stage IV epithelial ovarian, fallopian tube, or primary peritoneal cancer  The patient has previously untreated advanced (FIGO Stage IIIB or IIIC) epithelial ovarian, fallopian tube, or primary peritoneal cancer and
ONTINUA assessmerequisite  O No  ITIATIONassessmerequisite	Patient has an ECOG performance status of 0-2  To be given in combination with atezolizumab  ION – unresectable hepatocellular carcinoma ent required after 6 months s (tick box where appropriate)  evidence of disease progression  advanced or metastatic ovarian cancer ent required after 4 months s (tick boxes where appropriate)  The patient has FIGO Stage IV epithelial ovarian, fallopian tube, or primary peritoneal cancer  The patient has previously untreated advanced (FIGO Stage IIIB or IIIC) epithelial ovarian, fallopian tube, or primary peritoneal cancer  The patient has previously untreated advanced (FIGO Stage IIIB or IIIC) epithelial ovarian, fallopian tube, or primary peritoneal cancer  The patient has previously untreated advanced (FIGO Stage IIIB or IIIC) epithelial ovarian, fallopian tube, or primary peritoneal cancer  The patient has previously untreated advanced (FIGO Stage IIIB or IIIC) epithelial ovarian, fallopian tube, or primary peritoneal cancer  The patient has previously untreated advanced (FIGO Stage IIIB or IIIC) epithelial ovarian, fallopian tube, or primary peritoneal cancer  The cancer is sub-optimally debulked (maximum diameter of any gross residual disease greater than 1cm)
ONTINUA: -assessmerequisite O No -assessmerequisite -assessmerequisite	Patient has an ECOG performance status of 0-2  To be given in combination with atezolizumab  ION – unresectable hepatocellular carcinoma ent required after 6 months s (tick box where appropriate)  evidence of disease progression  advanced or metastatic ovarian cancer ent required after 4 months s (tick boxes where appropriate)  The patient has FIGO Stage IV epithelial ovarian, fallopian tube, or primary peritoneal cancer  The patient has previously untreated advanced (FIGO Stage IIIB or IIIC) epithelial ovarian, fallopian tube, or primary peritoneal cancer  Debulking surgery is inappropriate  O Debulking surgery is inappropriate

## HOSPITAL MEDICINES LIST RESTRICTIONS CHECKLIST

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.

Name: Name: Name: Name: Ward: NHI: MHI: MHI: MHI: MHI: MHI: MHI: MHI: M			
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 – 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 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			
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 patient has recurrent respiratory papillomatosis  The treatment is for intra-lesional administration  CONTINUATION – Recurrent Respiratory Papillomatosis			
The patient has recurrent respiratory papillomatosis  The treatment is for intra-lesional administration  CONTINUATION – Recurrent Respiratory Papillomatosis			
CONTINUATION – Recurrent Respiratory Papillomatosis  Re-assessment required after 12 months			
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)			
O Ocular neovascularisation  O Exudative ocular angiopathy			

I confirm that the above details are correct:	
Signed:	Date: