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.

	CRIBER PATIENT:		
ne:	Name:	. Name:	
rd:	NHI:		
vacizu	sizumab		
-assess	TION – unresectable hepatocellular carcinoma sessment required after 6 months quisites (tick boxes where appropriate)		
0.1	O Patient is currently on treatment with bevacizumab, and met all remaining criteria prior to commencing tre	atment	
or	O Patient has locally advanced or metastatic, unresectable hepatocellular carcinoma		
	and Patient has preserved liver function (Child-Pugh A) and		
	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		
	Patient has experienced treatment-limiting toxicity from treatment with lenvatinib		
	O No disease progression since initiation of lenvatinib		
	Patient has an ECOG performance status of 0-2		
	o i allone had an 2000 portormando dialado o o 2		
	and To be given in combination with stazelizumen		
	To be given in combination with atezolizumab		
DNITNU	To be given in combination with atezolizumab INUATION – unresectable hepatocellular carcinoma		
-assess	To be given in combination with atezolizumab INUATION – unresectable hepatocellular carcinoma sessment required after 6 months		
-assess erequis	INUATION – unresectable hepatocellular carcinoma sessment required after 6 months quisites (tick box where appropriate)		
-assess erequis	INUATION – unresectable hepatocellular carcinoma sessment required after 6 months quisites (tick box where appropriate) No evidence of disease progression		
-assesserequis	INUATION – unresectable hepatocellular carcinoma sessment required after 6 months quisites (tick box where appropriate)		
-assess erequis 1 TIATIO -assess	INUATION – unresectable hepatocellular carcinoma sessment required after 6 months quisites (tick box where appropriate) No evidence of disease progression TION – advanced or metastatic ovarian cancer		
-assess erequis 1 TIATIO -assess	INUATION – unresectable hepatocellular carcinoma sessment required after 6 months quisites (tick box where appropriate) No evidence of disease progression TION – advanced or metastatic ovarian cancer sessment required after 4 months quisites (tick boxes where appropriate) The patient has FIGO Stage IV epithelial ovarian, fallopian tube, or primary peritoneal cancer		
-assess erequis 1 TIATIO -assess	INUATION – unresectable hepatocellular carcinoma sessment required after 6 months quisites (tick box where appropriate) No evidence of disease progression TION – advanced or metastatic ovarian cancer sessment required after 4 months quisites (tick boxes where appropriate) 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 or The patient has previously untreated advanced (FIGO Stage IIIB or IIIC) 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 or the patient has previously untreated advanced (FIGO Stage IIIB or IIIC) 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 or the patient has previously untreated advanced (FIGO Stage IIIB or IIIC) 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 or the patient has previously untreated advanced (FIGO Stage IIIB or IIIC) 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 or the patient has previously untreated advanced (FIGO Stage IIIB or IIIC) epithelial ovarian, fallopian tube, or primary peritoneal cancer or the patient has previously untreated advanced (FIGO Stage IIIB or IIIC)	llopian tube, or primary	
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-assess erequis 1 TIATIO -assess	INUATION – unresectable hepatocellular carcinoma sessment required after 6 months quisites (tick box where appropriate) No evidence of disease progression TION – advanced or metastatic ovarian cancer sessment required after 4 months quisites (tick boxes where appropriate) 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, fa peritoneal cancer and Debulking surgery is inappropriate or The cancer is sub-optimally debulked (maximum diameter of any gross residual diseas		
TIATIO -assesserequis	INUATION – unresectable hepatocellular carcinoma seasment required after 6 months quisites (tick box where appropriate) No evidence of disease progression TION – advanced or metastatic ovarian cancer seasment required after 4 months quisites (tick boxes where appropriate) 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 or Debulking surgery is inappropriate The cancer is sub-optimally debulked (maximum diameter of any gross residual diseas and Devacizumab to be administered at a maximum dose of 15 mg/kg every three weeks		

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PRESCRIBER	PATIENT:		
Name:	Name:		
Ward:	NHI:		
Bevacizumab - continued			
CONTINUATION – advanced or metastatic ovarian cancer Re-assessment required after 4 months Prerequisites (tick box where appropriate)			
O No evidence of disease progression			
INITIATION – Recurrent Respiratory Papillomatosis Re-assessment required after 12 months Prerequisites (tick boxes where appropriate) Maximum of 6 doses			
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: